1	SUPERIOR COURT OF CALIFORNIA			
2	COUNTY OF ALAMEDA			
3	BEFORE THE HONORABLE WINIFRED Y. SMITH, JUDGE PRESIDING			
4	DEPARTMENT NUMBER 21			
5	00			
6	COORDINATION PROCEEDING) SPECIAL TITLE (RULE 3.550))			
7	ROUNDUP PRODUCTS CASE) JCCP No. 4953			
8)			
9	THIS TRANSCRIPT RELATES TO:)			
10	Pilliod, et al.) Case No. RG17862702			
11	vs.			
12	Monsanto Company, et al.) Pages 3457 - 3654 Volume 22			
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16	Wednesday, April 17, 2019			
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PROCEEDINGS

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(Proceedings commenced in open court out of the presence of the jury:)

THE COURT: So there are a couple things on the agenda. We don't have a lot of time. I know

Dr. Benbrook is here this morning. The jurors aren't all here yet so we have a few minutes.

I'm going to park the judicial notice for the moment just because it involved a lot of documents and probably a much longer discussion than we can have right before 9:00 o'clock. But I will say -- I didn't take the binders home, I took the briefs home just to review them last night.

And I looked at some of the documents this morning. I need more time to actually look at what it is you want me to admit. Because on the documents that I've already ruled on, I'm not inclined to change my rulings on those. Your authority didn't change my mind about the scope of judicial authority.

Now, having said that, the documents in your binders are a little different. Some of them are Federal Register, some of them are complete reports with short appendices. Generally speaking the reports would

be judicially noticeable but not the appendices of the underlying -- I still disagree about all the underlying data to be judicial noticeable.

So just sort of a broad stroke based on kind of a cursory review.

So I don't know which ones you think you need in order to cross-examine Dr. Benbrook. You can try to have a shorter conversation if that -- if we need to.

So what I would ask you to do is just identify the documents that you think you need today, and I can try to make a ruling on those. And then we can have a little longer discussion on the others.

I noticed that there was probably four or five Federal Registers -- documents from the Federal Registers. I don't think there's too much dispute about judicially noticeable the documents in the Federal Register. But the others I think are a little more complicated.

MR. WISNER: I think the one in the Federal Register, it's not a question of whether or not they're judicially noticed. It's the relevance. They're about food residues.

THE COURT: So relevance is a whole other story. You can object based on relevance.

MR. WISNER: That's right.

THE COURT: So judicially noticeable doesn't mean admissible. It just means judicially noticeable.

So you can have that conversation whenever -or make that objection whenever -- because there was
also another report that related to something in
pesticides, but I wasn't sure whether or not -- in your
brief whether or not they're actually contained in
Roundup or what the relevance is. I don't know.

So I couldn't determine that as one of my questions but probably not going to have a lot of time to talk about that this morning.

I do want to talk about the bench brief about Dr. Benbrook.

MR. WISNER: Should we excuse the witness?

THE COURT: Yes, please.

(Witness exited the courtroom.)

THE COURT: So I think Dr. Benbrook and Dr. Raj are simple. I can just deal with that. So I'm going to let the conversation about Raj come in, that's fine.

Rubenstein, I've looked at that and I've looked at it. My first instincts I think are correct because I think anecdotal information about his experience is not expertise. Just anecdotal conversation about what he's done. I'm not really going

to entertain a lot more conversation about it. 1 2 MR. MILLER: I understand, Your Honor. THE COURT: But I will tell you what I will 3 let in. MR. MILLER: All right. 5 This is 44 -- is it 44, 45, 46? THE COURT: 6 MR. MILLER: 31:17 to 25, Your Honor. 7 THE COURT: I don't think I was looking at 9 that. 10 MR. MILLER: It goes to 32:7 actually. THE COURT: So, you know, my concern about 11 this is that it's his conversation about -- even 31 to 12 13 32 is somewhat offhand. He's not really saying -there's no underlying data or opinion or information 14 15 that supports it. He just says, "Oh, yeah, they're 16 known." 17 And my concern is that he's a treating physician, and to allow him to make just general 18 statements like that, it lends a lot of support or 19 20 credence to the conclusions, which are, they're related, 21 when he has no underlying expertise or data to support what he's saying. So 31 and 32 is out. 22 23 I'm will allow him to answer the question when his interest begin --24 (Telephone interruption.) 25

THE COURT: -- which would be 7 through 12 on page 44. I'm sorry. Yeah, 7 through 12.

And the reason I'm eliminating the other lines is he's come down with blood cancers. That's not specific. That's not non-Hodgkin's lymphoma. That's just blood cancers in general.

Again, just sort of making general statements about what his patients came down with that are not related specifically to NHL or some expertise about it, again, lends, I think, some credibility to the conclusion that these -- it's like a short end to the conclusion which is, oh, yeah, blood cancers are related to exposure of pesticides. And then that's it.

So --

MR. WISNER: Your Honor, just keep going.

There's the lines 47:19 through 48:5 which is sort of a counter for that if you keep going.

There's actually a specific discussion about whether or not Roundup causes cancer. I assume that comes out as well.

THE COURT: Well, this is in any of the papers where you've written --

MR. WISNER: That part I don't mind up through 47:22. But from 47:23 to 48:5.

THE COURT: Right. That will come out.

Because, again, I just think it's general 1 conversation. So whatever is related to the more 2 3 general conversation about basically conclusory conversation about it as opposed to specific opinions and/or data that supports it. 5 When this interest begin, sir. 6 I would say January 1995. I did three months 7 in a row on the blood and bone marrow transplant unit. 9 Period. 10 MR. MILLER: I'm sorry. Where are we, Your Honor? 11 That's up to line 6 on page 45. 12 THE COURT: So then lines 4 through 10 -- I'm sorry -- line 4 13 through 8. I don't even know if you want to include 14 this because it kind of cuts him off mid sentence, but: 15 I did a lot of leukemia or some leukemia lymphoma time 16 17 Period. So that's it on that. too. How are we doing on jurors? 18

COURT ATTENDANT: Let me check right now.

MR. BROWN: Then, Your Honor, on the Benbrook issue, you indicated --

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THE COURT: Okay. So talking about

Dr. Benbrook, it looks like this bench brief really is sort of throwing up a smoke signal that there may be an issue based on what happened in Johnson. I gather

that's really the impetus for this.

2.

MR. BROWN: And as an adjunct to it,

Your Honor, is an article that was published this month
that was prepared by Robin Mesnage, Charles Benbrook,
and Michael Antoniou. And apparently it was published
in something called Food and Chemical Toxicology, and it
was published this month online and it came well after
the disclosures in this matter.

And in this document or in this paper, which is really not a scientific paper, but it goes into a lot of issues concerning GMOs, it goes into a lot of information concerning misleading information being provided by manufacturers. It goes into --

THE COURT: Is this an exhibit today?

MR. BROWN: Yeah, I anticipate -- I anticipate that they intend to use this. And it's -- this is not something that has been produced in discovery or --

THE COURT: Well, let me just stop you.

Are you going to use it?

MR. WISNER: Yeah, absolutely. It was already used with Dr. Sawyer extensively. It's been published to the jury and discussed at length.

THE COURT: That document?

MR. WISNER: Yeah. I'm not entirely sure what the objection is at this point. This is about the POEA

toxicity, and it was published by scientists including Dr. Benbrook, peer reviewed. Dr. Sawyer talked about it with the jury and talked about how POEA was more toxic than just glyphosate.

That was Dr. Benbrook's article. We showed it to the jury and we discussed it. That's the only part of that article that will be used today. We're not going to talk about GMOs, that's excluded. And we're definitely not going to talk about -- what was the other issue? Misleading? We're not going to talk about that stuff.

MR. BROWN: Misleading. There's information in here about poisoning cereal and children eating cereals.

THE COURT: So to the extent that it's been published, I don't think there's really an objection. If there's anything else that you're planning beyond what's already been published, then tell me now --

MR. WISNER: Absolutely not.

THE COURT: -- so that I can -- you know, we can talk about that before Dr. Benbrook gets on the stand and we're talking sidebars.

MR. BROWN: Yeah, but Your Honor, when you say "published," this article, I don't recall this article being published to the jury. And what he's saying is

it's Dr. Benbrook's article. And he's talking about the substance of what it is as opposed to the particular article which Dr. Benbrook is on here as an author.

And it's --

THE COURT: Why don't we first clarify whether or not any part of it has already been published. I think that might --

MR. EVANS: Your Honor, I believe in the context of Dr. Sawyer talking about POEA, the article was published with respect to POEA. But Dr. Sawyer was already talking about POEA, and this is a published article so I wasn't going to object because we had an agreement that published articles can be displayed.

THE COURT: Okay.

MR. EVANS: But that's a completely different discussion when you have a toxicologist. You know, Dr. Benbrook today, he's an economist. So the part of this article that he has expertise on is not POEA, and it's also cumulative to what Dr. Sawyer's already testified about.

I mean, I'm not -- I don't know why we're going to have an economist go through the POEA piece of the puzzle that we've already heard about, candidly from several people.

THE COURT: Okay. So what --

MR. WISNER: So I'm trying to get my objection straight here. So the first one was --

THE COURT: Okay. So it's already been published so that portion of published talking about POEA. I think we're talking relevance. I think Mr. Evans' objection at this point is relevance because he's an economist and he's here to talk about something else and not opine about POEA.

And I don't know if his expertise includes whatever that particular part of the article is. You'd have to show it to me so I can figure out is it scientific -- was there some part of it that was scientific that really wasn't his bailiwick but there was some part of it that was economic that was his bailiwick?

MR. WISNER: So, first of all, this idea that he's just an economist is just not true. He's published multiple peer-reviewed literature about the science involving glyphosate. He's currently a researcher on an ongoing project involving the toxicity of glyphosate in farm workers and animals in the Midwest. He is a scientist.

THE COURT: Just give me an overview. What's his background?

MR. WISNER: He has a Ph.D. in economics, but

then he proceeded to work basically his whole life in pesticides. He's been doing research both on the economics of it, the regulatory aspect as well as the science. He's published in this area. And we'll lay that foundation in spades today.

2.

One of those publications is the very one that they're talking, which is a scientific publication, peer-reviewed, and published in an academic journal where he talks about the toxicity of these various surfactants.

Now to address the issue of relevance, we're not going to be cumulative. We're going to touch on it very briefly. I'm going to ask him what they looked at when they came to these conclusions because the jury has already heard from Dr. Sawyer. He can talk about that since he wrote it. And that's about it.

So I think that it's valuable testimony insofar as understanding what this article supports, coming from what it's based on, and how they came to the conclusions that they did.

But finally, Your Honor, this discussion of POEAs and their toxicity is in Dr. Benbrook's report. So it's not like this is a new opinion. So all of this seems to be -- I'm not clear what the problems are.

THE COURT: Well, I'm assuming that whatever

the other objectionable portion of it is, which is GMOs and other things that are excluded, aren't going to be touched on.

MR. WISNER: That's right.

2.

THE COURT: So as long as -- well, let's just see. I mean, if you lay the foundation, you've laid the foundation, he can testify. If you haven't and there's no objection, I'll rule on it at the time. I don't think that preliminarily excluding something based on not enough information is something I'm going to do.

MR. WISNER: Sure.

MR. MILLER: Your Honor, I'm not rearguing
Rubenstein. Your Honor has ruled. But Your Honor ruled
yesterday as well that it was either going to be in or
out. So we --

THE COURT: I did exclude the other several lines at the end of the conversation.

MR. MILLER: Right. I just wanted to show
Your Honor, page 55, if we're not allowing
Dr. Rubenstein to say that pesticides are a risk, you
clearly can't have him say that smoking or autoimmune
disorders are a risk or being older without saying that
pesticides are a risk.

So we're asking the Court to strike page 55, lines 6 right through 56, line 25. It wouldn't make any

sense to have him be precluded from saying what he knows as an expert that pesticides are a known risk and yet have counsel go on and talk about every other risk factor that they want to talk about.

2.

THE COURT: Well, no, because he's talking about in your appearance as an oncologist and specifically someone who was working with non-Hodgkin's lymphoma, specifically PCNSL, you see other risk factors that are signals for this disease.

So he's talking about something that is within his expertise as a treating oncologist, where the other information is anecdotal in his history. That's the difference.

MR. MILLER: Squarely not, Your Honor.

Page 31, line 17. That's simply not accurate, with all due respect. He says at page 31, line 17: In the course of your treatment of Alberta Pilliod, that it's clear it's in the course of his treatment, and with your experience -- and no one has challenged he is the world's leading expert on this -- is pesticides a known association with non-Hodgkin's lymphoma like the one she had? And the answer is yes.

So it's impossible --

THE COURT: No. The answer is:

Well, pesticides are known to be

1	associated with blood cancer.	
2	MR. MILLER: Which is what she has.	
3	THE COURT: (Reading:)	
4	There's a lot of data in the	
5	literature that blood cancers, leukemias,	
6	and lymphomas are strongly associated with	
7	blood cancers.	
8	MR. MILLER: See the next question and answer.	
9	She has a blood cancer like the type	
10	of blood cancer she has.	
11	So for him, either we go all in or all out.	
12	Either he's allowed to say that or clearly we can't go	
13	on and have him say smoking or getting older or	
14	autoimmune disorders. Either it all comes in or it all	
15	comes out in any fair playing field. That's all I'm	
16	asking.	
17	THE COURT: I'm not beating a dead horse. I	
18	don't want to keep talking about this. I've ruled.	
19	Right or wrong, end of story.	
20	MR. MILLER: Has Your Honor ruled on page 55	
21	or 56, is all I'm asking?	
22	THE COURT: No, I'm not. End of story. I	
23	ruled.	
24	So let's get back to Dr. Benbrook.	
25	Is that it with respect to Dr. Benbrook?	

There were some other issues regarding his political affiliation and just --

2.

MR. WISNER: Not coming in. I met with counsel yesterday after I saw the bench brief, and I told them Dr. Benbrook is not going to talk about politics at all. So that's not an issue.

There was one topic that said anything that might suggest that Monsanto misled the EPA, and I spoke with counsel and I said, listen, that's an incredibly broad umbrella. I mean, there's facts that he's going to talk about that could reasonably lead to a conclusion that they misled the EPA. But he's not going to say that Monsanto misled the EPA. That's not coming out of his mouth.

We are going to talk about, for example, the tumor story and what happened there. And that story and the facts surrounding that, someone could conclude it falls under the "might suggest that Monsanto misled the EPA," but he's not going to say that Monsanto misled the EPA. He's going to say this happened, this happened, this happened.

That was the only consideration I had about the bench brief, it was vaguely overbroad and swept in a potential amount of stuff that would be --

THE COURT: I've forgotten the tumor story.

Is this the magic tumor?

MR. WISNER: I'm not allowed to call it magic,
but, yes. The magic tumor story.

MR. BROWN: Your Honor, what you have indicated that Dr. Benbrook can testify about is that Benbrook may provide testimony that summarizes and explains the regulatory framework for herbicide regulation, including Monsanto's legal obligations on registration matters, period.

The rest of the *Sargon* order excludes his testimony concerning all of these other issues. And the reason for it is, is because he has no expertise in any area that is relevant to this particular case.

Even the issue that this agricultural economics is not relevant within the context of this, and if the Court looks at -- again at page 5 of the -- I'm going to have the tentative order on *Sargon* motions, motion for summary judgment.

THE COURT: There was a final order, by the way.

MR. BROWN: Yeah, but -- and I just don't have that in my binder.

THE COURT: Okay.

MR. BROWN: But it clearly delineates what Dr. Benbrook is being called to testify about.

And, you know, we want to slice the bologna very thin here by saying how close can we walk to the line, and I think that that is something that we want to avoid.

And so we've got to read this in the true -
THE COURT: Let me just interrupt you to say
this, because I'm not oriented to precisely what he's
going to say that's objectionable about the tumor.

Why don't you walk me through what that looks like.

MR. WISNER: Your Honor, I will. But let me back up really quickly.

So his report is like 300 pages long, and it has a whole bunch of facts and stuff that he's proffered to offer testimony about.

Monsanto moved to exclude specific portions of that testimony. And Your Honor ruled on the specific portions of his report. This is not an inclusive statement of what he's going to testify. It is your ruling on what's in front of you, right, and you're going through the topics and saying granted or denied.

But if there's stuff that they didn't move to exclude, that isn't excluded by virtue of them not having raised the issue with the Court in the first place. That's a logical fallacy.

So Mr. Brown said he's only allowed to testify about that, and that's not what the Court had ordered.

If they had moved to exclude reference to the magic tumor story, right, which they did and they lost in a motion in limine.

So him talking about the magic tumor story to the extent he's qualified to talk about it, which he's entirely capable of doing, I think it's not even at issue in this order. They're trying to sweep in hundreds of pages of uncontested opinions and by saying, well, you only said he can testify..., that's just not correct.

THE COURT: Well, the motion was to exclude all of his testimony.

MR. WISNER: Sure.

THE COURT: And so I'm just looking at the order. So they did move to exclude everything. And then I said, okay, he can testify about the following three.

MR. WISNER: That's not correct. They didn't move to exclude everything. They moved to exclude him generally under *Sargon* as not being qualified.

THE COURT: Right.

MR. WISNER: You denied that. You didn't go down that road in that order.

THE COURT: No, I didn't. But I did 1 2 circumscribe what he could say. 3 MR. WISNER: Well, fair enough, about the things that they challenged. Your Honor didn't read his 4 300-page report. 5 6 THE COURT: No, you're right. I did not. They didn't challenge the 7 MR. WISNER: 300-page report. They didn't say this topic he's not qualified to do, this topic --9 THE COURT: I skimmed it because I wanted to 10 see what was in it, but I couldn't tell you right now 11 12 what's in the 300-page report. 13 MR. WISNER: Sure. But my point is if I had 14 known -- so I can only respond to what they challenged. 15 That's how a motions practice works. 16 THE COURT: Well, let me ask you this. How 17 does the tumor story work into just explaining the regulatory framework for herbicide regulation --18 19 MR. WISNER: It doesn't. 20 THE COURT: -- the legal obligations and 21 registration matters? MR. WISNER: It could fall under that, but I 22 23 don't think that's what the issue is here. Your Honor 24 didn't exclude him testifying about that.

THE COURT: No, I specified about what he

25

could talk about. That was my intent, was to specify what he could talk about.

There was a motion to eliminate him under Sargon. And I said no, I'm not going to eliminate him under Sargon. I will allow him to talk about the following few things, which are testimony that summarizes and explains the regulatory framework for herbicides including description of Monsanto's legal obligation on registration matters. The Court orders that Benbrook may not testify on whether Monsanto complied with legal obligation. Orders that he may not testify on the following other things.

MR. WISNER: Yeah.

THE COURT: And I think what you're telling me now is --

MR. WISNER: Because those things that you ruled on were things specifically challenged by Monsanto.

MR. BROWN: Your Honor, that's disingenuous.

MR. WISNER: Hold on. Let me finish my argument.

THE COURT: I have to go back to the Sargon motion now.

MR. WISNER: Please. Because I've read them closely, and they did not -- they did not specify topics

that you ruled on. And if they're -- you can't do that. You can't say -- they did challenge everything and you didn't strike him as unqualified across the board. So they lost the general -- that was a small portion of the challenge.

Then they said, okay, this opinion, this opinion, this opinion, this opinion, and you ruled on those opinions that they challenged.

They never challenged his opinions about the magic -- actually they did. They tried doing it in the motion in limine.

THE COURT: Well, what does that have to do with his expertise on -- whatever his expertise is?

What does the magic tumor story -- how does that relate to whatever expert opinion he has about registration and compliance? Maybe that's the question I'm really asking.

MR. WISNER: Sure. Fair enough.

He's an expert in the regulatory process at the EPA. He can walk us through how -- we have all these documents that are in evidence, that came into evidence already, they're already in. And he can talk about what they mean and what the obligations of Monsanto were at that point, what that classification meant. And so he can walk through the story and he can

explain from a regulatory perspective what was going on.

THE COURT: Okay.

MR. WISNER: So I think that's 100 percent within that framework.

THE COURT: So is this magic tumor story -I'm sorry, I keep calling it a magic tumor story.

MR. WISNER: I know. It's catchy.

THE COURT: But whatever, so is the tumor report part of that in some fashion?

MR. WISNER: Absolutely. Because the entire interchange is with the EPA. Right? And so they're saying, oh, we want you to do another study, data call in. What does "data call-in" mean? I need him to explain these things.

I need him to explain what happened -- for example, he's going to talk about that IBT and what that IBT document that's in evidence means. What does it mean when it says data call-in? What does it mean when it says it's invalid? I'm going to tie it to the current EPA report. I'm going to walk him through all the stuff that's come into evidence already so the jury has some context. And that's all in his report. And that wasn't even moved to be excluded.

So --

MR. BROWN: Your Honor, he keeps talking about

a report. And let me be clear that this is a report that was prepared not in this case. Okay? So it was prepared in some other case.

2.

And in anticipation of the testimony, it was -- the motion was filed.

And the Court is absolutely correct because the motion was to exclude Dr. Benbrook entirely because all he has done is read some literature and then he wants to render opinions about it.

Now, we've heard already from Dr. Sawyer.

We've heard from Chris Portier. We've heard from

Dr. Jameson. And those documents that he's referring to that have come in have been testified to by all of those people.

THE COURT: Okay.

MR. BROWN: And what the Court has said, and again going back to the order, is the Court reserves trial -- reserves for trial whether Benbrook may explain the context and possible meaning of regulatory technical documents, which have already been testified to on three occasions by the witnesses who've gone before Mr. or Dr. Benbrook.

And, again, Your Honor, Dr. Benbrook is not a toxicologist, he's not an epidemiologist, he's not a medical doctor, he is not -- he is an agricultural

economist. And that does not qualify him to simply read articles, formulate opinions that are consistent with his own personally held beliefs, and then to come in and proselytize to the jury.

THE COURT: I guess counsel is telling me that he is an expert because -- or he's going to attempt to lay the foundation as an expert, I guess, in this area. And I don't know whether or not he was offered for that in his -- you know, I'd have to go back do the designations --

MR. WISNER: Sure.

2.

THE COURT: -- to see how he was -- which I don't have in front of me right now either.

MR. WISNER: Just to be clear, so much factual inaccuracy here.

THE COURT: I'm just asking what was he offered as? Because it's 9:20 and the jurors are here.

MR. WISNER: Because that report that he said wasn't in this case was disclosed in this case specifically as part of the designation. So that's just completely untrue.

THE COURT: So what opinions did -- you know, if I'd known we were going to have this kind of fight, I would have suggested we stay later yesterday because now we're opening up a whole, you know, front that I had not

anticipated having to discuss.

But the designations, what exactly was he designated to testify about?

MR. WISNER: We served a 300-page report with all of his opinions in detail in this case. It's the same one from the MDL, and we served it in this case. So they had full knowledge about it. They chose not to depose him. I'm not sure why they chose not to.

Second, Your Honor, and this is another factual inaccuracy, the very documents that he says other experts have testified about, they haven't. They weren't even in evidence yet. They just came into evidence through the videos that were just played. So that's why they're going to be talked about now, we're going to explain the technical aspects of them.

This -- and that he's not an expert, he's just an economist, also factually untrue. He's published scientific literature on this very issue in peer-reviewed articles that they had access to and were fully disclosed before he gave his opinion.

So I'm not sure what Mr. Brown is talking about. Maybe he hasn't been involved in the litigation up to this point. But this is the facts.

MR. BROWN: I think I have. And by virtue of publishing an article does not make someone an expert on

any particular point. 1 2 **THE COURT:** Does anyone have his report, by 3 chance? Yes, I have it here, Your Honor. 4 MR. BROWN: THE COURT: Can I see it? If you have it. 5 6 MR. WISNER: And to be clear, Your Honor, we're not going to offer any opinions that you excluded, 7 right. He's not going to talk about Monsanto's motive 9 or any of that stuff. And really under CCP 10 MR. BROWN: Section 2034.010, incorporating a report from some other 11 12 case does not even satisfy the requirements of the 13 section. So if he's -- if he's citing to and saying, well, we're -- he's going to testify about everything 14 15 he's testified to over the last 15 years, is not in compliance with the code. He's supposed to say 16 17 specifically what he is going to be offered in this case and specifically what is being offered in the case. 18 MR. WISNER: Your Honor, we specifically gave 19 20 them this report --THE COURT: Did he testify to this stuff in 21 the Johnson case? Or --22 23 Yes. And we specifically served MR. WISNER: this very report as part of our CCP disclosures. 24

factually untrue. I'm sorry. I don't know why

25

1 Mr. Brown is not familiar with what's happened in this 2 litigation, but that's the facts of it, Your Honor. 3 I've been doing it for the last three years. MR. BROWN: Doing something. MR. WISNER: Not making stuff up. 5 THE COURT: Okay. So I'm going to take a 6 break and see where the jurors are. 7 And I'm not exactly sure. I feel like I got 9 blindsided this morning, and I don't appreciate it. MR. WISNER: Me too, Your Honor. 10 11 MR. BROWN: I take responsibility for that, Your Honor. It's my fault. I apologize. 12 (Recess taken at 9:23 a.m.) 13 (Proceedings resumed in open court out of the 14 15 presence of the jury at 9:34 a.m.) 16 **THE COURT:** I don't have time to sort through 17 this. It's just really a lot. So I'm going to say this. 18 The order that I issued and the order that 19 Judge Karnow issued, which was one of the bases on which 20 21 I reviewed this, was in the order it states specifically: Benbrook is offered as an expert -- this 22 23 was my summary of how he was offered -- as an expert on whether Monsanto's conduct is as a pesticide 24

manufacturer and registrant comports with its obligation

25

and stewardship responsibilities.

Had no idea, concept that was also being offered was expert testimony on the science. That's not part of what I understood his proffered expertise was on, that this was limited to laying of conduct as a pesticide manufacturer, registrant, and comports with an obligation and stewardship responsibilities.

Now I looked basically at the table of contents, and, yes, there is a lot more in his report.

But what else is in his report has a lot to do with mouse oncogenicity. That was not my expectation that he was going to touch on those subjects. I thought he was going to touch on a fairly narrow subject because there are all kinds of experts on all the other stuff.

MR. WISNER: Sure.

THE COURT: So I'm not sure where you're going to with this, where you want to go with this. I can't -- I'm not going to make any preliminary rulings because we have to wing it. Here we are. But that would be my expectation with respect to Benbrook's testimony.

You know, whether or not -- whether there's factual testimony that touches on whether they misled the EPA or not, that's a fine line. I'll just entertain objections if we get too close to the third rail, if

that's what I think is happening.

MR. WISNER: Sure.

THE COURT: But that's the general scope of which I'm expecting him to testify and pretty much where I expect you to go today.

MR. WISNER: Okay. I mean, your Honor, I just want to clarify. I actually double-checked the disclosures, and in our disclosures in this case we not only specifically referenced the report that you just got handed, we attach it and serve it. So I just want to make clear for the record there's no question that they were given this.

The other issue, Your Honor, is, you know, there's a sort of timeliness issue here. Right?

Because as part of the *Sargon* process, they need to tell us what in his disclosed opinions they're moving to exclude. And they chose to focus on the issue that you just discussed, stewardship obligations and whether or not he complied. And your ruling specifies what he can and cannot do with regards to that. And that's fine.

My understanding from the very beginning because of the way this process works is if they had other opinions of his that they sought to exclude, they had to move to exclude them. And so when you read the Sargon briefing, it's focusing on this exact issue and

Your Honor ruled on that issue.

But if they chose not to challenge anything else -- I mean, I understand they're moving to exclude his opinion in its entirety, but then they go on to say his only opinion is stewardship, which is false as you can see from his report.

And so at a certain point -- I agree we have to wing it at this point, and you can tell me if we've gone too far and we'll move on and do our thing. But at a certain point, there has to be some frankness and candor to the Court from the defense counsel about what they're seeking to exclude.

And if they were seeking to exclude his opinions about the oncogenicity studies from 1983 because he wasn't able to offer opinions about it, which is clearly in his report, they should have moved to exclude it. But they didn't. And in so failing to take action, they waive that pretrial ruling.

Now if they want to argue he can't offer the testimony at trial, I'm going to lay a foundation that he's qualified to do it, and if he is, he is and we're done.

But, I mean, at some point, you know, they want to micromanage what we do, and that's fine, but the Sargon process was their way to do that. And their

failure to take action has to mean something. Instead they're sandbagging us the morning before he takes the stand.

And I'm sorry if I got a little frustrated there because I felt like I'm in a land I didn't understand. Because I understand what we briefed and what we fought and we argued and I got it. But it was never about these things. And so now they're saying, oh, included that too, gotcha. And that's not how I litigate. I don't think --

THE COURT: Did he offer scientific opinions in the San Francisco case?

MR. WISNER: Absolutely.

THE COURT: And Judge Bolanos didn't -- there was no objection to that? I mean, I guess I'm having a problem with -- this morning I realize, well, there seems to be a fundamental disagreement about the Sargon motion altogether, which I didn't think there was at the time I ruled on it. And I wish I had known that. We would have had this conversation yesterday. Or at least I would have been alerted to it. Because my understanding and my expectation is that he's going to testify about a fairly narrow area in which he's going to talk about the registration and the --

MR. WISNER: Sure.

THE COURT: And so I had no sense that he was going to be testifying about any science. But --

MR. WISNER: Probably part of the confusion, Your Honor, is -- it's all context-dependent. Right? So he's not going to go up there and say: I looked at the pathology slide from the mouse tumor in 1983 and concluded there was no tumor. That's not his -- he's not a pathologist, he's not going to say that. Right?

It's -- he's a science and a science opinion -- he's going to say, though, that when you add that tumor to the control group, it no longer makes it significant. We've already heard that. That's not a new opinion really. It's not even controversial.

And then we're going to talk about the significance of it being a class C oncogen on the ability of Roundup to be used in farm product at the time in 1983. Because he's going to talk about the Delaney Clause and how it affects food tolerances.

All of that is clearly in his report. I mean, and it's all -- he wrote the statute essentially. I mean, he's an expert in this area.

So when you say is there science opinion, there's some factual predicates to the science, but he's not going to be interpreting raw data or anything like that. And he's not going to be offered for that.

THE COURT: All right. Well, we're going to have to -- I'm going to have to fly blind a little bit here. But I'll just entertain objections as we go along and we'll just see whether or not it's admissible and we'll go from there.

MR. BROWN: Yeah, and Your Honor, as we move forward in terms of what we're doing, I am hoping that we're going to do it within the confines of the order --within the confines of the order the Court has made post Sargon because that does structure this. And, again, I don't see how anyone could read it to include the things that counsel is talking about.

And, secondly, let me say that he has indicated that what things about the mouse study he says are already in and in front of the jury, he's absolutely correct --

THE COURT: Well, let me just stop you there. Let me hear it because if he's talking about what it meant in terms of -- if it's this versus that, and this is what it meant in the regulatory framework, it may very well be relevant. They've already heard it. They've heard a lot of things in different ways from different scientists. So that's really not of any particular concern.

If it's not relevant to his framework, his

regulatory framework, and I'll just be looking at it through that lens. When you say post order, what are you talking about specifically?

MR. BROWN: Well, I'm talking about the days after the Court issued its Sargon ruling.

And what I'm saying is we've got to be going pursuant to what the Court has included in the order.

THE COURT: Yeah, and I'm saying I'm looking at the -- I will be hearing the testimony sort of through that lens. But I would also say to you that if there is -- if in the context of that, the status of the science makes a difference, then that may very well be relevant and not objectionable.

And so if we're talking about -- if we're talking about the science in that context, that may be relevant. If we're talking about pure science, I can't imagine you want them to hear it a 19th time, but that's up to you. You can bore your jury to death if you want to. But you understand what I'm saying.

MR. WISNER: Absolutely.

THE COURT: It's 9:40. And it's time for the jury to come in. And we will take it from here.

MR. WISNER: Your Honor, here's the binder.

I'll give it to you now.

MR. BROWN: Do you want to keep that? You're

```
1
       welcome to it.
 2
                  THE COURT: You know what, no. Do you need it
 3
       to listen, to follow along? I'll hold on to it for a
       minute. It might be helpful as I listen to his
 4
       testimony.
 5
                       (Recess taken at 9:43 a.m.)
 6
                   (Proceedings resumed at 9:44 a.m.)
 7
                  (The following proceedings were heard in the
 9
       presence of the jury:)
                  THE COURT: Good morning, ladies and
10
11
       gentlemen.
                  We are going to proceed now with plaintiffs'
12
       next witness.
13
                 Mr. Wisner will introduce him.
14
15
                 MR. WISNER: Thank you, Your Honor.
                 At this time the plaintiffs call Dr. Charles
16
17
       Benbrook.
                              If you would, please.
18
                  THE CLERK:
                  THE COURT: Would you like to stand to be
19
       sworn first.
20
21
                  THE WITNESS:
                                Sure.
22
                  THE COURT: Thank you.
23
                  THE CLERK: Please raise your right hand.
       ///
24
       ///
25
                                                               3495
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CHARLES BENBROOK, 1 called as a witness for the plaintiffs, having been duly 2 3 sworn, testified as follows: THE WITNESS: I do. 4 THE CLERK: Thank you. Please be seated. 5 And would you please state and spell your name 6 for the record. 7 THE WITNESS: Charles Benbrook. 9 C-H-A-R-L-E-S, B-E-N-B-R-O-O-K. 10 MR. WISNER: May I proceed, Your Honor? 11 THE COURT: Yes, you may. 12 **DIRECT EXAMINATION** BY MR. WISNER: 13 Dr. Benbrook, could you please introduce 14 Q. yourself to the jury. Tell them where you're from and 15 16 where you currently live. 17 I am -- I'm -- I live in Troy, Oregon, Α. which is in the northeast corner of the state. I've 18 19 lived and worked there for 16 years. I've spent a 20 number of years working in the area of pesticide use and risk and regulation, mostly through my small consulting 21 business. 22 23 All right. Sir, I want to talk to you a lot 0. about your background for a minute. Let's start off 24 from the beginning. 25

Where did you go to college?

- A. I went for undergrad, my undergrad education was at Harvard. And my graduate degrees were from the University of Wisconsin Madison.
 - Q. And what did you study at Harvard?
 - A. Economics.

- Q. And what did you study at the University of Wisconsin?
 - A. Agricultural economics.
 - **Q.** What is agricultural economics?
- A. It's a field of study that tracks the economic consequences of decisions farmers make. It strives to understand the impacts of policy and institutions and new technology on agriculture production in the food system and the cost of food.

Agricultural economists are often among the people that get heavily involved in the study of various policy issues, including things like pesticide regulation.

So that was sort of what got me into the overall policy arena, impacting decisions made by farmers and how government actions shape the way farmers farm and control pests, for example.

Q. All right, Dr. Benbrook. Two things. I'm bringing you a bottle of water. Okay?

1	A.	Thank you.
2	Q.	And the second one is I'd like you to slow
3	down.	
4	A.	Okay.
5	Q.	Okay. Great. So let's go through your
6	backgroun	d.
7		If you look in your binder, Exhibit 3091. Is
8	that a fa	ir and accurate copy of your CV?
9	A.	Wow, there's a lot of tabs here.
10	Q.	Sorry.
11	A.	39?
12	Q.	3091.
13	A.	Okay. It's kind of near the end. There it
14	is.	
15		Yes, it is.
16	Q.	Okay.
17		MR. WISNER: Your Honor, permission to
18	publish.	
19		THE COURT: Any objection?
20		MR. BROWN: No, Your Honor.
21		THE COURT: Granted.
22		(Exhibit published.)
23	BY MR. WI	SNER:
24	Q.	All right. So we're looking here at a copy of
25	your CV.	And we talked about that a second ago. I just
		3498

kind of want to go through the first part here.

We have here obviously your personal information. Do you see that?

A. Yes, sir.

- Q. And you talk about your education. You said you have a Ph.D. in agricultural economics. That's what we were talking about a second ago.
 - A. Yes, sir.
- Q. All right. Quickly, hobbies: Raising rabbits, what is that?
- A. I raise rabbits. Some people raise dogs, some people raise pigeons. I raise rabbits. And we take them to shows just like a national dog show, and they're judged on a standard of excellence. And, yeah, it's my hobby.
- Q. So you go to shows and, like, show rabbits like dogs?
- A. Well, you don't lead them around on a leash. They're put up on a table, and the judge feels their fur and checks out their body conformation. You know, some of us would be a little heavy, some might be a little long.

But, yeah, so it's a wonderful hobby. It gets you away from the computer and out of the house.

Q. All right. So let's talk about your

employment history. And start off from when you got your graduate degree.

So the first thing -- I'm sorry. So it looks like from 1979 to 1981, was that during the time you were getting your Ph.D.?

A. Finishing it, yes.

- Q. You were at the Council on Environmental Quality, at the Executive Office of the President. What is that?
- A. It's a very small policy shop within the Executive Office of the President. At that time Jimmy Carter was President. And it advised the White House and the executive branch on environmental issues in all areas of the economy.

And I was hired as an agricultural expert and worked on a number of projects that CEQ, that's the acronym for the Council on Environmental Quality, had undertaken during the Carter administration and wanted to get done before the inaugural bringing in the next president.

- Q. And then you have you left there and worked at the subcommittee on department operations research and farm agricultural -- Committee on Agriculture, U.S. House of Representatives.
 - A. Correct.

Q. What was that?

A. So I was very fortunate because I was able to move from working on a cluster of agriculture environment, public health issues in the -- for the administration to a professional staff job in the Congress, of which there were relatively few. And with the presidency going from Democrats to Republicans, there were a lot of people out of work.

MR. BROWN: I'm going to object. That's irrelevant. I'm going to ask to move to strike and subject to the Court's prior ruling.

THE COURT: Yeah, let's move past the politics.

MR. WISNER: Oh, sure.

- Q. Let's not talk about the politics.
- A. Okay, I'm sorry.
- Q. What did you do there?
- A. I was the staff director of the congressional subcommittee that had jurisdiction or responsibility for the national pesticide regulatory law called FIFRA.
 - Q. And what is FIFRA, sir?
- A. It is the Federal Insecticide, Fungicide and Rodenticide Act. And it establishes the criteria upon which the EPA is supposed to make decisions about whether a pesticide can be used and how it can be used

and under what conditions. And so the statute addresses all aspects of the regulatory process.

- Q. And as part of your work at this committee, did you have to investigate the implications of EPA regulations on pesticide use?
 - A. Yes.

- Q. And how did you go about doing that?
- A. You hold hearings. And we had, in the three years that I was the staff director of the subcommittee, we had a very extensive set of hearings on pesticide issues because there was a lot of demand for changing national policy at the time.

We probably held over two dozen hearings in the three years. At essentially every hearing, a representative of the EPA would be invited and would explain what they're doing, how they're implementing the law, and whether they -- they thought there were aspects of the law that needed to be clarified or changed.

- Q. Did you, in that capacity, also interface with scientists?
- A. Oh, for sure. At every hearing that we scheduled, it was sort of my job as the staff director to identify and invite outside scientists that had expertise in the particular issues.

For example, we held a hearing on

cancer-causing pesticides and how EPA was conducting its risk assessments and evaluating the risk. And for that hearing, we invited a scientist from the National Cancer Institute, for example.

Q. Was that scientist Dr. Aaron Blair?

- A. Yes, it happened to be Dr. Aaron Blair.
 That's when I first met him.
- Q. All right. So then moving on to that, you began work for, it looks about six years at the National Research Council, National Academy of Sciences.

What is the National Academy of Sciences?

A. The National Academy of Sciences is an independent advisory body set up actually by President Lincoln to provide scientific and technical advice to the federal government. It's not part of the government, but it's set up to provide independent scientific and technical assistance to different federal agencies.

I was recruited actually from my staff director position on -- in the U.S. Congress where we were doing, in effect, studies on various agricultural issues including pesticides. I was recruited to come and build a new program in the National Academy of Sciences on agriculture.

So I became the executive director of what was

called the board on agriculture. And we designed and carried out, in my seven years there, oh, maybe 50 projects of which, you know, a quarter of them either were directly about pesticides, pesticide risk, and pest management, or in which pesticides and pest management was an important part of the project.

2.1

- Q. And when you were working at the National Academy of Sciences and specifically on these pesticide projects, were you reviewing, considering, and discussing or writing reports about, like, scientific articles and publications?
 - A. Oh, yes. That was part of the job.
- Q. And as part of that job, how were you able to keep up with the scientific aspects of it?
- A. Well, one of the terrific things about the National Academy of Sciences is it's considered in the scientific community a great honor to be invited to serve on a committee. All of our projects were done by committees of independent scientists, most of them academics working at different universities.

We would have between 10 and 15 members. And so these people would come to Washington two or three times a year for two or three days in producing their report. And so I got to spend days with the top scientists in every area that we were conducting a

project on.

I was young, it was an early part of my career, and I was just a sponge for knowledge. And it was a terrific place to work to learn about, you know, what was going on in the area.

- Q. Following your time with the National Academy of Sciences, we have on your résumé here, it looks like you worked for about 10 years -- I'm sorry -- for about six years with the Organic Center, chief scientist. Do you see that?
- A. So when I left the National Academy of Sciences towards the end of 1990, I started my own little one-person consulting firm, Benbrook Consulting Services. And really I continued working on the same cluster of issues that we had worked on during my years in the National Academy of Sciences when I was the executive director of the board on agriculture.

And actually the biggest issue that I worked on in that next decade from 1991 through 2000 was the Delaney Clause and the impact of federal law on the regulation of cancer-causing pesticides. Over half of my contracts involved that in one way or another.

Q. After that it looks like for about three or four years you were on the faculty of Washington State University; is that right?

A. Correct.

- Q. And what did you teach there?
- A. I didn't have a teaching appointment. I had a research appointment. And I was -- my mission there was to develop new analytical systems to quantify the impact of agricultural production systems and technology on the environment, on human health, on wildlife, et cetera.
- Q. And during this time that you've been a consultant for Benbrook Consulting Services, have you participated in any sort of hard scientific projects?
- A. Well, yes. I've been -- either through my own individual research or as part of a broader team, I've published, you know, over 30 peer-reviewed papers in a wide range of journals that have reported original analytical work on, again, trying to evaluate how agricultural production systems, pest management systems, different pesticides, different policies impact things that, you know, we all care about: The safety of food, rates of cancer, birth defects, and the cleanliness of water, and the productivity of our agricultural systems.
- Q. Can you tell the jury about any current projects you're working on, scientific projects that relate to glyphosate?
 - A. Well, I've just published a new paper with two

colleagues from King's College in London on the surfactants that are mixed in with glyphosate to produce Roundup and other, I'll use the term glyphosate-based herbicides, or I'll say GBHs. So that refers to a herbicide often made by a company other than Monsanto that contains glyphosate.

For the first, you know, 30 years of my professional career, Monsanto was the sole manufacturer and completely responsible for everything to do with Roundup.

MR. BROWN: Excuse me, Your Honor. I'm going to object. There's no foundation. This is all speculation and argumentative. And it's also nonresponsive to the question.

THE COURT: I'm going to so strike "for the first 30 years." And why don't we go on.

BY MR. WISNER:

- Q. Sir, have you looked at who manufactured and sold Roundup in the United States for the first 30 years of your career?
 - A. Yes.
- Q. Who manufactured and sold Roundup for the first 30 years of your career?
 - A. Monsanto.
 - Q. Okay. Now I want to go through some of these

1 publications that you mentioned. Here we go.

So that publication you just mentioned, is that the one right here that just came out?

A. Yes, sir.

2.1

- Q. And was that a publication that specifically looked at the toxicity of surfactants?
- A. Yes. It tried -- it tries to report to the broader scientific community several factors why there's a lot of confusion among scientists about the toxicological properties of glyphosate-based herbicides and Roundup. Because scientists outside of the industry, scientists that don't work for one of the companies, don't have access to the confidential statements of formula --
- MR. BROWN: Excuse me, Your Honor. I'm going to object. It's nonresponsive to the question. It exceeds the scope of this witness's purported expertise. And it lacks foundation.
- MR. WISNER: He literally published an article about it.
 - THE COURT: Hold on just one second.

 (Pause in the proceedings.)
- THE COURT: I'm going to strike "because scientists outside of the industry," from that point forward is stricken.

MR. WISNER: Okay.

Q. So in this article -- we're going to get to your article later. I just wanted to know generally what it was about. That's fine.

Let's move on to some other articles in here.

I don't want to know what they're about, just very
general statements about what they're about; okay?

A. Okay.

Q. So we have here an article, "How the U.S. EPA and IARC reached diametrically opposed conclusions on genotoxicity of glyphosate-based herbicides."

Do you see that?

- A. Yes.
- Q. Was that also published in a peer-review journal?
 - A. Yes.
- Q. And it looks like we have discussions about -the next one about the track and control of pesticide
 risks. Do you see that?
 - A. Yes.
- Q. And then there's -- I mean, Doctor, I see a lot of articles here about pesticides. I guess my question is to you: How many times have you published in peer-reviewed scientific journals about pesticides?
 - A. Maybe 20 papers, something like that.

- Q. And were some of those papers looking at the toxicological profile of pesticides?
- A. Oh, most of them. And also the use, how many pounds were applied. And I've also published a number of papers that get at the impact of policy on pest management systems, which indirectly then impacts pesticide use.
- Q. Now, this is something that I think -- I got to clear up.

So you're talking about how policy and science sort of relate; is that right?

A. Yes.

- Q. To be able to talk about that relationship, do you have to be able to understand the science?
 - A. Well, it certainly helps.
- Q. What happens if you don't understand the science and start talking about how it applies in policy?
- A. Well, for one thing, you can make a fool of yourself pretty quickly because, you know, the science is absolutely integral to, for example, the regulatory process. The regulatory process, it's really managing of the evolution of scientific knowledge that relates to pesticide impacts and pesticide risk.
 - Q. All right. Doctor, so I want to talk about

Roundup and glyphosate. Okay? And I want to start from the beginning.

Let's start -- I kind of want to go back in time here and start at the beginning of the story of glyphosate in Roundup. Okay.

So my first question is: When was -- when was glyphosate first discovered?

- A. In 1950 by a small Swiss pharmaceutical company, I believe it was Cilag, C-I-L-A-G. They were looking for a new drug and were unable to identify any medicinal applications of it. And so it -- they really didn't do anything with it.
 - Q. That was in the 1950s; right?
 - **A.** 1950, correct.

2.1

- Q. Then what happened next?
- A. The Cilag sold a number of --

MR. BROWN: Excuse me, Your Honor. I'm going to object. It's all hearsay. And there's no foundation.

THE COURT: Overruled. He can answer.

THE WITNESS: Cilag sold a number of molecules to a company called Aldrich which was another chemical company. And they tried to develop and find uses for various molecules. But they also did not recognize any valuable commercial use for glyphosate.

So it was sort of -- it was on the shelf by 1 2 this Aldrich company, but it was not being actively 3 developed and nor had its remarkable properties as a herbicide been recognized yet. 4 BY MR. WISNER: 5 Well, what happened next? Was it ever used --6 Q. did it have any use before being recognized as an 7 herbicide? 9 No. No commercial uses to my knowledge. 10 Q. Okay. Was it patented in 1961 for use in cleaning industrial boilers? 11 12 THE COURT: So, counsel, I do want you to lay a foundation. 13 MR. WISNER: Sure 14 15 Are you familiar with whether or not 16 glyphosate has ever been used as a descaling agent? 17 It's been -- it has been explored for that Α.

A. It's been -- it has been explored for that use, and I think it did have some uses and there was a patent around 1960 for that particular use.

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- Q. Okay. So the 1960s, patented as descaler; is that right?
- A. Yeah, that would be a correct term to describe it, descaling agent.
 - Q. So when did it become an herbicide?
 - A. So Aldrich sold a suite of molecules, I think

1	there were over 100, to Monsanto. And at the time	
2	MR. BROWN: Excuse me, Your Honor. The answer	
3	is nonresponsive.	
4	MR. WISNER: Actually it is.	
5	THE COURT: Why don't you just approach for a	
6	second.	
7	(Sidebar held but not reported.)	
8	MR. WISNER: May I proceed, Your Honor?	
9	THE COURT: You may proceed.	
10	BY MR. WISNER:	
11	Q. Sir, have you researched the history and	
12	origins of glyphosate in the U.S.?	
13	A. Yes.	
14	Q. Have you researched the patents at issue?	
15	A. Many of them.	
16	Q. Okay. So you were describing to the jury a	
17	second ago how it went from being patented as a	
18	descaling agent to an herbicide. Can you please finish	
19	your answer.	
20	A. So what	
21	MR. BROWN: Your Honor, I hate to object. But	
22	there's no foundation. And it calls for hearsay.	
23	THE COURT: I'm going to overrule and let him	
24	answer the question.	
25	Go ahead.	

THE WITNESS: So Monsanto purchased a set of molecules. They were actually also looking for descaling agents. It was an active area of their research. And a scientist that worked for Monsanto at the time named Dr. Franz decided to take it into the greenhouse and see if it had any activity on weeds, which was a common thing for a scientist working in the industry at the time to just see if by any chance it had herbicidal activity.

And, lo and behold, Dr. Franz discovered the potent effectiveness of the glyphosate molecule to kill plants.

BY MR. WISNER:

- **O.** And when was that?
- **A.** 1970.
- Q. Okay. All right. I wrote glyphosate becomes herbicide. Maybe I should say --
 - A. Recognized as a herbicide.
 - Q. Yeah, I realize.

All right. Following 1970, what's regulatory steps -- well, back up.

Have you investigated the early regulatory history of glyphosate and its registration with the EPA?

- A. Yes, I have.
- Q. All right. Before we get to that, when did

the EPA actually come into existence?

A. In 1970.

- **Q.** When in 1970?
- A. I don't remember the month.
- Q. Okay. So in the same year that glyphosate was recognized as an herbicide, the EPA was actually created?
 - A. Correct.
 - Q. All right.

And at what point did the EPA have an authority to specifically regulate pesticides?

- A. Well, certainly by 1971 they were in control of the pesticide regulatory process, which actually had just been moved en masse from the USDA into the newly formed EPA.
- Q. Okay. So between 1970 and 1970 -- well, when was Roundup -- glyphosate approved?
- A. The first regulatory actions approved experimental uses of glyphosate-based herbicide -- it hadn't been named Roundup yet -- I think cotton was one of the first and soybeans was one of the first. These are very limited applications involving only usually a few acres of crop that are done to, first of all, see if it works on the field scale, but to also help the company understand how to formulate it, what the

application rate ought to be, how it needs to be applied.

So an experimental use permit, or EUP, is always the first action, followed very soon thereafter, and I believe the first tolerance petitions from Monsanto to EPA to sanction residues of glyphosate in the crops that it's been sprayed on, were submitted to the agency in late 1971 or '72.

- **Q.** Okay. So in 1971-72, it had this experimental application?
 - A. Right.

- Q. What I'm interested in is when did it become Roundup as we know it today? Or get approved as Roundup as we know it today?
- A. Well, the first commercial sales of Roundup were in 1974.
- Q. Okay. So by 1974 Roundup is approved; is that right?
 - A. Yeah, named and approved. Yep.
- Q. Okay. And when you say Roundup, that includes both glyphosate, the molecule we've been talking about, as well as the other ingredients?
 - A. The other surfactants, yes, correct.
- Q. All right. So between 1970-1974 so between the discovery of glyphosate as an herbicide and Roundup

being approved, were there any scientific studies done on glyphosate to get approval?

A. Oh, yes, definitely.

2.

- Q. And what are some of the studies that are required before an herbicide like Roundup can be approved?
- A. So in order for when Monsanto submitted to the EPA what's called a tolerance petition, it's a request to the EPA to establish a legal limit of glyphosate in soybeans or glyphosate in corn, they had a set of toxicological data requirements that they had to fulfill.

Two long-term animal studies, which you've heard a lot about, the mouse studies and the rat studies. There would be studies on teratogenicity which would look at birth defects, there would be mutagenicity studies. There would be a large number of metabolism studies in different animals to try to understand when a person is exposed to glyphosate, how it moves through their body.

And there's also a large number of studies in plant metabolism to help EPA figure out where to set that tolerance. You know, could there be one part per million in the soybean when it's harvested or five parts per million.

So between 1970 and 1974, there were probably a few hundred studies submitted by Monsanto to the newly formed EPA to set the regulatory foundation for the commercial uses of Roundup that started in 1974.

- Q. Now, you've mentioned this concept of tolerances a couple of times. Can you just explain to the jury what that is.
- A. So for all uses of a pesticide on a food crop, apples, oranges, spinach, soybeans, corn, there has to be established, before the EPA will approve a label making it legal to say spray Roundup on a soybean field, there has to be a tolerance covering the residues of glyphosate that are going to remain on the soybean after the farmer harvests the crop.

And those tolerances are set at a level sufficient to cover the residues measured in soybeans based on the way the label allows the pesticide to be used.

So they're a -- they're both a way to control the dietary risks from pesticides in food, and they're a way to enforce compliance with the label directions.

Because the body of data that Monsanto developed and submitted to the EPA showed the maximum amount of glyphosate that would remain on soybeans if the product was legally applied.

So if there was ever higher residues, that would be evidence that an illegal application had been made.

- Q. Okay. So going between 1970-1974 where these studies are being done to support the registration for Roundup, did the EPA itself conduct any of those studies?
 - A. No.

2.1

- Q. Well, who does conduct the studies?
- A. The registrants either conduct them in their own in-house laboratories by scientists that work for them. Or more typically they would contract with outside commercial testing laboratories that are set up to meet the needs of the drug industry, the pesticide industry, the oil and gas industry for testing of chemicals.
- Q. Between 1970-1974, were there long-term animal cancer studies done with glyphosate?
 - A. Yes.
 - Q. Who conducted those studies?
- A. A contract lab in Northbrook, Illinois called Industrial Bio-Test Laboratories, IBT.
- Q. Are you familiar with the events surrounding the IBT laboratory?
 - A. Oh, yes.

- Q. Why are you so familiar with them?
- A. So, remember, from in 1981, '82, '83, I was the staff director of this congressional subcommittee that had jurisdiction and responsibility to track what's going on with pesticides.

Right at that time, the events involving this IBT Lab producing fraudulent data in support of the registration of a bunch of pesticides broke into, you know, public view. And the EPA was in a terrible position. The companies were in a bad position. The public was frustrated that there was no valid data supporting the use of a couple hundred pesticides.

It was probably the largest sort of scandal, if you will, in the history of pesticide regulation in the U.S.

- Q. And I want to talk about the IBT story a little bit. I understand a report was prepared by the EPA in 1983.
 - A. Yes.

- Q. Are you familiar with that report?
- A. It was done at our request and submitted to the subcommittee.
- Q. Okay. If you look into your binder, is Exhibit 1364 a fair and accurate copy of that report?

1 That's the one. Α. 2 MR. WISNER: Permission to publish, 3 Your Honor. It's 1364. 4 MR. BROWN: Objection, Your Honor. Exceeds 5 the scope. And also not relevant in terms of the order 6 that the Court has previously made. 7 THE COURT: Overruled. You can publish. 9 (Exhibit published.) BY MR. WISNER: 10 All right. Doctor, looking here at this 11 Q. report, and I'll call out the front, this is the Summary 12 of the IBT Review Program, Office of Pesticide Programs, 13 July 1983. Do you see that? 14 15 Α. Correct. And what was going on in 1983 with regards to 16 17 this IBT review program? Oh, there was a major federal trial in 18 Α. Chicago --19 20 Q. Oh, please don't talk about that. 21 Α. Oh. I just want to talk about what this report is 22 Q. 23 about. Okay. This report identifies all of the 24 Α. toxicology studies supporting currently registered, 25 3521 currently used pesticides that rested upon IBT studies and then identified which of those IBT studies had been deemed to be fraudulent or invalid.

Q. Okay. So if we go into this report, and we go specifically to page 19 of the report.

Well, actually, let's start off with the intro. It's bottom right page 12; do you see that?

- **A.** Page 12?
- Q. Yeah.

2.1

- A. Yes, I'm there.
- Q. First paragraph talks about, Exhibit A, quantitatively presents the database of the chemical compounds for which studies were conducted by Industrial Bio-Test Laboratories. The IBT studies are designated by the letter O, studies in the EPA database done by labs other than IBT are designated by the letter X.

Do you see that?

- A. Yes.
- Q. Okay. And if you turn to page 19, we have the data here specifically for glyphosate. Do you see that?
 - A. I do.
- Q. All right. Well, I'll just keep it so everyone can see it.

And under here we have this category "Oncogenicity." Do you see that?

Α. Yes. 1 2 What is oncogenicity? 3 Α. It's the study of the capacity of a chemical to cause cancer. 4 Okay. And if we go down to glyphosate, right 5 Q. 6 here, we have one zero. Do you see that? Α. Yes. 7 What does that one zero mean? 0. 9 That is one IBT study had been submitted that Α. 10 had been done at IBT on glyphosate. And it was specifically about whether or not 11 Q. it causes cancer? 12 13 Α. Correct. All right. If we look over, there's another 14 Q. 15 that says mutagenicity. Do you see that? Α. Yes. 16 17 And if we go down to mutagenicity, it looks Q. like there's one X and four O's? 18 Correct. 19 Α. What does that mean? 20 Q. That means that at that point, the EPA had 21 Α. received from Monsanto five mutagenicity studies, four 22 of which came from IBT and one of which came from a 23 different lab. 24

All right. And if you look at the far right,

25

Q.

it has this phrase "data call-in." Do you see that?

A. Yes.

- Q. What does "data call-in" mean?
- A. "Data call-in" is the term of art in the EPA regulatory program where it requires a company to submit a new study. It may be a new study to support a new type of pesticide or a new application of a pesticide.

But in this case, it's a data call-in to replace studies that had been deemed to be invalid.

- Q. All right. If we go to Exhibit B, which is the next portion of this report, we're on page 30 if you want to follow by paper, there's a section here defining some terms. Do you see that?
 - A. Yeah.
- Q. And it says, "Review designed to determine if the information in the final report was supported by the raw data from the study." Do you see that?
 - A. Yes.
 - Q. What is that referring to?
- A. These are the criteria or the classifications in which EPA placed the different studies that had been submitted to it from the IBT lab. It was sort of their effort to determine which ones were invalid, which ones were poor quality but acceptable, and which ones were fine.

- Q. Now it says right here "was supported by the raw data." Is it your understanding that the EPA collected the raw data from the study?
- A. They visited the lab and asked to see the records, yes.
- Q. And then they compared those records to what the results were?
- A. The results as reported in the physical report that had been submitted to the agency by Monsanto.

 Monsanto having received it from IBT.
- Q. Okay. In the first category here, it says, "Invalid. The information in the final report was not supported by the raw data from the study."

What does that mean?

- A. It's sort of self-explanatory. The EPA went in and investigated the raw data as stored in the files at the laboratory, and it didn't support the data that appeared in the report. So they would have no basis to know what the study showed or even if there was a study.
- Q. In the original IBT cancer study that was submitted in support of registration, did that study report no oncogenic effect?
 - A. That's what it reported, yes.
- Q. So let's go and see what they said about glyphosate. This is on page 37. And as you can see

1 right here, there is a bunch of studies involving glyphosate by IBT. Do you see that? 2 3 Α. Yes. All right. And now if we go to the one that 4 0. says carcinogenicity. 5 Yes, I see it. 6 Α. You see it? I'll highlight it for the jury. 7 Q. Do you see that? 9 Yes. Α. All right. What letter did the EPA give it? 10 Q. "I." 11 Α. What does that mean? 12 Q. Invalid. 13 Α. 14 So the EPA reviewed the raw data and concluded Q. 15 this study, the only study that supported glyphosate, 16 was actually invalid? 17 Α. The only two-year cancer study supporting 18 glyphosate, yes. 19 All right. And if we go into here, there's Q. also discussions about mutagenicity. Do you see that? 20 21 Α. Yes. So, for example, we have this mutagenicity 22 Q. 23 mouse study. Do you see that? 24 Α. Yes. And, again, what letter did the EPA give it? 25 Q.

1 Α. "I" for invalid. 2 All right. Let's find the rest of them. Here's another one, mutagenicity mouse. 3 you see that? 4 Α. Yes. There should be two more. 5 6 Okay. So it looks like -- then there's Ames. Q. Here we go? 7 The Ames study is a mutagenicity study. 8 Α. So of those four IBT studies that we saw in 9 ٥. the earlier of the document, how many of them were 10 deemed invalid, meaning not supported by the raw data? 11 All of them. Α. 12 Okay. Do you know when the IBT scandal first 13 Q. broke, when people first learned about it? 14 In 1976. 15 Α. 16 Okay. If you go to the beginning of this Q. 17 document -- well, not the beginning. Sorry. On page 9 it says right here: 18 19 1976, during a routine lab inspection of one of IBT's facilities, FDA discovered 20 deficiencies in the manner in which 21 studies were being conducted and 22 23 discrepancies between those studies and their raw data. 24

25

Α.

Yes.

- Q. Have you investigated actually what was discovered in 1976?
 - A. Oh, yes.

- Q. And you read the reports at the time?
- A. I actually spoke to the scientist that did the inspection.
- Q. And what did you learn about what was happening at the IBT Laboratories during this time?
 - A. It was a mess.
 - Q. Please describe.
- A. They had grown very fast. They were founded in the early 1950s, and by the early 1970s they had hundreds of studies underway and were trying to take care of 15- to 20,000 animals and -- you know, rats and mice in the cages.

And they had installed what at the time was a modern watering system that would automatically provide water to the rodents in their cages and also help clean out the feces. And it never worked right and it got plugged up. And it sprayed water where it wasn't supposed to go.

At one point there was a couple inches of water on the floor of the lab. Some animals were getting no water and died of thirst. Other animals were in water in the bottom of their cages --

1 MR. BROWN: Excuse me, Your Honor. 2 **THE WITNESS:** -- and it led to a very high 3 rate of --MR. BROWN: Excuse me, Your Honor. I'm going 4 to object. This is hearsay. It lacks foundation. 5 6 THE COURT: I'm going to sustain any objection with respect to IBT's internal issues. The fraudulent 7 data issue is relevant. 8 9 MR. WISNER: Fair enough. 10 Q. Putting aside the watering thing and what happened at IBT, I quess my question, the bottom-line 11 question is: Was it concluded, sir, that IBT had 12 engaged in scientific fraud? 13 14 Α. Yes. And when it was discovered in 1976 and the 15 years following that these studies were fraudulent, did 16 17 Monsanto remove Roundup from the market? 18 Α. No. 19 Q. Did they ever repeat the mouse study? 20 Α. Yes. Okay. And when did they repeat that mouse 2.1 Q. study? 22 23 It was begun in I believe 1981 at a lab called Α. Biodynamics. 24 And when did they finally have the results of 25 Q.

1 that study? 2 Α. 1983. So it wasn't until 1983 that Monsanto then had 3 Q. a valid mouse study assessing the carcinogenicity of 4 Roundup -- of glyphosate? Sorry. 5 That's correct. 6 Α. All right. So between 1976, and we're up 7 Q. to -- where's my markers? 1970 -- sorry, 1983. that's approximately how many years? 9 10 Α. Seven. Okay. So for seven years they didn't have any 11 Q. valid data, but was it still being sold the whole time? 12 Objection. Argumentative. 13 MR. BROWN: THE COURT: Overruled. I think he established 14 15 that already. THE WITNESS: Yes, it was still being sold. 16 17 BY MR. WISNER: Okay. And I guess my other question, and this 18 Q. 19 is kind of important, between 1976 and 1983, did 20 Monsanto ever warn about their lacking cancer data on their label? 21 22 Α. No. 23 Objection. The Court has ruled in MR. BROWN: terms of it exceeds the scope in terms of the Sargon 24 ruling. 25

1 THE COURT: Approach. 2 (Sidebar held but not reported.) 3 MR. WISNER: May I proceed, Your Honor? THE COURT: 4 Yes. BY MR. WISNER: 5 Dr. Benbrook, during this time period between 6 Q. 1970 -- sorry -- 1976 when the IBT fraud was discovered 7 and 1983 when they now had a new mouse study, did Monsanto ever disclose on its label that they had no 9 valid cancer data? 10 11 Α. No. Now, this new study, what was it called? 12 Q. 13 Α. I remember it as the Biodynamics 1983 mouse 14 study. 15 Okay. Now the jury has seen this before. 16 This is the tumor charts that Dr. Portier put together. 17 And just for clarity's sake, when you say Biodynamics, are you referring to the Knezevich and Hogan study from 18 19 1983? 20 Α. Yes, sir. Okay. I guess I actually have a follow-up 21 Q. question about this IBT~Labs story. You understand that 22 23 certain scientists were implicated in this fraud; is that right? 24

25

Α.

Yes.

- Q. One was by the name of Paul Wright?

 A. Correct.
 - Q. And the other one was by the name of Dr. Keplinger?
 - A. Correct.
 - Q. Did Dr. Keplinger sign off on this cancer study?
 - A. Yes.

- Q. So in 1983, we have the -- I'll call it Biodynamics because that's what you like -- study involving mice; is that right?
- A. Correct, yes.
- Q. And that was submitted to the EPA in what year, sir?
 - A. I think it was in the fall of 1983.
 - Q. All right. And actually I keep forgetting stuff I want to cover.

Before we get to that study, I want to talk about it briefly, but after the IBT scandal was kind of learned about 1976, did the EPA have the authority to just force the withdrawal of these pesticides off the market?

- A. Actually they did not.
- Q. So if we look at this document on page 4, it says right here --

- A. That's page 3 -- oh, bottom of 3, okay.
- Q. Yeah, it's confusing. There's like 10 numbers on everything.

But it says:

2.1

The IBT case caused serious concern and uncertainty about the potential hazards of the hundreds of pesticides involved, both for EPA and the public. Although it was advocated by some that all 212 pesticides tested in whole or in part by IBT be removed from the market pending retesting, that option was not available under current law.

Do you see that?

- A. That's correct.
- Q. Can you please explain to the jury why that option was not available under current law.
- A. The basic FIFRA statute, this is the federal statute that lays out the rules that EPA has to follow to cancel an existing registration of a pesticide, states that the EPA must conclude based on valid evidence that the risks associated with the ongoing use of the pesticide exceed the benefits to the farmers from the use of the pesticide. That's the basic standard embedded in the statute.

But lacking a valid study, they had no way to 1 2 quantify the risks. And so they knew they would be 3 unsuccessful if challenged in court because they had no basis to even say there might be any risk. 4 So because of that, they had no legal 5 6 mechanism to drive the products off the market. Who, other than -- so the EPA didn't have the 7 Q. ability. Who had the ability to take it off the market 9 pending test? 10 Α. The registrants. And that was who? 11 Q. 12 Α. Monsanto. 13 Q. All right. So we're now up to 1983. this Biodynamics study. By when did the EPA review this 14 15 study? I'm sorry, I didn't hear you, Brent. 16 Α. 17 Q. Sorry. By when did the EPA review the study that was submitted to it? 18 MR. BROWN: Calls for speculation. 19 It --20 THE WITNESS: MR. BROWN: Lacks foundation. 21 THE COURT: Overruled. 22 23 The EPA review of the THE WITNESS: Biodynamics study was -- it went through many phases, of 24

course, but in 1984.

1 BY MR. WISNER: 2 Okay. I want to look at a document that's 3 actually already in evidence. If you turn to your binder, Exhibit 868, is 4 that one of the original memos discussing the EPA's 5 assessment of that study? 6 Yes, it is. 7 Α. MR. WISNER: Permission to publish, 9 Your Honor? THE COURT: It's in evidence? 10 MR. WISNER: Yes. 11 12 THE COURT: Granted. 13 (Exhibit published.) 14 BY MR. WISNER: All right. So we're looking at this memo. 15 16 The jury actually saw this in the context of a 17 deposition. But I want to walk you through some of it just to make sure I understand some of the technical 18 19 aspects of it. Okay? So it's dated here April 3rd, 1985. Do you 20 see that? 21 22 Α. Yes. 23 And it's from William Dykstra, Ph.D., Q. Toxicology Branch. Do you see that? 24 Yes, sir. 25 Α.

- Q. What was the Toxicology Branch back in 1985?
- A. That was the part of the Office of Pesticide
 Programs which was the part of the EPA responsible for
 pesticide regulation. The Office of Pesticide Programs
 had several branches, and they took on different aspects
 of the scientific review. The Toxicology Branch
 evaluated all of the cancer studies, the mutagenicity
 studies, the birth defect studies on pesticides.
 - It says right here under conclusions:

 Glyphosate was oncogenic in male mice causing renal tubule adenomas, a rare tumor, in a dose-related manner. The study is acceptable as core minium data.

 Do you see that?
 - A. Yes.

Q.

- Q. All right. We're going to break down that sentence. It says glyphosate was oncogenic in male mice, causing these tumors. What does that mean?
- A. It means that there was a statistically significant increase in renal tubular adenomas in the male mice that were administered glyphosate in their feed compared to the control group of male mice that did not receive any glyphosate in their feed.
- Q. What does it mean when it says here "core minium data"; what does that mean?

- A. It means that the study was deemed to be valid and that it satisfied one of the data requirements supporting both glyphosate registrations and also the tolerances covering glyphosate residues in food.
- Q. Now if we go into the study a little bit farther to actually look at what the study showed, if we go to on page 2 of this document, it talks about the renal tubular adenomas that occurred in male mice, do you see that?
 - A. Yes.

2.

- Q. Briefly explain to the jury what's the significance of this zero, zero, one, three.
- A. So that's the core result of the Biodynamics mouse study in the males as reported to EPA, both in the Monsanto summary of the -- of the Biodynamics study, as well as the Biodynamics study itself, which of course was submitted in full to the EPA.
- Q. All right. So we go to the front page again. It says this information oncogenicity of glyphosate was evaluated by the Toxicology Branch AD ad hoc committee which concluded that this was an oncogenic response.

What is a Toxicology Branch ad hoc committee?

A. It's a team of the scientists working within the Toxicology Branch that span the range of expertise required to evaluate cancer studies. So it would be

1	statisticians, pathologists, toxicologists.
2	Q. Okay. And have you seen a copy of the
3	consensus report mentioned here?
4	A. Yes.
5	Q. Is that Exhibit 875 in your binder?
6	THE COURT: So we're going to take a break
7	this morning for about 10 minutes.
8	MR. WISNER: Sure.
9	THE COURT: Is this a good time?
LO	MR. WISNER: Right after this document, it's
L1	perfect.
L2	THE COURT: Okay.
L3	THE WITNESS: Yes, the 875 is the consensus
L4	report of this review committee.
L5	MR. WISNER: Your Honor, it's in evidence.
L6	I'm going to publish.
L7	(Exhibit published.)
L8	BY MR. WISNER:
L9	Q. So we're looking here at Exhibit 875. And we
20	have here a listing of all these different scientists.
21	Do you see that?
22	A. Correct.
23	Q. One of them, of course, is Dr. Dykstra from an
24	earlier memo.
25	A. Yes.

Q. We have a statistician, Herbert Lacayo. 1 2 you see that? 3 Α. Yes. All right. And then if we go to the final 4 0. conclusion here, it states: 5 Classification of glyphosate. 6 In accordance with EPA proposed quidelines, 7 the panel has classified glyphosate as a category C oncogen. 9 Correct. 10 Α. What is a category C oncogen? 11 It is a chemical that is considered to 12 Α. possibly pose cancer risks to animals including humans. 13 And is there a significance in a regulatory 14 capacity of a finding as a class C oncogen? 15 16 Α. Oh, yes. 17 MR. WISNER: Great. Let's take a break, Your Honor. We can talk about that after the break. 18 THE COURT: We're going to take 10 minutes 19 because we will be breaking for lunch so we'll take a 20 short break. 21 (Proceedings continued in open court out of 22 23 the presence of the jury:) 24 THE COURT: You can step down, Doctor, 10 minutes. 25

1	MR. WISNER: Yes, Your Honor.
2	THE COURT: In terms of just timing, how much
3	longer?
4	MR. WISNER: I was planning to go till about
5	2:00 o'clock today.
6	THE COURT: 2:00?
7	MR. WISNER: Yeah, but that was with the
8	understanding that we'd start at 9:00. So we didn't
9	start until almost 10:00. So I'll probably go to 2:30,
10	2:40.
11	THE COURT: Okay. So then will he be done
12	today?
13	MR. BROWN: Yes.
14	THE COURT: I'm just trying to get an idea of
15	the schedule.
16	MR. WISNER: He's available to come tomorrow
17	if we have to.
18	(Recess taken at 10:45 a.m.)
19	(Proceedings resumed in open court in the
20	presence of the jury at 10:59 a.m.)
21	THE COURT: You may proceed, Mr. Wisner.
22	MR. WISNER: Yes, Your Honor.
23	Q. All right. Doctor, just before the break, we
24	were talking the category C classification.
25	Now, earlier we talked about this concept of

tolerances.

2.

So going back in time, back to 1985 when this memo was written, what, if any, was the significance of a category C carcinogen on the ability to sell it and market Roundup?

- A. You mean EPA classifying glyphosate as a possible human carcinogen?
 - Q. That's right.
- A. It had very significant regulatory implications because that would prohibit the EPA from establishing higher tolerances that would be needed to expand the use of Roundup on a wide range of crops. So it had a very direct effect on the market potential for future Roundup sales.
- Q. Now, you said -- are you familiar with something called the Delaney Clause?
 - A. Yes.
 - Q. How are you familiar with it, sir?
- A. Probably -- the impact of the Delaney Clause on pesticide regulation and in particular the regulation of cancer-causing pesticides has been probably the public policy issue I've spent the most time on in my entire career.
 - Q. Have you written books about it?
 - **A.** I've written books and reports and articles,

yes.

- Q. Have you actually testified before Congress about this issue?
 - A. Multiple times.
- Q. Okay. So walk the jury through what the Delaney Clause is and specifically how a category C classification is implicated.
- A. Okay. The Delaney Clause was added to the Food, Drug, and Cosmetic Act, which is another major federal statute that governs things like food additives. So food additives, like a coloring agent or a preservative that General Mills would put in a cereal or whatever.

The Delaney Clause is really a very simple clause. It says thou shalt not add a known carcinogen into food as a food additive.

So it was passed into the -- the Delaney
Clause amended Section 409 of the Food, Drug, and
Cosmetic Act. Section 409 of the Food, Drug, and
Cosmetic Act is the authority under which a certain
class of pesticide tolerances are established, and in
particular, tolerances for residues in processed foods.
Foods other than -- when an apple is harvested off of a
tree or a grape taken, that's a fresh raw food in its
fresh form.

And, yes, there are tolerances covering the residues of pesticides that might have been legally applied on the apple tree, on the apple and on the grape. But, remember, a lot of us eat grapes as raisins. So if you -- what happens between a fresh grape and a raisin, you take the water out. So it's much lighter, but the pesticides stay in it.

So if you were to measure the level of a pesticide in a raisin compared to the grape, it will be much higher. And to cover those higher residues, a food additive tolerance needed to be established under Section 409 of the Food, Drug, and Cosmetic Act. But that provision was -- had as part of it the Delaney Clause.

So if there was evidence of possible oncogenic effect, the EPA would be blocked, it would be illegal to establish those tolerances.

So this was the significance of the EPA's judgment that this 1983 mouse study showed that glyphosate, you know, was a possible carcinogen, it really would block the approval of the next batch of tolerances that Monsanto had to get on the books in order to allow the use of Roundup to expand.

Q. All right. So they categorized it as a class C, and shortly after that the EPA and Monsanto had

a conversation; is that right? 1 2 Α. Yes. 3 Q. And you know about this because -- have you seen this in a memo before? 4 Oh, there's extensive documentation in the 5 Α. 6 record of the case of what happened actually the very day that Monsanto found out that Dykstra's review was 7 going to --9 MR. BROWN: Excuse me. THE WITNESS: -- find that --10 Excuse me, Your Honor. There's no 11 MR. BROWN: question pending. This is speculation. 12 Lacks foundation. Calls for speculation. 13 THE COURT: The question was: And you know 14 15 this because you've seen this in a memo. So that's the question. 16 17 MR. WISNER: Okay. Doctor, I want to draw your attention to an 18 Q. exhibit that's actually in evidence, Exhibit 73 in your 19 binder. 20 Do you have it, sir? 21 Yes, sir. 22 Α. 23 It's also on the screen if you want to use Q. that as well, whatever you prefer. 24 Now, is this one of the memos that you're 25

1	referring to?
2	A. Yes.
3	Q. And this is dated February 22nd, 1985. So
4	this is approximately 10 days after the consensus
5	statement by the EPA?
6	A. Correct.
7	Q. All right. And if we go down here, we see a
8	bunch of individuals who are present. Do you see that,
9	sir?
10	A. Yes.
11	Q. And some of them have names, for example: Ted
12	Farber, a branch chief. Do you see that?
13	A. Correct.
14	Q. And above that it has Bill Burnam, and it says
15	Assistant Chief OPP Toxicology Branch?
16	A. Correct.
17	Q. Who are these individuals within the
18	regulatory context?
19	A. These are the top people in the Tox Branch
20	that were responsible for evaluating cancer studies like
21	this 1983 Biodynamics mouse study.
22	Q. And these three individuals right here, are
23	they with Monsanto?
24	A. They were the
25	MR. BROWN: Excuse me. Lacks foundation.

1	Calls for speculation. And hearsay.
2	THE COURT: Sustained.
3	BY MR. WISNER:
4	Q. Do you see the name Lyle Gingerich there?
5	A. Yes.
6	Q. And this is a memo by Monsanto; right?
7	A. Correct.
8	Q. Let's look and see who signed this document.
9	Who signed it, sir?
LO	A. This memo was prepared by
L1	MR. BROWN: Excuse me, Your Honor. I'm going
L2	to object. The document speaks for itself. The
L3	question is argumentative.
L4	THE COURT: Sustained.
L5	MR. WISNER: The question is who signed it? I
L6	can't ask that question?
L7	THE COURT: It speaks for itself.
L8	MR. WISNER: Okay.
L9	Q. Was the document signed by Monsanto's
20	employee?
21	A. Yes.
22	Q. When we go into this document, there is a
23	couple of things that I want to ask you about, about
24	what they mean.
25	So we see here: Concerns of the Toxicology

1 Branch. Do you see that? 2. Yes. Α. 3 Q. And it says right here: Oncogenic in mouse, IARC ranking C. Do you see that? 4 Α. Yes. 5 Do you know what an IARC ranking C is back in 6 Q. 1985? 7 Possible human oncogen. Α. And it says down here: Biologically 9 Q. significant rare tumors. Do you see that? 10 11 Yes. Α. And statistically significant at the .5 level. 12 0. Do you see that? 13 14 Α. Correct. Do you know what that's referring to? 15 16 Yes. The -- remember from the table that Α. 17 Mr. Wisner showed before, there was zero, zero, one, three renal tubular adenomas in the male mice. 18 19 It's that trend that was statistically 20 significant, showing that there was a response to the feeding of glyphosate. 21 So I'm actually going to go a few pages ahead. 22 Q. 23 For example, I just want to visually depict this, sir. 24 So if we have a -- kind of plot out the four 25 groups. Okay.

1	We have the control, low, mid, and high dose;
2	right?
3	A. Correct.
4	Q. And here there's no tumors?
5	A. Zero.
6	Q. Here there's no tumors?
7	A. Zero, yep.
8	Q. How many are in the middle dose?
9	A. One.
LO	Q. So we'll put one there. And how many in the
L1	high dose?
L2	A. Three. You put a zero there.
L3	Q. I'll do that as a tumor.
L 4	So here we have three; right?
L5	A. Yes, sir.
L6	Q. And if we draw a line through this, is that
L7	the slope you're talking about?
L8	A. Essentially, yes.
L9	Q. Okay. And then if you go into this document
20	here, it reads "FJ asked." Do you see that?
21	A. Yes.
22	Q. And if we go to the beginning of the document,
23	who here has the initials of FJ?
24	A. Fred Johannsen.
25	Q. Okay. And to the best of your knowledge, was
	3548

1	Fred Johannsen a Monsanto employee?
2	MR. BROWN: Objection. Lacks foundation.
3	THE COURT: Sustained.
4	BY MR. WISNER:
5	Q. Do you know if Fred Johannsen was a Monsanto
6	employee?
7	A. Yes.
8	Q. Was he a Monsanto employee?
9	A. Yes.
LO	Q. And it says here:
L1	Short of a new study or finding
L2	tumors in the control groups, what can we
L3	do to get this thing off group C.
L4	Do you see that?
L5	A. Yes.
L6	Q. All right. So finding tumors in the control
L7	group, how would that have any influence on this?
L8	A. If there was
L9	MR. BROWN: Excuse me. Calls for speculation.
20	Lacks foundation.
21	THE COURT: Overruled.
22	THE WITNESS: Given that EPA and exercising
23	its regulatory responsibilities has to statistically
24	analyze the occurrence of tumors in the control and the
25	three feeding groups. Whether there is a consistent

slope in that line is absolutely essential. It's what determines whether the study is positive for renal tubular adenomas in the male mice or negative.

So you can imagine if there was one tumor in the control group, that line shifts up and becomes probably an equivocal finding at that point.

- Q. So if we draw in this tumor, it would change the line more flat; is that right?
 - A. Correct.

- Q. Okay. Following -- well, actually earlier in this document, it said -- and I had that section blown up for a reason -- it says right here: We'll ask to resection tissues, consider crystal formations, et cetera. Do you see that?
 - A. Yes.
 - **O.** What does it mean to resection tissues?
- A. The Biodynamics scientists would pull the kidneys out of the deep freezer, thaw them out and then cut fresh slides from the kidney to provide a second reading of whether -- whether there are, in this case, renal tubular adenomas in the various kidneys from the control group, the low, medium, and high treatment group.
- Q. Have you reviewed the documentation related to the resection of this?

- A. Yes.
- Q. Did Monsanto hire anybody to relook at these kidney tumors?
 - A. Yes.
 - Q. Who did they hire?
- A. A Dr. Kuschner was the principal pathologist that they hired. But then there were several others that were also hired and asked to express their opinion about whether there was actually a tumor in the particular male control mouse that has been talked about, I'm sure, during the course of this trial.
- Q. And when Dr. Kuschner reviewed these tumor slides, did he discover this tumor in the control group?
 - A. He reported that he saw one.
 - Q. Okay. How do you spell Kuschner?
 - A. K-U-S-C-H-N-E-R, I believe.
- Q. And when he found that -- when he claims to have found that tumor, did it affect the slope?
 - A. Yes.
- Q. Now I want to back up a little bit because before that whole tumor thing occurred, there was a report issued by an EPA statistician. Are you familiar with that?
 - A. Yes, I am.
 - Q. Turn to Exhibit 1375 in your binder. Is that

1 that report? Yes, it is. 2 Α. 3 Q. Now, it's already in evidence so I'm going to put it up on the screen. 4 (Exhibit displayed.) 5 BY MR. WISNER: 6 And we have here a report, you see it's 7 written by Herbert Lacayo, a statistician. Do you see 9 that? 10 Α. Yes. 11 And if we go into what he's doing here, the Q. background, it's -- is it talking about this mouse 12 study? 13 14 Yes, it is. Α. 15 Q. It reads here: The registrant, Monsanto, claims that 16 17 such tumors are unrelated to treatment. In support of that, they provide 18 19 historical data from Biodynamics and two other laboratories. 20 Do you see that? 2.1 22 Yes. Α. 23 And then if you look down here, if you go to Q. the summary, there is some remarks by the statistician. 24 Do you see that? 25

1	A. Yes.
2	Q. Okay. The first thing I want to point out is
3	right here, and this is on page 3 where Dr. Lacayo has
4	done sort of a probability analysis. Do you see that?
5	A. Yes.
6	Q. And just don't get into too many details, we
7	don't need to get into it, but what generally was
8	Dr. Lacayo doing?
9	A. He was trying to assess
10	MR. BROWN: Excuse me, Your Honor. I'm going
11	to object. Calls for speculation.
12	MR. WISNER: Your Honor, I'm just trying to
13	explain the document and what it says.
14	THE COURT: Overruled. He can answer if he
15	knows.
16	MR. WISNER: Yeah.
17	Q. Sir, do you know what this analysis is
18	generally doing?
19	A. Yes, he's trying to quantify the odds that
20	there would be a tumor in a particular number of mice in
21	a control group.
22	Q. He writes:
23	Another way of saying this is that if
24	glyphosate were truly unrelated to kidney
25	production, we would expect to see four

1 more tumors in less than one out of 100 2 experiments of the type sponsored by 3 Monsanto. Do you see that? 4 Yes. 5 Α. 6 Can you explain what that means? Q. So, remember, in the chart there were four 7 Α. renal tubular adenomas identified by the Biodynamics pathologist that read the original slides. And what 9 Dr. Lacayo did was he calculated the odds of it in a 10 11 study with this number of mice in each of the groups, started out with 50 in each of the groups, what are the 12 odds of there being four tumors. And that's what this 13 14 table shows. 15 And he goes: Thus glyphosate is suspect. 16 you see that? 17 Correct. And with regard to suspect, is that because 18 19 the likelihood of seeing four tumors, these rare tumors, was astronomically small? 20 21 Α. Very small. Okay. And then he talks about something 22 Q. 23 called false positives. Do you see that?

What is a false positive?

24

25

Yes.

Α.

Q.

- A. A false positive is a study that appears to contain data suggestive of a biological response or of cancer when in fact there is no underlying impact.
- Q. Okay. And Dr. Lacayo goes on to discuss this concept of false positive. And I want to sort of look at this part right here. Well, look at the bottom part. It says:

Viewpoint is a key issue. Our viewpoint is one of protecting the public health when we see suspicious data. It is not our job to protect registrants from false positives. We sympathize with the registrant's problem, but they would have to demonstrate that this positive result is false.

Do you see that?

A. Correct.

- Q. And this is something that I think is important from a regulatory context. Does the EPA have to prove that it's a negative study or a nonpositive result?
 - A. No.
 - O. How does it work?
- A. When the registrants submit the various data requirements, and in this case a two-year mouse cancer

1 study, the EPA looks at the results of the study, and if 2. there is a statistically significant upward trend in the response of a particular tumor, and in this case these 3 renal tubular adenomas, then that is evidence that 4 there's a possible oncogenic response from exposure to 5 6 the pesticide or the chemical at question. And once EPA has one study that shows such a 7 response, it has typically been the position of the

agency that it will be regulated as a possible oncogen.

Now, it says right here the registrant will Q. have to demonstrate that this positive result is false.

Do you see that?

Α. Yes.

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- Would one way of demonstrating that it's false Q. be finding a tumor in the control group?
 - Α. Yes.

MR. BROWN: Objection. The question is argumentative.

THE COURT: Overruled. The answer will stand.

BY MR. WISNER:

- And if you look at the date on this, sir, at the very beginning, this is February of 1985; is that right?
 - Correct. Α.
 - Okay. When was Dr. Kuschner hired to look at Q.

these kidney tumors?

- A. I believe it was in April.
- Q. So this statement about demonstrating falseness, that was before Dr. Kuschner found -- or claims to have found this tumor?
 - A. Correct.
- Q. Okay. All right. Well, I don't want to go into the whole story. I just want to break this down because these documents were in evidence, and I wanted the jury to understand it.

But there is one sort of other issue that I want to clear up. Following this ordeal with this tumor, did the EPA ever order Monsanto to do a new mouse study?

- A. Yes.
- Q. Please explain to the jury what happened.
- A. Because of the --

MR. BROWN: Excuse me. The question is vaque.

THE COURT: Overruled. You can answer.

THE WITNESS: Because the pathologists in the EPA and working for Monsanto disagreed, they simply disagreed. The EPA people didn't see the additional renal tubular adenoma in that male mouse, and the Monsanto-hired pathologists all did. They were at an impasse.

And so whenever that happens in -- in the vast majority of cases when that happens in the case of assessing the risks of a particular pesticide, the agency asks for a new study and often a better study.

And that's what EPA did in a -- what's called a registration standard document that was issued in 1986. It did a data call-in for a new mouse study.

BY MR. WISNER:

- Q. All right. So going back to our timeline here, 1985, EPA classifies glyphosate as class C; is that right?
 - A. Correct.
- Q. And then there was this fight about the tumor.

 And by 1986 EPA orders new mouse study; is that right?
 - A. Correct.
- Q. Now, did they also order a new rat study as well?
 - A. Yes.
- Q. What happened to the -- well, let's pull up the board. But we have the rat studies here.
- And this study from 1981, that was a Monsanto study; is that right?
 - A. Correct.
- Q. What was wrong with this study? Why wasn't this enough?

Α. It was another IBT study, I believe. 1 2 No, this is 1981. 0. 3 Α. Oh, the 1981. Okay. So they asked for a new -- do you know why 4 0. they asked for another rat study? 5 I don't remember. 6 Α. That's fine. We don't need to get into it. 7 Q. But they ordered a new mouse study; right? 8 9 Yes. Α. And this chart here, the jury has seen it 10 Q. quite a bit, these are the mouse studies that we know 11 about related to glyphosate; right? 12 Correct. 13 Α. And that was the 1983 study; right? 14 Q. 15 Α. Yes. And actually just to clarify, Dr. Portier 16 Q. 17 identified a spleen composite lymphosarcoma which is a type of lymphoma in the Knezevich and Hogan study. 18 your review of the EPA documents, even back in 1983, did 19 20 they see that? 21 Α. Yes. Okay. So moving forward from 1983 onward, did 22 Q. 23 Monsanto conduct any of these studies? 24 Α. No. So after the EPA ordered Monsanto to conduct a 25 Q.

new mouse study to resolve this issue, did they ever do it?

A. No.

- Q. Did the EPA make any accommodations to Monsanto for doing that mouse study?
- A. EPA, Dr. Dykstra, who had been the EPA toxicologist most involved in this, actually interacted with Monsanto scientists to design a special study designed to definitively determine whether there was an increase in the renal tubular adenomas in male mice. It was kind of a mouse study on steroids in effect. He wanted 250 animals per group instead of 50, which would increase the statistical power of the study.

And the EPA called for Monsanto to only do the histopathology on the liver and the kidney to keep the costs down because they were asking for a lot of animals to be in the study. And if there was no evidence of any problems in the liver and kidney, then Monsanto wouldn't have to do any other diagnostic work.

So it was a big, very powerful -- statistically powerful study designed just to resolve the issue over renal tubular adenomas in male mice.

- Q. Did that study ever get done?
- A. No.
- Q. And every study afterwards, is it your

understanding that they found malignant lymphoma in mice exposed to glyphosate?

A. Yes.

Q. Going through this process from 1986, was -- well, let me back up.

Are pesticides just approved one time and they're good forever? Or is there a reregistration process with the EPA?

A. The regulatory status of pesticides evolves in two principal tracks. One is the company wants to get registrations to allow more crops to be treated or different circumstances where an application can be legal. The company might want to start selling the product into the home market for people like the Pilliods to use.

So as the diversity and number of uses of a currently registered pesticide expands, the company often is required by EPA to submit -- to develop and submit new data to support those uses.

The other track and circumstance in which EPA asks for new data and does a fresh assessment is through what's called the reregistration process. And one of the things when I was the -- back in the day when I was the staff director of the subcommittee with jurisdiction over the FIFRA statute, the nature of that

reregistration process, what had to be done with it, who paid for the data, was issues that were constantly under debate.

But so in the early 1980s when all of the debate was going on about these mouse studies, et cetera, glyphosate was in what's called the registration review process. And that provided EPA with sort of an opportunity to take a new look at all the science that was available at the time.

And in 1986 they issued what's called a registration standard, and that document set forth what EPA had to do in order to qualify for or get EPA approval of existing and new uses of glyphosate.

Q. When was --

MR. BROWN: Excuse me, Your Honor. I'm going to object and move to strike. The answer was nonresponsive.

THE COURT: Overruled. The answer stands.

BY MR. WISNER:

- Q. Now, Doctor, when was glyphosate -- so the original registration, right, that's back in 1974; right?
 - A. Correct.
 - Q. And that was based on the IBT fraud?
 - A. The IBT cancer study was the only two-year

cancer study available to the agency.

- Q. And four of the genotox studies; right?
- A. Right.

- Q. So the next time after 1974 that glyphosate is reregistered by the EPA is what year?
 - **A.** 1986.
 - Q. That's the quidance document; right?
- A. Well, that guidance document set out the terms and conditions for continuing the registrations that were currently on the books, but also to -- if EPA were to approve any new registrations for additional uses.
 - Q. But when did the final reregistration happen?
- A. Well, it would kind of depend on what product you're talking about because the actual approval of these different labels, they go through the EPA process on different time schedules.
- Q. Sure. Are you familiar with, you know, a 15-year cycle of reregistration?
- A. That is in the basic FIFRA statute as the goal, that every 15 years currently registered pesticides will be reassessed by the EPA and the underlying database supporting current uses will be brought up to current scientific standards.
 - Q. That's exactly what I'm talking about.
 So following 1974, when did that actually

happen next?

- A. It happened in the early '80s and led to the issuance in 1986 of this registration standard document.
- Q. Okay. Are you familiar with the 1993 RED document?
 - A. Correct. That would be the next cycle.
 - Q. Oh, so this is actually part of that cycle?
- A. No. They changed the name of this review process a couple of times. There's been registration standards. That was kind of in the '80s.
 - Q. Gotcha.
- A. And then they went to reregistration. And then they went to what's called a RED, which is a reregistration eligibility document, RED. Very confusing.
 - O. And that was in 1993?
- A. Yeah. By 1993, they called them -- I believe they called them a RED at that point. But that was the next reregistration cycle.
- Q. And in 1993 the only animal data that they had were the two rat studies and the mouse study; right?
 - A. Correct.
 - Q. And they didn't really have any epidemiology?
- A. Well, EPA never requires pesticide registrants to do epidemiological studies. And I don't think any of

1 the glyphosate non-Hodgkin's lymphoma epidemiological 2 studies had been published at that point. So that was in 1993. When did the next 3 Q. Okay. cycle for reregistration get set for? 4 I believe it started in 2005 or '6. Α. 5 6 Has it completed yet? Q. No. 7 Α. So the last official reregistration that's Ο. finalized was in 1993? 9 Correct. 10 Α. Before the epi? 11 Q. 12 Α. Correct. And if you look at the mouse boards, just look 13 Q. at the dates here, it was before -- well, actually it 14 was before Atkinson too; right? 15 16 Α. Correct. 17 Q. So all these studies were after? Correct. 18 Α. 19 And if we look at the mouse charts, all of Q. these malignant lymphoma findings were after? 20 21 Α. Correct. In the current status of the reregistration, I 22 Q. 23 want to talk a little about that. You said started 2005; is that right? 24

25

Α.

Around then, yes.

1 Q. Are you familiar with something called CARC? 2 Yes. Α. 3 What is the CARC? Q. It's the Cancer Assessment Review Committee, 4 Α. which is -- it's an internal committee within the Office 5 6 of Pesticide Programs composed of their senior 7 scientists with expertise in, again, biostatistics, pathology, toxicology, oncogenicity. 8 So if you turn to Exhibit 547 in your binder, 9 it's in evidence already, and we turn to the second 10 page, there's this e-mail exchange between Daniel 11 Jenkins and William Heydens and Jennifer Listello. 12 you see that? 13 14 Α. Yes. 15 It says down here: Q. Jess called me out of the blue this 16 17 morning. Do you see that? 18 19 Yes. Α. And this is from April of 2015; right? 20 Q. 2.1 Yes. Α. And this is -- I quess this is just shortly 22 Q. after the IARC classification? 23 24 Yeah, very shortly after. Α. All right. Go down to the e-mail. 25 It says Q.

1 right here --2 MR. BROWN: Excuse me, Your Honor. I'm going 3 to object. Foundation. Speculation. THE COURT: There's no question pending. 4 The document is in evidence. 5 MR. BROWN: Yeah, there's no foundation. 6 This witness isn't qualified to testify concerning this 7 document. 9 THE COURT: I don't know what the question is 10 yet. And so wait until the witness is asked a question 11 about it. BY MR. WISNER: 12 All right. So it says down here: 13 Q. I am the chair of the CARC and my 14 15 folks are running this process for 16 glyphosate in reg review. I've called a 17 CARC meeting in June. 18 We talked about CARC a second ago. Is that your understanding of that CARC there? 19 20 Α. Yes. MR. BROWN: Objection, Your Honor. 21 There's no foundation. This is hearsay as to this witness. 22 23 THE COURT: I'm sorry. The question was: Is this the CARC you're talking about, what are you 24 referring to? 25

MR. WISNER: A second ago, I asked him what 1 2 CARC was. 3 THE COURT: Right. MR. WISNER: And now I'm asking if that's the 4 same CARC. It's a document in evidence. I'm trying to 5 6 clarify what these terms mean. Sustain the objection. There's no 7 THE COURT: foundation for what that is by that author. 9 MR. WISNER: Okay. BY MR. MILLER: 10 Well, fair enough. 11 Q. 12 Are you familiar with something called reg review? 13 14 Α. Yes. What is that from a regulatory perspective? 15 That's this every 15 year rereview. It's the 16 Α. 17 regulatory review of all registered pesticides. And this, of course, is referring to the reregistration 18 review of glyphosate. 19 And in April of 2015 shortly after the IARC 20 classification, is your understanding that CARC was 21 doing a reg review for glyphosate? 22 23 Yes, it was. Α. And are you familiar with who was in charge of 24 Q. that reg review? 25

Yes, I am. 1 Α. And was that person by the name of Jess 2 0. 3 Rowland? Yes. 4 Α. Okay. Turn to Exhibit 705 in your binder. 5 Q. Is that a copy of the CARC review? 6 The report following the review, yes, it is. 7 Α. What is it dated, sir? 0. October 1, 2015. 9 Α. So it was after this e-mail that we looked at? 10 Q. 11 Correct. Α. And is Jess Rowland a signatory to this 12 0. document? 13 It's from Jess Rowland. 14 Α. Okay. One of the things that came up earlier 15 16 during Dr. Portier's testimony was a discussion of one 17 of these mouse studies. It was the Kumar study from 2005. Are you familiar with that study, sir? 18 19 Yes, I am. Α. And there was a discussion about the EPA 20 report dismissing this finding because of this supposed 21 viral infection. Are you familiar with that issue? 22 23 Yes, sir. Α. 24 In this CARC report that's in front of you, and if you want I can direct you to the page, is there a 25

1 discussion of that study, sir? 2. Yes, there is. Α. 3 Q. And is there language specifically related to a viral infection in the Kumar study? 4 Α. Yes. 5 Now, if you turn to Exhibit -- do you have the 6 Q. 7 page in front of you where it says that? 8 I'm -- what page is it? Let me find it for you. It would be I think 9 Q. 10 on page 40. 11 Α. It's a long one. 12 0. Yeah, I know. It's not page 40. Give me one second. 13 14 It will be page 39. Sorry. Do you have 39 there? 15 16 Yes, sir. Α. 17 And if you look at the bottom bullet point Q. there, do you see the discussion? 18 19 Yes. Yes. Yes. Α. And it talks about -- well, just read through 20 Q. that and make sure it's fresh in your mind. Okay? 21 22 Do you see that? 23 All right. So I want to turn to Exhibit 3036. 24 Okay? Is that the most recent version of the EPA's 25 3570

1 issue paper related to glyphosate? 2 Α. Yes, it is. 3 MR. WISNER: Permission to publish, It's actually been published before. 4 Your Honor. Any objection? THE COURT: 5 MR. BROWN: Not to 3036. 6 BY MR. WISNER: 7 So here we have Exhibit 3036. And just to 9 give the jury a quick background, what is an issue paper in the context of a CARC review? 10 It's where the EPA is explaining in detail the 11 Α. studies that it looked at in the area of animal 12 bioassays, in epidemiology, in genotoxicity, in 13 mutagenicity, in reaching its classification decision 14 about glyphosate relative to its potential to cause 15 It's like the full report, the most complete 16 17 articulation of the scientific basis for EPA's decision. Is it a final report? 18 Q. 19 Α. Not at this point. And this was in December of 2017? 20 Q. 21 Correct. Α. 22 Q. It's been over almost two years now. No, a year and a half? 23 Correct. 24 Α. And has the EPA reregistered glyphosate yet? 25 Q.

1 Α. No. 2 If you turn to page 70 in this document, we 3 have this discussion of the Kumar study at bullet Do you see that? 4 point 4. Α. Getting there. 5 6 Q. Sure. 7 Α. Yes. And it says here that this study was not Q. included due to the presence of a viral infection in the 9 colony which confounded the interpretation of the study 10 11 findings. Do you see that? Α. 12 Yes. It also notes that malignant lymphomas were 13 Q. 14 reported in this study in all dose groups. Do you see that? 15 16 Α. Yes. 17 Q. Now, it has a citation there, 14. Do you know where that goes to? 18 19 Citation to what? Α. 20 Q. It has a footnote 14. Oh, yes, it goes to the Greim's study. 2.1 Α. Okay. So they're citing a publication; 22 Q. 23 correct? Yeah, peer-reviewed publication. 24 Α. And have you reviewed that Greim study before? 25 Q.

Α. Yes. 1 Let's look at Exhibit 2064 in your binder. 2 Q. 3 Α. 2064. No, that's not it. Well, I don't have it. So we'll just move on. 4 0. We'll come back to that later. I don't want 5 6 to spend time on it until I have it. All right. One of the things that has been 7 discussed is -- well, on that viral infection, does the 9 Greim paper discuss it? Yes, it does. 10 Α. And does it characterize it as a speculative 11 Q. assertion? 12 13 Α. That's a fair characterization, yes. And yet the EPA report does not do that, does 14 Q. 15 it? It states it as the reason that they 16 Yes. 17 discounted the positive tumor finding in the study. All right. If you go back to the EPA report, 18 Q. I actually want to ask you about another thing that I 19 20 found interesting. 21 So we have here on page 85. Do you see that, sir? 22 23 I'm getting there. Α. Okay, I'm there. 24 And we have the Reyna and Gordon study from 25 Q.

1 1973. Do you see that? 2 Α. Yes. 3 Q. That study, Reyna and Gordon from 1973, are you familiar with that study? 4 Α. Yes. 5 All right. Let's go to the reference. 6 7 believe it's 156. And I can just pop it up here 8 quickly. 9 Reyna and Gordon, do you see that? 10 Α. Yes. 11 And this is the study that was performed by Q. Industrial Bio-Test Laboratories; is that right? 12 Yes, it is. 13 Α. Okay. I mean, is this the same study? 14 Q. The same invalid study, yes. 15 Α. 16 Hold on a second. Let's just verify this. Q. 17 So it says right here B569. Do you see that? Correct. 18 Α. 19 Let's go back and look at that EPA document Q. 20 that we were looking at about the studies. And if we go to glyphosate, and we look at the carcinogenicity study, 21 that's the one that was invalid; right? 22 23 Right. Α. 24 What's the identifier? Q. B569. 25 Α.

MR. BROWN: I'm sorry, which exhibit number is 1 2 that? 3 MR. WISNER: It's 1364. So that's the same study? 4 Q. Yes. 5 Α. And this is the study that was deemed invalid; 6 Q. is that right? 7 Correct. Α. 9 So if we go back to the actual paper and look Q. at what the EPA said about this study and it's 10 discussing Reyna and Gordon, does it mention anything 11 about the EPA reviewing the raw data and comparing it to 12 the results that conclude it's invalid? 13 14 Α. No. So, sir, based on what they say here in their 15 own documents, EPA is citing as evidence of a lack of 16 17 carcinogenicity a study based on fraud? Objection, Your Honor. 18 MR. BROWN: Lacks It's also argumentative. 19 foundation. 20 THE COURT: Sustained. 21 (Sidebar held but not reported.) BY MR. WISNER: 22 23 All right. So we were looking at this study Q. and we were talking -- and we were talking about the 24 Knezevich -- I'm sorry -- the Reyna and Gordon study. I 25

1 just want to establish this is in fact the IBT study that was deemed invalid? 2 Yes, it is. 3 Α. And having reviewed this document, have you 4 0. seen any concession or acknowledgment by the EPA that 5 this was in fact an invalid study? 6 There is none. Α. 7 All right. I want to talk a little bit about 0. 9 some of the data in here. I understand you've actually published an analysis comparing what the EPA did with 10 what IARC did; is that right? 11 12 Α. In the area of genotoxicity, yes. 13 Q. Exactly. Can we look -- if you look in your binder, 14 Exhibit 2349, is that a copy of your article, sir? 15 16 That's a copy of my, yeah, scientific paper Α. 17 published in Environmental Sciences Europe. MR. WISNER: Permission to publish, 18 Your Honor? 19 20 THE COURT: Granted. BY MR. WISNER: 21 All right. So we're looking at your 22 Q. 23 publication here. And it looks like the title of it is: 24 "How did the U.S. EPA and IARC reach diametrically opposed conclusions on the genotoxicity of 25

1	glyphosate-based herbicides?"
2	Do you see that?
3	A. Yes.
4	Q. Sir, why did you write this article?
5	MR. BROWN: Well, I'm going to object,
6	Your Honor. It's irrelevant. It also exceeds the scope
7	and it lacks foundation.
8	THE COURT: I think we're getting afield.
9	BY MR. WISNER:
10	Q. Okay. Let me ask you this. What did you do
11	in this paper? Don't tell me the conclusions. Just
12	tell me what you did.
13	MR. BROWN: Objection. Irrelevant.
14	THE COURT: Sustained.
15	Counsel come to the sidebar, please.
16	(Sidebar held but not reported.)
17	BY MR. WISNER:
18	Q. All right. Sir, I guess we can just cut to
19	the chase on this. Are you familiar with something
20	called a registrant study?
21	A. Yes.
22	Q. What is a registrant study in the context of
23	regulations?
24	A. It's a study either conducted in a laboratory
25	of a pesticide registrant or a contract laboratory on
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behalf of the registrant that is submitted to the EPA in fulfillment of a data requirement needed to support EPA approval of a pesticide use or a pesticide tolerance.

- Q. Are registrant studies made publicly available to be subject to peer review?
 - A. Typically not.
- Q. So, for example, all of these mouse studies that were done either by the registrant or a contract laboratory, were these published in peer-reviewed literature to be peer-reviewed at the time they were done?
 - A. No.

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- Q. So who does look at them when they're completed?
- A. The -- within the Toxicology Branch of EPA, any of these new studies would be assigned to a particular reviewer. In the case of the 1983
 Biodynamics study, to Dykstra, and he would look at the 2,000-page report and all the individual tables on the different mice and do a review, his own fresh review of the results reported in the study, and then prepare a memo to -- you know, other people in the Office of Pesticide Programs about his determination relative to what that study shows.
 - Q. Now, that article we had up a second ago, was

that article subject to peer review?

A. Yes.

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- Q. How much peer review?
- A. The journal actually had it reviewed by ten different peer reviewers.
 - Q. And walk the jury through what that means.
 - A. So when a --

MR. BROWN: Excuse me, Your Honor.

THE WITNESS: -- scientist submits --

MR. BROWN: Excuse me, Your Honor. This is irrelevant.

THE COURT: Overruled. He can answer.

THE WITNESS: When a scientist submits a paper to a scientific journal, it's the journal editor sends it out to other scientists with expertise in the field for their assessment of the validity of the data, the appropriateness of the analytical methods that are used to interpret the data, the conclusions that are drawn from the data, and any discussion of the implications, say, for the regulatory status of a pesticide or a policy or whatever.

BY MR. WISNER:

Q. And as the paper goes through this rigorous peer-review process, does it make generally the publication stronger?

- A. Oh, absolutely.
- Q. Why is that?

A. Because other scientists who often have much deeper expertise than -- I mean, I'm not a genotoxicologist, but my paper was reviewed by several of them, some of the top ones in the country, and they shared their suggestions on how I can improve the paper, make it clearer.

I had 26 pages of comments on this paper, and I believe strongly that the peer-review process made it a more -- a more solid and well-grounded paper.

- Q. Now, as part of this paper and as part of your work investigating in this case, have you looked at how often the registrant studies that are not subject to peer review are positive and how often the peer-review studies that are subject to that are positive?
 - A. Yes.
 - Q. And what is the difference?
 - A. It's quite substantial.
 - Q. What is it?
- A. Well, in the case of the genotoxicity data that I analyzed in this paper, the registrants -- the registrant studies covered 95 different assays. So this would be an individual test of whether glyphosate technical, the chemical that's in Roundup, or the

formulated product, which would be the glyphosate and the surfactants, triggered a genotoxic response in a given assay or a given test.

There were 95 of those registrant studies done and submitted to EPA, and one of them reported a positive response and 94 of them didn't.

- Q. So one out of 94; is that right?
- A. Correct.

- Q. And these are the ones that only the registrant and EPA look at?
 - A. Correct.
- Q. What about the ones that the rest of the world gets to look at?
- A. So in the -- both the EPA analysis and the IARC analysis, there were about 122 different assays considered that have been published in peer-review journals typically by scientists not working for pesticide manufacturers. And of those, 73 percent reported one or more positive response.
- Q. Do you know what the number was, since we did a fraction?
 - A. Yes, 89.
- Q. Okay. And I think that's approximately
 1 percent; is that right?
 - A. Correct.

So I just want to make sure I understand the 1 Q. 2 significance of this finding. The study is done by the 3 manufacturers that no one gets to see. Only 1 percent 4 are positive. The question is argumentative. 5 MR. BROWN: THE COURT: Sustained. 6 THE WITNESS: Well, the EPA gets to see them. 7 MR. WISNER: Don't answer. 9 THE COURT: Sustained and stricken. Move on. BY MR. WISNER: 10 Fair enough. I just want to clarify, though, 11 Q. the question I was trying to articulate is: This group 12 of studies, they're performed by the manufacturers; is 13 that right? 14 Either by them or under contract for them. 15 Α. And these ones are primarily done by 16 Q. 17 independent researchers? Typically university-based scientists, yes. 18 Α. MR. WISNER: Okay. I'm moving on to another 19 topic, Your Honor. This might be a good time for lunch. 20 That's fine. We'll take an hour 21 THE COURT: for lunch. Be ready at 5 of. 22 23 Don't discuss anything you've heard today or any other time during the trial. Enjoy lunch and we'll 24

see you in an hour.

(Recess taken at 11:56 a.m.)

(Proceedings resumed in open court in the presence of the jury at 1:04 p.m.)

THE COURT: You can continue, Mr. Wisner.

MR. WISNER: Thank you, Your Honor.

- Q. Doctor, just before lunch, we were looking at the 2017 EPA issue paper draft; correct?
 - A. Correct.
- Q. Okay. And I want to ask you a quick question before we move on from the document. If you look at -- on page 68 of the document at the very end of the epidemiological section, it says:

Based on the weight of the evidence, the agency cannot exclude chance and/or bias as an explanation for observed associations in the database. Due to study limitations and contradictory results across studies of at least equal quality, a conclusion regarding the association between glyphosate exposure and risk of NHL cannot be determined based on the available data.

Sir, does that mean that the EPA says it doesn't cause NHL?

A. No.

- Q. What does that mean?
- A. It means that it can't draw a definitive conclusion one way or the other from the data that it has available to it.
 - Q. It then says:

The agency will continue to monitor the literature for studies, and any updates to the AHS will be considered when available.

Do you see that?

A. Yes.

2.1

- Q. Since December of 2017, have there been new epidemiological studies published?
 - A. Yes.
 - Q. Can you give me -- what are you familiar with?
- A. There's a study in the journal Mutation

 Research by Professor Zhang and two other coauthors.

 That was a very sophisticated meta-analysis of all the epidemiological studies that had data associating exposure to Roundup and non-Hodgkin's lymphoma.

In addition, there was a very large study published by a team of French scientists that pooled data from Scandinavia, France, and the Agricultural Health Study into a combined data set which again, as I've said before, when you have more animals in an

1 animal study, the more people you have in an epi study, 2 the greater the statistical power is. And by that, that 3 means it has a greater chance of detecting a statistically significant increase in a disease outcome 4 like non-Hodgkin's lymphoma. 5 And that Zhang article that you mentioned, did 6 Q. that meta-analysis actually include AHS? 7 Α. Yes. 9 And did that meta-analysis show a Q. 10 statistically significant elevated rate for non-Hodgkin's lymphoma? 11 Yes, it did. 12 Α. Objection, relevance. 13 MR. BROWN: THE COURT: Sustained. 14 MR. BROWN: Move to strike. 15 16 THE COURT: Stricken. 17 MR. WISNER: Well, my question, Your Honor, I asked the question because I want to get to this point 18 which is: 19 20 Since the Zhang article came out showing the results that it showed, has the EPA issued a new 21 22 statement? 23 Objection. Relevance. MR. BROWN: Sustained. 24 THE COURT: 25 MR. WISNER: Okay.

I guess what I'm trying to ask is, is this 1 Q. 2 2017 issue paper the most recent iteration of the 3 reregistration process which is still ongoing? Yes, by EPA. 4 All right. I want to move on to a different 5 Q. topic now. I want to specifically talk about Dr. James 6 Parry. Are you familiar with who that is? 7 Yes, sir. And if you look in your binder, I believe it's 9 Q. Exhibit 38, is that one of Dr. James Parry's reports? 10 You said 38? 11 Α. 12 0. That's right. Yes, it is. 13 Α. MR. WISNER: Your Honor, this document is 14 already in evidence, and I will put it up on the screen. 15 MR. BROWN: Your Honor, I'm going to object. 16 17 It's not relevant. There's no question pending. 18 THE COURT: 19 (Exhibit published.) 20 BY MR. WISNER: Sir, have you reviewed this document? 2.1 Q. 22 Α. Yes. All right. And at the end of the document are 23 Q. a series of -- part of it says "Key issues concerning 24 the potential genotoxicity of glyphosate, glyphosate 25

1	formulations and surfactants, recommendations for future
2	work." Do you see that?
3	A. I'm getting there.
4	THE COURT: What page are you on, counsel?
5	MR. WISNER: It's on page 32 of the document
6	if you look at the bottom right-hand corner.
7	THE WITNESS: Yes, I'm there.
8	MR. WISNER: Are you there, Your Honor?
9	THE COURT: Yes. Go ahead.
10	BY MR. WISNER:
11	Q. So this these recommendations for future
12	work, have you studied them?
13	A. Yes.
14	Q. And have you looked to see whether or not the
15	recommendations and actions proposed by Dr. Parry back
16	in this time frame were ever done?
17	MR. BROWN: Objection. Relevance.
18	THE COURT: Overruled. He can answer.
19	THE WITNESS: Yes, I have looked at that.
20	BY MR. WISNER:
21	Q. Okay. And just to give some context, if you
22	look on the second page it says right here
23	"Recommendations for action." Do you see that?
24	"Actions recommended." Do you see that?
25	A. Yes.

- Q. And it has A, B, C, D, E. Do you see that?
- A. Correct.

- Q. And it goes on to the next page; correct?
- A. Through "I."
 - Q. Okay. Great.

If you look in your binder, sir, there's a chart that I believe you created, Exhibit 3093. It should be towards the end.

- A. Yeah.
- Q. What does that chart reflect?
- A. I have felt all along as I've reviewed the records in the case that the EPA and Monsanto handling of the genotox data were very important and the different view, for example, between EPA and IARC of the overall database. So I paid particular attention to Dr. Parry's report who had been hired by Monsanto to advise it on --
- MR. BROWN: Excuse me, Your Honor. The answer is nonresponsive.

THE COURT: What does the chart reflect?

MR. WISNER: It's the foundation for him getting to the answer, Your Honor. He's explaining what he did to get to this chart.

THE COURT: Get there now.

MR. WISNER: Okay.

THE WITNESS: In the deposition of a senior toxicologist for Monsanto, Dr. Donna Farmer, her --

MR. BROWN: Your Honor, I'm going to object.

This is irrelevant. Lacks foundation. And it's all hearsay.

THE COURT: Overruled. He can answer that question.

THE WITNESS: As part of that deposition,

Monsanto entered exhibits that were developed by

Dr. Farmer specifying in great detail what genotox

studies Monsanto did in response to Dr. Parry's report,

I mean down to the individual studies.

When I learned of that exhibit, I requested it. And that provided me with a definitive record of what Monsanto did in response to Dr. Parry's report, according to Dr. Farmer, the senior toxicologist that oversaw that area of work.

BY MR. WISNER:

- Q. And I guess the bottom-line question is: Did Monsanto do all of the recommendations that Dr. Parry made?
 - A. No, they did not.
- Q. And did you tabulate how many tests it would have taken to fulfill those recommendations?
 - A. I did.

1	MR. BROWN: Objection. Relevance.
2	THE COURT: Sustained.
3	MR. WISNER: Well, Your Honor, permission to
4	publish the chart?
5	MR. BROWN: Objection, no foundation.
6	THE COURT: It's a demonstrative.
7	MR. BROWN: As to this witness?
8	THE COURT: I understand he prepared this.
9	Did Dr. Benbrook prepare this chart?
10	MR. WISNER: Yes.
11	MR. BROWN: He prepared this?
12	MR. WISNER: Yes. I mean he just testified.
13	THE COURT: It's a demonstrative. Go ahead,
14	you may.
15	MR. WISNER: Okay. It just occurred to me,
16	let's make sure we're looking at the same one. I'm
17	looking at 3092. It's not 3093. Maybe that's the
18	confusion.
19	MR. EVANS: Yeah.
20	MR. WISNER: It's one page away.
21	BY MR. WISNER:
22	Q. Is that you're looking at 3093; right?
23	A. Yes, sir.
24	Q. Sorry, I think there was a miscommunication.
25	A. Well, no, I'm looking at 3092. It's the table

1	that I prepared.
2	Q. Okay. So 3092 is the exhibit. Is that the
3	table you prepared?
4	A. Yes, sir.
5	MR. WISNER: All right. Permission to publish
6	that, Your Honor?
7	THE COURT: I have a question I think I need
8	to ask you at sidebar.
9	MR. WISNER: Sure.
LO	(Sidebar held but not reported.)
L1	MR. WISNER: May I proceed, Your Honor?
L2	THE COURT: Yeah.
L3	BY MR. WISNER:
L4	Q. So, Doctor, let's keep it simple. Did
L5	Monsanto conduct the studies or all of the studies
L6	that Dr. Parry recommended?
L7	A. No.
L8	Q. Approximately what percentage of the studies
L9	that he recommended did they actually do?
20	MR. BROWN: Objection. Relevance.
21	THE COURT: Overruled. He can answer.
22	THE WITNESS: Less than half.
23	BY MR. WISNER:
24	Q. All right. There was well, let me ask you
25	something. You're familiar with regulatory obligations;
	3591

1 right? 2 Yes, sir. Α. 3 Q. And we have this report here. And if you look at it, it's Exhibit 38 in your binder and we were just 4 showing it, that was published in -- well, it was 5 submitted -- do you know about when this was submitted, 6 sir? 7 In the second half of 1999. And if you actually look at Exhibit 37, this 9 Q. is the one before, this is also in evidence, this is the 10 original report by Dr. Parry; is that right? 11 Α. Correct. 12 And if we go to the second page, you see the 13 Q. cover letter from him right here? 14 15 Α. Yes. Under EPA regulations, was Monsanto required 16 0. 17 to disclose these reports to the EPA? 18 Α. Yes. 19 Q. Why? Objection. 20 MR. BROWN: Relevance. Overruled. 21 THE COURT: He can answer. There's a provision in FIFRA, 22 THE WITNESS: 23 the statute governing pesticide regulation, 24 6(a)(2)/(b), it's called the adverse effects reporting requirement. And it places on the registrant 25

the responsibility to provide to the agency, after the agency has approved the pesticide, has looked at the studies, granted the labels, approved the tolerances, it requires the registrant, if they come into possession of any information that might shed new light on the risks associated with using the pesticide -- a poisoning episode, information they get from their manufacturing plant, a study that they're running where they get a preliminary report -- if any of this information suggests a new or higher risk than what has already been submitted to the agency, the registrant is bound by law to submit that information to the EPA within a specified time period. And that's what this 6(a)(2)/(b) requires.

BY MR. WISNER:

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Q. And if we turn to, for example, on page 11 of this document where it says right here:

The overall data provided by the four publications provide evidence to support a model that glyphosate is capable of producing genotoxicity both in vivo and in vitro by a mechanism based on production of oxidative damage.

This kind of conclusion from an independent expert that Monsanto hired, is that something that would

1	have been required to be disclosed?
2	A. Yes.
3	Q. All right. I want to talk to you about the
4	sort of use of glyphosate over time.
5	Sir, have you published about have you
6	published the rate or the amount through which
7	glyphosate is being used in the United States?
8	A. Yes, I have.
9	Q. And how many publications have you done on
10	this?
11	A. Two that have appeared in peer-reviewed
12	scientific journals.
13	Q. I understand as part of your analysis of it,
14	you actually prepared a chart sort of documenting the
15	usage of glyphosate in the United States?
16	A. Yes, I prepared a chart based on use data from
17	the Environmental Protection Agency.
18	Q. And I believe that chart is 3093; is that
19	right?
20	A. I suspect that's the correct number.
21	Q. Okay.
22	MR. WISNER: Your Honor, permission to
23	publish?
24	THE COURT: This is what the doctor was first
25	looking at?
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MR. WISNER: That's right.

MR. BROWN: Your Honor, I'm just going to interpose an objection. It's not relevant in the scope of this witness's expertise.

THE COURT: Overruled.

BY MR. WISNER:

Q. All right. Doctor, let me get up it up on the screen here.

(Exhibit published.)

MR. WISNER: Your Honor, permission for the witness to just come down and walk us through what this chart says?

THE COURT: Sure.

THE WITNESS: So periodically the EPA issued a public report on the pounds of different pesticide active ingredients applied by both by farmers and by non-agriculture users. And they literally provided a ranking from the most heavily applied pesticide in the United States in a given year to the number 25.

What I've done in this chart is beginning -this was the first year they issued this report in 1997.
Glyphosate was part of the report. It ranked 17 out of
the top 25 with 6 to 8 million pounds of glyphosate
active ingredient in the form of Roundup applied in the
United States by farmers. Which accounted for most of

the use.

The number one herbicide that year was a corn herbicide -- the number one pesticide was a corn herbicide called atrazine. See the rank here. This is 71 to 76 million pounds. In 1987, farmers planted approximately 70 million acres of corn so it was about a pound per acre of this most widely used corn herbicide applied.

Now watch what happens over the years. So the next report issued by the -- the years that I have the data, these are the years EPA put the report out. So six years later glyphosate had risen up to 11 with the use approximately doubling.

So, you know, there were 10 more pesticides used more wildly, but it was moving up in the rank.

Atrazine is still first. It jumps up to seventh. And two years later -- EPA put out the report every two years now for a number of years -- as you can see, atrazine stays number one all the way to here, 2001.

And look at how the use of glyphosate jumps from 25 to 30 million pounds, to 34 to 38, to 67 to 73. That's in only two years the use doubled, making it by 1999 the second most heavily applied pesticide in the United States by U.S. farmers. And it reached number one and where 85 million to 90 million pounds of

glyphosate were applied.

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And then if we can go to the other half --BY MR. WISNER:

- Dr. Benbrook, before we move on from this time 0. period, this time period right here, '93 to 2001, is that the same time period that the AHS was beginning to collect data about usage of glyphosate?
 - As a matter of fact, it is, yes.
- Okay. So then we go down to the next part, Q. starting 2001. Walk us through this, sir.
- So continuing with the periodic report that Α. EPA put out every two years through 2009. Then the last one they put out came out in 2012. You'll see that the growth of use of glyphosate was not over. 85 to 90 million, just two years later up to 128 million to 133 million. Two years later, now we're 147 million to 167 million.
- So atrazine is still -- it's still the second But by 2005, note there's twice as much glyphosate applied as atrazine. So not only has it jumped to the number one position --
 - MR. BROWN: Excuse me, Your Honor.
- **THE WITNESS:** -- but now there's twice as 23 much --
- 25 MR. BROWN: Excuse me, Your Honor. This is

1 all argumentative. 2 THE COURT: Overruled. But just stick with the numbers. 3 THE WITNESS: I'm almost done. 4 THE COURT: Just stick to the numbers. 5 6 THE WITNESS: So, again, continuing the data. Glyphosate remains number one. By 2012, total 7 agriculture usage is in the range of 270 million pounds to 290 million pounds, four times -- four times the 9 10 volume --MR. BROWN: 11 Excuse me. THE WITNESS: -- of the number 2. 12 13 MR. BROWN: Excuse me. It's argumentative. THE COURT: Sustained. 14 Sustained. 15 MR. WISNER: Can I have a sidebar? I actually 16 don't understand the objection. 17 (Sidebar held but not reported.) MR. WISNER: I'll continue, Your Honor. 18 Okay. So, Doctor, fair to say that the use of 19 Q. 20 glyphosate has substantially increased since at least 1987? 21 Correct. 22 Α. 23 And in your opinion, has this substantial increase affected the ability of scientists to properly 24 measure exposure in epidemiological studies? 25

1	MR. BROWN: Calls for speculation.
2	Foundation.
3	THE COURT: Sustained.
4	MR. WISNER: I'm a bit fish out of water here,
5	Your Honor. I need some guidance from the Court, sorry.
6	Because I don't want to do something that I can't do.
7	THE COURT: Dr. Benbrook is not an
8	epidemiologist.
9	MR. WISNER: No, no, that's not the issue.
LO	Okay. Well, I'll just keep going and see where it goes.
L1	Q. Are you familiar with the Farm Family Exposure
L2	Study?
L3	A. Yes, sir.
L4	MR. WISNER: Your Honor, am I allowed to
L5	inquire about that?
L6	THE COURT: You haven't inquired yet.
L7	MR. WISNER: Okay.
L8	Q. Have you reviewed the Farm Family Exposure
L9	Study?
20	A. Yes, I have.
21	Q. Have you discussed it in your own
22	peer-reviewed published literature?
23	A. Yes.
24	Q. And I want to draw your attention to the
25	study. It's Exhibit 1582 in your binder. It's also up
	3599

1 on the screen, sir. 2 I've got it. Α. 3 Q. Okay. And in this study -- you understand who performed this study? 4 Yes, sir. Α. 5 Who was it? 6 Q. The lead author was John Acquavella. 7 Α. And who was he with, sir? Q. 9 He was a Monsanto Company scientist. Α. All right. Is it your understanding that this 10 Q. study was in fact paid for and conducted by Monsanto? 11 Yes, sir. 12 Α. Okay. Now if we go down to the discussion in 13 Q. 14 result section. It says right here: The results of our analyses suggest 15 16 that modifying specific practices should 17 be effective in minimizing glyphosate exposures for farmers, spouses, and their 18 19 For farmers, the use of rubber children. gloves when mixing and loading pesticides 20 or when repairing equipment was associated 2.1 22 with measurably reduced urinary 23 concentrations. 24 Do you see that?

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Α.

Yes.

- Q. Sir, in your understanding of the EPA regulations, finding that in fact you could reduce exposure using gloves, does that trigger an obligation of Monsanto to either amend the label or inform the EPA about this finding?
- A. It certainly wouldn't trigger a positive requirement for EPA -- for Monsanto to change the label, but it's clearly this study generated data that is highly pertinent to the EPA's evaluation of mixer, loader, and applicator exposures and would be taken into account by the EPA in assessing whether the provisions on the label are adequate.

For example, do they require wearing gloves.

- Q. Now, remember earlier we were talking about the 1986 guidance?
 - A. Yes.

- Q. I'd like to go to that document. It's Exhibit 830 in your binder. Is that a copy of the document?
 - A. Yes, it is.
- Q. All right. It's been shown previously to the jury.
- And this is that guidance document you were discussing with the jury before?
 - A. Correct.

Q. Okay. And, you know, it occurred to me when we were discussing whether or not Monsanto would have been ordered to do another mouse study, I want to see what it says here about that so you can explain what that means.

It says here -- it says:

Therefore, in order to fully address this question, the agency is requiring that this study be repeated with a larger number of animals in each test group so that the statistical power of the study is increased.

- A. Yeah, that's what I described earlier.
- Q. Okay. I want to understand something. How is it possible that the agency could require it but it never happened?
 - MR. BROWN: Calls for speculation.
 - THE COURT: Overruled.

THE WITNESS: The EPA in this registration standard document and all other similar documents applying to other pesticides make certain requirements of the registrant to do new studies or clarify the language on the label or impose new protective clothing provisions to reduce exposures like wearing gloves.

And it assumes and in most cases it is correct

in assuming that the registrant will make those changes in order to have no trouble getting its future labels approved so that the pesticide can continue to be used consistent with the provisions laid out in the registration standard.

But sometimes the registrants choose not to do what is -- what they are asked to do by EPA and sends them a letter and challenges the substantive basis or the justification for the request.

For example, Monsanto submitted a letter in response to this registration standard document to EPA laying out the reasons that it felt that a repeat mouse study was not justified.

It also sent another communication addressing a series of worker safety provisions in this document that Monsanto was supposed to put on all Roundup labels.

MR. BROWN: Excuse me, Your Honor. The answer is nonresponsive.

THE COURT: Starting with it "sent another communication addressing a series of worker safety provisions" will be stricken.

THE WITNESS: Okay, sorry.

MR. WISNER: We'll get to that in a second.

Q. I was just trying to make sure I understood how --

1 I'm sorry. I did a -- may I finish answering Α. 2 your question? 3 Q. Sure. No. Your Honor. 4 MR. BROWN: THE COURT: Ask another question because I 5 struck the last portion of it. 6 BY MR. WISNER: 7 Please finish answering my question as it 9 relates to the mouse study. Okay? Right. If a company chooses not to do a study 10 Α. 11 requested in a document like this, the EPA has basically It can initiate a cancellation action to two choices. 12 try to drive the pesticide off the market, which is a 13 14 long and arduous process, or it can just let it go. Now, you talked about requirements for 15 16 labeling; right? 17 Α. I'm sorry, I didn't hear you. Briefly -- actually it was struck. 18 Q. 19 Let's go to page 32. 20 Α. 32? Yeah. Using the bottom right number, not the 21 Q. middle number. 22 23 And you see that point 4 at the very bottom? Yes, sir. 24 Α. 25 It says: Q.

1 The following worker safety rules 2 must appear on end-use products containing 3 glyphosate except for those labeled for homeowner use only. 4 Do you see that? 5 Yes, sir. 6 Α. And then if we turn to the next page, there's 7 Q. a bunch of language. Do you see that? 9 Yes. 10 Q. As, for example, wear goggles or face shield, chemical-resistant gloves, chemical-resistant apron, and 11 chemical-resistant shoes, shoe coverings or boots. 12 13 Do you see that? 14 Α. Yes. And it says wear the following protective 15 clothing during application. Do you see that? 16 17 Α. Yes. What is the EPA doing -- saying in this 18 Q. 19 document? 20 Α. It's saying that --2.1 MR. BROWN: Objection. The document speaks for itself. And it also exceeds the scope, Your Honor. 22 23 THE COURT: Overruled. He can answer that. 24 THE WITNESS: This is the language that the EPA is requiring or directing Monsanto to add to all of 25

1 the labels for end-use products. And those are the 2 products that a farmer would buy or a homeowner of a 3 Roundup product. So this is the language that was supposed to 4 appear on all of the labels. 5 BY MR. WISNER: 6 Did that language ever make it onto the label? 7 Q. Α. No. 9 Now are you familiar with something called Q. material safety data sheet? 10 Yes, sir. 11 Α. And have you reviewed the material safety data 12 0. sheet specifically for glyphosate or Roundup? 13 Well, actually several of them, yes. 14 15 Are you familiar with Monsanto's material safety data sheet? 16 17 I have reviewed that one too. What is Monsanto's material safety data sheet? 18 Q. MR. BROWN: Objection, relevance. 19 Exceeds the

THE COURT: Let me ask this question. Is it required in the regulatory process to submit safety data sheets?

scope in terms of regulatory.

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MR. WISNER: Yeah, I'll lay the foundation.

THE COURT: Lay the foundation first.

BY MR. WISNER:

- Q. Doctor, what is a material safety data sheet?
- A. It is a summary of the properties of a chemical, any risks that are associated if you spill some on you or get some in your eyes, directions for what should be done in the case of an accidental spill or exposure, numbers to call for medical assistance in an emergency, and information useful for the safe storage of the chemical, what temperatures it can be stored at.

So it's best practices for handling the chemical in a safe way.

- Q. Now, is there a regulatory structure that governs the creation of an MSDS?
- A. They're -- different agencies use them for different purposes. OSHA is obviously interested in what they -- what is in a material safety data sheet because they are primary documents in occupational safety and health programs at chemical plants or formulating plants or anywhere where chemicals are handled.
- Q. And does -- well, Monsanto manufactures glyphosate. Do they have a material safety data sheet for their own employees?

A. Yes.

Q. Have you reviewed that? 1 2 I have. Α. 3 Turn to Exhibit 3094. Is that a copy of the Q. Monsanto materials safety data sheet from 1995? 4 30, what was it, 94? 5 Α. 3094, yes. 6 Q. The last one. Ah, yes. 7 Α. MR. WISNER: Permission to publish, 9 Your Honor? MR. BROWN: Foundation. 10 THE COURT: Lack of foundation? 11 12 Is this related to the product that was used by plaintiffs? 13 14 MR. WISNER: Yes. 15 THE COURT: Okay. Overruled. BY MR. WISNER: 16 17 So we're looking at the material safety data Q. sheet. And if you see up here, lawn and garden 18 19 products. Do you see that? 20 Α. Correct. And you can see up here, this is the Monsanto 21 Q. lawn and garden products specific MSDS; right? 22 23 Correct. Α. 24 And if we go down here to the handling and storage -- let me find that section. 25

1 Okay. There's a section here that says 2 exposure control personal protection. Do you see that? Yes. 3 Α. And, for example, here under "Skin 4 Q. Protection, " it says: 5 6 Wear chemical-resistant gloves. Ιf there's a significant potential for 7 contact, wear face shield, wear 9 chemical-resistant clothing/footwear. Do you see that? 10 11 Α. Correct. And this is what Monsanto is telling their own 12 0. 13 employees about how to handle lawn and garden products 14 involving Roundup? 15 Α. Correct. 16 Do they give that same instruction to regular 0. 17 consumers? MR. BROWN: Objection, Your Honor. 18 Exceeds 19 the scope. THE COURT: Overruled. He can answer. 20 THE WITNESS: No, they didn't put these 21 provisions on the label. 22 23 BY MR. WISNER: 24 All right. The last thing I want to talk to Q. you about -- well, there's two last things I want to 25 3609 talk to you about, but they're pretty quick.

I understand you recently published an article about POEA; is that correct?

A. Correct.

- Q. Tell the jury a little bit about -- don't tell me what you said in your article, but describe how you went about preparing that article.
- A. I wrote this article with two scientists at King's College in London who are well-known experts in pesticide toxicology. They've done a lot of research over many years on the formulated products which includes the active ingredient glyphosate plus the surfactants, which the U.S. EPA regards the surfactants as inert ingredients.

And what we did in this paper is we tried to help the scientific community that's doing research on glyphosate-based herbicides like Roundup that's got a certain amount of glyphosate in it and a certain amount of surfactant, to help scientists understand exactly what the test substance they were working with.

Because the identity and the concentration of the surfactants are classified as confidential business information and are not disclosed on the label and are not available anywhere publicly.

So scientists will purchase Roundup Pro from a

retail dealer, and they'll know the percentage of glyphosate in it, but they won't know what the surfactants are.

The only thing that the label will disclose is the percent of the product by weight, that's the surfactant. And what a large body of science has shown over the last 20 years is that the surfactants in Roundup --

MR. BROWN: Excuse me, I'm going to object. This is beyond the scope.

THE COURT: Does this relate to registration?

MR. WISNER: I'm going to get there, but I --

MR. BROWN: No, Your Honor.

MR. WISNER: Let me ask the next question.

THE COURT: Let's get to the relationship with that to registration.

MR. WISNER: Okay.

- Q. The jury has actually already seen your article with Dr. Sawyer. They've seen what you said about POEAs.
 - A. Okay.
- Q. So we actually don't need to get into that fact.

What I want to ask you about is what is -- well, let me start off with the first question.

Under the regulations, is Monsanto required to 1 do a long-term animal cancer study on POEA? 2 3 Α. No. Are they required to do -- well, okay. 4 Q. the other question I have for you is: Has the POEA 5 formulation evolved since Roundup first came on the 6 market in the '70s to what it is today? 7 MR. BROWN: Objection, relevance. THE COURT: Overruled. You can answer. 9 THE WITNESS: Yes, it has. 10 BY MR. WISNER: 11 And as they changed the formulations of this 12 0. 13 POEA surfactant, has Monsanto been obligated to do cancer studies on each one of those new formulations as 14 15 it comes out? 16 Α. No, they have not. 17 Q. How do we know they're safe? Objection. Calls for speculation. 18 MR. BROWN: It's argumentative. 19 20 THE COURT: Sustained. BY MR. WISNER: 21 Let me just be clear. The type of Roundup 22 Q. 23 that existed in 1982, did that use the original POEA formulation? 24 The percentage of surfactant and the exact mix 25 Α. 3612

1	of POEA molecules in the primary Roundup products on the
2	market in 1982 differed a little bit from the products
3	initially registered and sold in 1974, but not
4	substantially.
5	Q. And that type of POEA that was around in 1982,
6	is that type of POEA allowed to be used in Europe?
7	A. Not anymore.
8	Q. And is that based on some of the research that
9	you reported on in your article?
10	MR. BROWN: Objection. Relevance.
11	THE COURT: Sustained.
12	BY MR. WISNER:
13	Q. Let me clarify the question. I think it was a
14	confusing question.
15	The studies that you document and discuss in
16	your article that the jury has already seen, are those
17	the studies that were referenced when it was banned in
18	Europe?
19	A. Correct.
20	MR. BROWN: Objection. The question is
21	(Interruption.)
22	THE COURT: Just wait a moment.
23	You interposed an objection to the question.
24	MR. BROWN: I'll repeat it, Your Honor.
25	THE COURT: I'm looking at it. I'm sustaining

1 the objection. 2 BY MR. WISNER: 3 Q. All right. Doctor, in Europe, the surfactants that are used, they're not POEA; is that right? 4 MR. BROWN: Objection. Relevance. 5 THE COURT: How does this relate to 6 registration here in the United States of the Monsanto 7 products? 8 9 MR. WISNER: It's a foundational question to 10 the next question. Okay. Go ahead. I'll just 11 THE COURT: overrule the objection subject to the next question. 12 THE WITNESS: The principal surfactants used 13 in Monsanto brand Roundup products in Europe are 14 15 different from the ones now incorporated in Roundup in 16 the United States. 17 BY MR. WISNER: And has Monsanto, to the best of your 18 Q. knowledge, attempted to replace those surfactants used 19 20 in Europe with the current ones being used in the U.S.? 21 Could you ask that again? Α. Sure. From a regulatory respect -- let me ask 22 Q. 23 you a simple question. 24 From a regulatory perspective, would there be anything stopping Monsanto from using the less toxic 25

stuff in Europe here in the U.S.?

A. No.

2.

Q. Okay. The last thing I want to ask you about, sir, and actually we'll just do this orally.

I want to talk about obligations that Monsanto had, okay, under the EPA regulations.

If Monsanto knew that Roundup was genotoxic to humans, is that something that they would have been required to warn about?

- A. They would be required to submit the underlying basis for that insight or conclusion, whether it was a study that they commissioned or a report from a recognized expert like Dr. Parry.
- Q. Sure, sir. I'm asking a very simple question.

 I'm talking about obligations to warn. If Monsanto

 knows its product can be genotoxic in humans, do they

 have to warn?
- A. There's no obligation in the FIFRA statute to do that.
 - Q. What about if they know it causes tumors?
- A. Again, there's no requirement that they put that on the label of their product. It does have to go on the -- on an OSHA information sheet that's governed by regulations under the OSHA statute.
 - Q. Sir, I think we're talking past each other

1 here. Under FIFRA, if Monsanto knew that their product 2 could cause cancer, do they have to warn people? 3 MR. BROWN: Your Honor, I'm going to object. Calls for a legal opinion. 4 THE COURT: Reframe the question. 5 MR. WISNER: Sure. 6 Under the regulatory framework, if Monsanto 7 Q. knows that their product can cause cancer, do they have an obligation under the regulatory standards to warn? 9 MR. BROWN: Calls for a legal conclusion. 10 They have a --11 THE WITNESS: THE COURT: Overruled. He can answer. 12 13 THE WITNESS: They have a regulatory obligation to provide the basis for that insight to the 14 15 EPA. 16 BY MR. WISNER: 17 Does the same thing apply to genotoxicity, Q. oncogenicity, oxidative stress? 18 Yes, sir. 19 Α. Okay. And to the best of your knowledge, 20 Q. 21 sitting here today, has any label related to lawn and garden products ever warned about any of that for 22 23 Roundup? Not that I'm aware of. 24 Α.

Thank you.

MR. WISNER:

1 No further questions, Your Honor. 2 THE COURT: Do you need a break or want to go straight into cross-examination? 3 I'm good if we're good. 4 MR. BROWN: Okay. We'll probably take a break 5 THE COURT: 6 in the next 20 minutes. 7 **CROSS-EXAMINATION** BY MR. BROWN: Since we're talking about labeling 9 Q. obligations, as I understand it, the EPA -- we've never 10 met. I'm Gene Brown, by the way. 11 I'm Dr. Benbrook. 12 Α. You're Mr. Brown. 13 Q. Okay. Does the EPA approve labeling? 14 Α. Yes. 15 And you can't put the product on the shelf without the EPA-approved label; correct? 16 Well, not legally. 17 Α. Right. And so the EPA registers the product; 18 Q. correct? 19 20 Α. Yes. 2.1 And it approves the label? Q. That's a part of the process, yes. 22 Α. 23 Now, Dr. Benbrook, you were asked some Q. questions about the work that you did while you were an 24 adjunct professor at Washington State University; 25

1 correct? 2 Α. Yes. And what is an adjunct professor? 3 Q. I had a position as an adjunct faculty member 4 Α. for three or four years prior to receiving an 5 6 appointment as a research professor which was 100 percent time paid position. 7 As an adjunct faculty member, there's no 9 compensation or no teaching role typically. So I had two different positions at WSU with 10 the adjunct position preceding the three years that I 11 served as a research professor. 12 Okay. And when you were working at Washington 13 Q. State University, did you have an office there? 14 No, I did not. 15 All right. So you were -- and Washington 16 Q. 17 State is in Pullman; right? 18 Α. Right. So how often did you go to campus? 19 Q. About three or four times a year. 20 about 90 miles south of Pullman. 21 And when you were at Washington State, were 22 Q.

you actually working for the Center for Sustainability?

23

24

25

A. That's not exactly the title of it. But, yeah, my program was affiliated with a center on natural

resources and environment or something. I could refresh my memory by looking at my résumé to get the exact name of the center. But, yes, I was affiliated with that center.

- Q. And then you told us that you were working for the National Research Council 1984 to 1990; is that correct?
- A. I spoke about that earlier. That was an earlier part of my career, yes.
- Q. Okay. And that was in 1984 to 1990.

 And you said that was an independent body that offered advice to the government.
 - A. Correct.
 - Q. And you left there in 1990.
 - A. Yes.

- Q. Okay. And why did you leave that job?
- A. It was time to move on. We had built a very effective program. And I ran what was called the board on agriculture. And the National Academy of Sciences wanted to see the board move in some different directions. And I agreed that I wasn't the right person to run it and so I moved on.
- Q. Okay. And who was the head of that organization at the time that you left?
 - A. A gentleman named Frank Press.

- Q. And you and Mr. Press had some disagreements?
- A. I don't know if I would characterize them exactly that way. But he was the president of the Academy at the time, and he and I reached an agreement that it would be best if I moved on.
 - Q. Did he ask you to leave?
 - A. Yes.

Q. Sorry about that.

And you have been working at Benbrook Consulting Services since 1990; correct?

- A. Yes.
- Q. And that's your primary work; is that correct?
- A. It has been over the period of time, although I had the three -- the three years that I was a research faculty member at WSU, was in there. And I also served for about six years as the chief scientist of a nonprofit organization called the Organic Center.
- Q. And the Organic Center was an advocacy group; is that correct?
 - A. It was a research science group.
- **Q.** And you testify for plaintiffs in litigated matters; is that correct?
- A. Over the years, yes, I have done other expert witness work.
 - Q. And do you also testify for defendants, folks

1 who are sitting on my side of the table? No. Most -- I think all the cases I've worked 2. 3 with the plaintiffs' attorneys. And, again, as I understand it, you are not a 4 Q. medical doctor? 5 Correct. 6 Α. You're, in fact, not an expert in physical 7 Q. 8 sciences; is that correct? I have no advanced degree in any of the 9 physical sciences. 10 11 You are not a toxicologist. You don't hold Q. yourself out as a toxicologist. 12 Α. Correct. 13 14 You're not an epidemiologist. Q. 15 Correct. Α. 16 You're not an industrial hygienist. Q. 17 Are we going to go through them all? Α. 18 Q. Yes, we are. 19 No, I'm not. Α. You're not a pathologist. 20 Q. 2.1 Correct. Α. 22 You have no formal training or degree in Q. 23 exposure assessments. 24 That's correct. Α.

25

Q.

You're not a lawyer.

Α. That is true. 1 2 You're not a geneticist. Q. 3 Α. Correct. And so what you do, Dr. Benbrook, is, for 4 Q. purposes of this case is, you've read literature; 5 correct? 6 I've read a significant body of literature, 7 Α. yes. All right. And you draw opinions from your 9 review of that literature. 10 Among other sources of information, yes. 11 Α. Right. You consider information, you read 12 0. literature. And then you -- that helps refine your 13 14 opinions; correct? 15 Α. Yes. 16 Okay. And then you come into court and you 17 tell people about what you read and how it -- and what you relied on to render your opinions; correct? 18 19 In effect, yes. Α. Now the material that you have reviewed to 20 Q. render your opinions in this case, is that information 21 available to us? 22 23 Α. Yes. 24 And we can get it just -- I would assume and I'm not going to go there. I was going to talk about 25

2 that. Is it available in libraries? 3 Well, certainly the entire discovery record 4 Α. would not be available in a library. 5 Well, not the entirety of it, but you can 6 Q. certainly find EPA publications. 7 Of the documents that I reviewed specifically 9 for this case, the ones that would be publicly available 10 were the peer-reviewed published scientific studies and 11 the primary EPA reports and the EPA memos that have gone 12 through the EPA clearance process to be made publicly 13 available. And those categor -- those groups of documents would be perhaps a quarter of the documents 14 that I reviewed in preparation for my testimony today. 15 All right. Have you reviewed any documents in 16 0. 17 preparation for your testimony today that reflect that glyphosate or Roundup are not registered by the EPA? 18 19 Α. Could you ask that question --20 Q. I'll make it easier. Is Roundup registered by the EPA? 21 MR. WISNER: Objection. Asked and answered. 22 THE COURT: Overruled. He can answer. 23 24 MR. WISNER: Okay.

THE WITNESS:

doing research on it, but I'm not going to talk about

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Yes, Roundup is registered by

1	the EPA.
2	BY MR. BROWN:
3	Q. And you said something about glyphosate being
4	banned in Europe. Do you recall that testimony?
5	A. Certain formulations of Roundup herbicide in
6	Europe are no longer marketed because European
7	regulators instructed Monsanto to alter the surfactants
8	in the formulation.
9	Q. All right. And has EPA requested that
LO	Monsanto alter the formulas here?
L1	A. No, they have not.
L2	Q. Now you talked about earlier in your
L3	testimony you talked about Dr. Franz. Do you recall
L4	that?
L5	A. Dr. who?
L6	Q. Franz.
L7	A. Yes.
L8	Q. All right. And how glyphosate was kind of
L9	just discovered.
20	A. The herbicidal activity of glyphosate was
21	discovered.
22	Q. Okay. And where did you learn that
23	information?
24	A. Oh, many different places. The history of
25	glyphosate as a herbicide has been covered in
	3624

innumerable papers and reports. There's an excellent detailed timeline on the Monsanto website about the history of glyphosate that I've read many times over the years.

You know, glyphosate is the most widely used herbicide in the world. So there's a lot of information about it and a lot of people interested in it.

- Q. There's a lot of information, there's been a lot of studies about glyphosate; is that correct?
 - A. Well, certainly several, yes.
- Q. You also talked to us about some work that was done at a lab called IBT?
 - A. Correct.

- Q. And what was IBT?
- A. It was a private commercial testing laboratory established in 1950 in Northbrook, Illinois that provided testing services for the pharmaceutical industry, the pesticide industry, the oil and gas industry, to generate the data that government agencies were starting to ask for. Because at that time we -- scientists had become concerned about the impact of chemical exposures on certain diseases.
 - Q. Okay. And this is an independent lab?
 - A. Yes, it's privately owned.
 - Q. All right. And at the time that this

1 difficulty with IBT was discovered, I think you told us, 2. and I would like for you to remind me, how many separate 3 clients or companies were using IBT's services? I don't know the exact number, but it would be 4 several dozen. 5 6 All right. So this wasn't just Monsanto that Q. IBT had a problem? 7 Oh, certainly not. Α. 9 Okay. And as a matter of fact, did any Q. governmental agencies use the services of IBT? 10 I don't know if they did or not. 11 Α. 12 0. But, again, this wasn't just a Monsanto 13 problem. Oh, yes, most definitely. 14 Α. And as a result of the difficulties with IBT, 15 0. 16 the companies were requested to redo studies; correct? 17 Either they realized they had invalid data and Α. they started the repeat studies themselves, or they 18 waited until they were asked to repeat them. 19 20 Q. All right. And that would include Monsanto? 2.1 Α. Correct. Now, you were asked some questions about 22 Q. Dr. Parry and recommendations. Do you recall that? 23 I do. 24 Α. All right. And you said that Dr. Parry had 25 Q.

been retained by Monsanto; correct?

- A. That is correct, yes.
- **Q.** All right. And Dr. Parry was an independent investigator?
 - A. He was a university professor in the UK.
- Q. All right. And was there any -- are you aware of any obligation being imposed by anyone that Monsanto follow all the recommendations that were handed down by Dr. Parry?
 - A. I know of none.
- Q. As a matter of fact, Monsanto had retained Dr. Parry and presumably paid him; correct?
 - A. They did.
- Q. Okay. And then you were asked a question about reregistration. And then you were asked a question about whether or not -- and I'm paraphrasing -- but whether or not the EPA could remove the registration for companies; correct?
 - A. Yes, I addressed that issue.
 - Q. Okay. And they do have that ability; correct?
 - A. They can initiate the process, yes.
- Q. Okay. And it would be fair to say that they would initiate that process if they saw a problem that was significant enough and posed a significant enough hazard to the citizens of the United States that they

1	would go ahead and initiate it in order to provide
2	protection; correct?
3	MR. WISNER: Objection. Speculation.
4	THE COURT: Sustained.
5	BY MR. BROWN:
6	Q. But no such action has been taken against
7	Monsanto, glyphosate or Roundup; correct?
8	A. No action to cancel the registrations, that's
9	correct.
10	MR. BROWN: Your Honor, if I could consult
11	with my colleagues, if you'd like to take a break.
12	THE COURT: Yes, sure. We can take a break
13	now. 15 minutes and we'll resume at 25 after the hour.
14	(Recess taken at 2:12 p.m.)
15	(Proceedings resumed in open court in the
16	presence of the jury at 2:27 p.m.)
17	THE COURT: Mr. Brown.
18	MR. BROWN: Thank you, Your Honor.
19	Q. Okay. Dr. Benbrook, just a few more
20	questions.
21	A. Okay.
22	Q. You were talking about earlier EPA
23	classifications. Do you recall that?
24	A. Classifications of what?
25	Q. Of, you know, for possible carcinogens,

1 probable carcinogens. 2 Okay. I understand. Α. 3 Q. Okay. And just so -- I made a note here of the dates you're talking about. Is it true that in 4 1985, the EPA classified glyphosate as a class C 5 possible human carcinogen? 6 7 Α. Correct. And then in 1986 it was reclassified as a 0. 9 group or class D; is that correct? No, that's not correct. 10 Α. Okay. When did that happen? 11 Q. 1991. 12 Α. 13 Q. 1991. And in 1991, what was the group or 14 classification? It would be not likely to pose oncogenic risk. 15 Α. 16 Okay. Evidence of noncarcinogenicity for 0. 17 humans? Correct. 18 Α. 19 Okay. And that was in 1991. Q. And is that the classification that exists 20 2.1 now? 22 Yes, it is. Α. 23 And we've been talking about the EPA and tests Q. and Monsanto and tests for a couple weeks now. 24 these tests and examinations of the various chemicals 25

and substances are done by lots of different people; correct? Not just Monsanto and the EPA.

- A. Well, all registered pesticides have to fulfill all the data requirements. So all the different pesticides companies have to do it. Are you referring to like the data requirements for drugs and other chemicals? Or I don't quite understand your question.
- Q. I'm just talking about regulatory agencies, governmental agencies, they're doing different tests on different substances for purposes of regulation, et cetera.
- A. Government agencies do very few tests themselves. They tend to require the private companies that are marketing chemicals to do the testing.
- Q. Now, did you have an opportunity to review documentation indicating in or about May, May 1st, 2013, the EPA released a pesticide residue tolerance approval for glyphosate?
- A. I'd be glad to look at the Federal Register notice for whatever specific tolerance action there was. I mean, there are several of them per year. So I don't know exactly which one you're referring to.
- Q. You are familiar with the September 12, 2016, EPA draft preliminary risk assessment?
 - A. Yes.

Q. And do you recall the conclusion of that 1 2 assessment? 3 Α. Yes. What is that? 4 0. That in terms of oncogenicity, I assume that's 5 Α. what your question is in reference to? 6 7 Q. Yes. That under current and expected levels of dietary exposure for the general public, exposure to 9 glyphosate does not pose a significant cancer risk. 10 And, again, the EPA, we know about EFSA and 11 Q. ECHA, but the EPA is the entity that regulates the 12 industry here in the United States; correct? 13 Yes. 14 Α. We talked a little bit about rodent studies; 15 0. do you recall that? 16 17 Α. Yes. And, again, you don't conduct animal assay --18 Q. 19 bioassays? 20 Α. That's correct, I do not. Okay. And are you familiar with Dr. Jameson? 21 Q. I know that he's an expert in this litigation. 22 Α. 23 I've never met him. And are you aware that a mice study was done 24 Q. where five pathologists were asked to review pathology 25

in a blind review to determine the existence or not of tumors?

- A. I have no idea what you're talking about.
- Q. Finally, in terms of the EPA, because we talked a lot about the EPA today, when they review a substance for registration, the review is done and assessed by the staff there at the EPA?
 - A. Correct.

- Q. And the staff, the staffs are made up of epidemiologists, toxicologists, a whole series of scientists?
- A. Correct. In each of the branches of EPA, depending upon the aspect of pesticide risk assessment that the branch is responsible for, it hires scientists with the requisite technical skills and training.
- Q. All right. And you don't have any reason to doubt the technical skill, expertise, or training of any of those scientists, do you?
- A. That's an awfully broad question. I've not done an analysis of the résumés of all EPA staff. So I really have no basis to answer that question.
- Q. But you have no reason to doubt that they're competent to do what they've been hired to do?
- A. I think like in any organization, there's a range of skills that are -- that different people bring

1 to the job. You know, I'm not going to -- I'm not going 2 to say that universally every scientist that works for 3 the EPA is at the cutting edge of their field and has done, you know, top quality work. I'm not prepared to 4 say that. 5 But at the same time, you cannot -- you can't 6 Q. say that they are not; correct? 7 That's also correct. I'm not rendering a 9 judgment one way or another. 10 MR. BROWN: Thank you. 11 THE COURT: Any questions on redirect? 12 MR. WISNER: Yes, Your Honor. Very few. 13 REDIRECT EXAMINATION 14 BY MR. WISNER: On cross-examination, Mr. Brown asked you a 15 16 question about the documents that you reviewed as part 17 of your analysis in this case. Correct. 18 Α. You said something like 25 percent of them are 19 Q. 20 public documents; is that right? 2.1 Α. It's a rough estimate, yes. What are the other 75 percent? 22 Q. 23 They're internal documents, recording Α. communications between Monsanto scientists and managers, 24 and Monsanto PowerPoints and training PowerPoints, 25

1 marketing materials, descriptions of Monsanto's 2 stewardship pledge and commitment to safety, and all 3 other sorts of information that Monsanto provided in response to the discovery requests in this litigation. 4 And to be clear, sir, those 75 percent of 5 Q. 6 documents that you looked at, are those available to EPA? 7 Most of them would not be, no. 9 Okay. There was a question about IBT; do you Q. recall that? 10 I recall that there was a question, but I 11 Α. don't remember exactly what it was. 12 13 Q. Sure. There've been a few. 14 Α. There was a question specifically about 15 16 whether it was related to Monsanto or not; right? 17 Correct. Α. Okay. And I just want to clarify something. 18 One of the scientists that was implicated in the IBT 19 20 scandal was Dr. Paul Wright; right? 21 Α. That's correct. And Paul Wright, before he worked at --22 Q. 23 MR. BROWN: Excuse me, Your Honor. Exceeds 24 the scope. The question was, was 25 THE COURT: Sustained.

1	Monsanto the only company.
2	MR. WISNER: No, Your Honor, there was a
3	question specifically about it being an independent lab.
4	That was the question.
5	THE COURT: The question was: Was IBT an
6	independent lab? And the answer was yes.
7	MR. WISNER: Exactly, and I'm probing that
8	point.
9	THE COURT: Okay. Be careful.
10	MR. WISNER: Sure.
11	Q. Dr. Paul Wright, before he went to IBT and was
12	involved in this fraud, he actually worked at Monsanto?
13	A. That's correct.
14	MR. BROWN: Objection. Argumentative.
15	THE COURT: Sustained.
16	MR. BROWN: Move to strike.
17	THE COURT: Stricken. MIL.
18	MR. WISNER: Your Honor, this
19	THE COURT: MIL.
20	MR. WISNER: That's fine. Yes, Your Honor
21	we'll talk about it later. I don't mean to violate any
22	MIL. I do believe it came in already. That's fine.
23	Q. Doctor, there were questions about labeling.
24	Do you recall that?
25	A. Yes.

Let me clarify something. Is Monsanto 1 Q. prohibited from proposing amendments to the label? 2 3 Α. No, they're not. Under the regulations, is Monsanto prevented 4 0. from warning consumers that it's genotoxic or oncogenic? 5 No, they're not. 6 Α. Is Monsanto prevented from conducting animal 7 Q. tests, for example, the ones that Dr. Parry recommended? 8 9 MR. BROWN: Objection, Your Honor. 10 exceeds the scope. I'm going to allow the answer to 11 THE COURT: that question, but we're moving afield from the cross. 12 13 **THE WITNESS:** No, they're not prevented. MR. WISNER: Fair enough. Your Honor, he 14 15 asked a question about whether or not they're required 16 to do it. I'm just asking if they were prevented. 17 Okay. Finally, Doctor, I guess that was close Q. to my last question, and that is simply put: Based on 18 your review of the record, has Monsanto ever made a 19 20 request to amend the Monsanto Roundup label to warn 21 about cancer? MR. BROWN: Calls for speculation. 22 Lacks 23 foundation. THE COURT: Lacks foundation regarding "ever 24 made request." Does he know that?

1	MR. WISNER: Fair enough.
2	Q. Have you reviewed all the regulatory
3	submissions from Monsanto to the EPA?
4	MR. BROWN: Calls for speculation.
5	THE COURT: Overruled. He can answer the
6	question.
7	THE WITNESS: Since 1974, Monsanto has
8	BY MR. WISNER:
9	Q. Answer my question first. Have you reviewed
LO	the records, what they sent to the EPA?
L1	A. I have reviewed a great number of records, but
L2	I have certainly not reviewed every document in that
L3	record.
L4	Q. Okay. And in the records that you have
L5	reviewed, spanning from 1974 to the present, has
L6	Monsanto, in those records that you've reviewed, ever
L7	proposed amending the Roundup label to warn about
L8	cancer?
L9	A. No.
20	MR. WISNER: No further questions, Your Honor.
21	THE COURT: Any more questions?
22	MR. BROWN: No, Your Honor.
23	THE COURT: Okay.
24	MR. WISNER: Thank you, Dr. Benbrook.
25	THE COURT: Thank you, Dr. Benbrook.

1 THE WITNESS: Thank you, Your Honor. 2 Do I get to keep my binder? 3 MR. WISNER: I'll take care of it, sir. THE COURT: Take anything that you brought, but leave everything else. 5 6 THE WITNESS: Okay. (Witness excused.) 7 MR. WISNER: Okay. Your Honor, before we move on to the next witness, we're going to read two 9 admissions into the record. 10 Admission number 6: 11 Admit that Monsanto has never 12 13 conducted any animal carcinogenicity study of any of the glyphosate-containing 14 formulations sold in the United States. 15 16 Response: Monsanto admits that it 17 has not conducted a long-term animal carcinogenicity study on any formulated 18 pesticide products. 19 Admission number 7: Admit that 20 2.1 Monsanto is not precluded by any applicable law, regulation, or ordinance 22 23 from conducting a long-term animal 24 carcinogenicity study on a glyphosate formulation. 25

1	Response: Monsanto admits that it is
2	not precluded by any applicable law,
3	regulation, or ordinance from conducting a
4	long-term animal carcinogenicity study on
5	a glyphosate formulation.
6	And with that, Your Honor, we will call
7	Dr. Kavitha Raj, treating physician for Mr. and
8	Mrs. Pilliod.
9	This was taken on January 8th, 2019, in
10	Pleasanton, California.
11	The entire video lasts one hour and 28
12	minutes sorry the whole thing lasts two hours. The
13	direct is one hour 28 minutes, the cross is 26 minutes.
14	THE COURT: Okay.
15	(Video excerpts from the deposition testimony
16	of Kavitha Raj played in open court; not reported
17	herein.)
18	THE COURT: There's nothing on my screen.
19	There's nothing on my screen.
20	MR. MILLER: We'll fix that.
21	MR. WISNER: May I just look at that screen to
22	see if it's this one?
23	THE COURT: Sure.
24	MR. WISNER: It's not there either.
25	THE COURT: If it doesn't work, I can look

1 over there. It's not a problem. Don't move a lot of 2 things around. 3 MR. WISNER: It's weird because that one is not working either. 4 It's the same -- given that size 5 THE COURT: it's about the same as looking at that. So that's fine. 6 We'll just check it later at the break. 7 Just keep going. 9 (Video excerpts from the deposition testimony 10 of Kavitha Raj resumes playing in open court; not 11 reported herein.) 12 THE COURT: We're going to take a quick 13 10-minute break. How long have we -- we're going to quit at 4:30 so I thought we needed to take a quick 14 15 10-minute break right now. 16 (Recess taken at 3:24 p.m.) 17 (Proceedings resumed in open court in the presence of the jury at 3:39 p.m.) 18 THE COURT: Okay. We're going to continue 19 with the video. 20 MR. WISNER: Yes, Your Honor. 21 I just want to clarify something because it 22 23 occurred to me that I hadn't explained it and I just talked to counsel about it. 24

The way this deposition was taken was the

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first portion of it, I did the deposition, and then
Monsanto did it, but it was about Mrs. Pilliod.

And then the second portion, I did it. So it actually kind of goes back and forth a couple of times, but the first part is about Alberta and the second part is about Alva.

THE COURT: That's fine. Good to know because actually the jurors need to know.

MR. WISNER: Yeah.

(Video excerpts from the deposition testimony of Kavitha Raj resumes playing in open court; not reported herein.)

THE COURT: Let's end for the day and start tomorrow.

Ladies and gentlemen, we're going to break now because we'll start with testimony about Mr. Pilliod separately fresh tomorrow.

So thank you for your time and attention today. We will be in session at 9:00 o'clock tomorrow. I know we started a little bit late the last couple of days. We're going to try to start on time. And thank you for everything.

And don't talk about anything, don't talk about the case to anyone. And have a good evening, forget you're a juror, and enjoy your family.

(Jury excused for the evening recess.)

(Proceedings continued in open court out of the presence of the jury:)

THE COURT: So tomorrow we start with the remainder of her testimony, the Pilliods, and then what's happening with this, is it Mr. Mills or Dr.?

MR. ISMAIL: It's Mr. Mills.

THE COURT: Is he the economist?

MR. MILLER: He is.

MR. ISMAIL: So Mr. Mills had two components of his testimony, one of which we've resolved by stipulation, you need not address, that's the net worth issue. The parties have covered that part.

The issue that he's being called for tomorrow relates to his calculation of medical expenses for Mrs. Pilliod. And the issue, as we've been sort of sounding this alarm for a while now with respect to Mr. Mills is that under California law, of course, for both past or future medical expenses, it has to be based on what has actually been paid by the plaintiff for the drug. I don't think there's any dispute under Howell and its progeny.

Mr. Mills, in his report, uses a basis for medical expenses that is a one-page printout from the Internet from drugs.com, which on its face says this

does not apply -- this price does not apply to anyone who has insurance. It's sort of the marked-up rate.

So we've been asking for quite some time what -- how much of Mrs. Pilliod actually -- how much has been paid either by her or on her behalf for her relevant treatment which informs Mr. Mills' calculations.

And we've never gotten an answer. There are no documentations that show what she has paid for her Revlimid and indeed we know for certain it's not the number that Mr. Mills is using in his calculation.

So we are --

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THE COURT: This was all the subject of a motion in limine; right?

MR. ISMAIL: It was. And if he can lay the foundation -- and Your Honor has said several times

Howell says what it says, and he has to give a number that's compliant with the law.

THE COURT: Right.

MR. ISMAIL: So here we are the day before.

We still don't have the number of what Mrs. Pilliod's -what cost --

THE COURT: Is there a number, Mr. Miller?

MR. MILLER: Your Honor, there is a number.

\$21,000 per month. California law does not say in the

future we have to deduct or figure out what she might be able to have paid. That's not California law. That's why Your Honor ruled and denied their motion in limine.

2.

To be clear, here's the evidence, Your Honor. In the past -- she needs Revlimid to stay alive.

Dr. Rubenstein says it and Dr. Nabhan says it. And no one disputes it. It costs \$21,000 a month. In the past, she has had insurance pay a lot of it, some charities have paid some. And we've stipulated past medical expenses. We have a stipulation that the parties have worked out.

This is future testimony. And we don't know in the future if she'll have health insurance, if health insurance will cover it, if charities will continue to help her. That's why under California law,

Your Honor -- and the Court's already ruled on this, and I don't know what they're talking about asking for more information. The Court has already ruled and that's

California law. In the future, we don't deduct for what insurance she might have in three years. That's not the law.

MR. ISMAIL: We're not asking for them to deduct what insurance has paid. We're asking them to base any medical expenses calculation on what is actually -- what the drug costs her, not a theoretical

person.

THE COURT: I understand what you're saying.

So are you making an assumption, Mr. Miller, that she's not going to have insurance and therefore the gross amount or whatever the posted amount -- you know, the maximum amount, which is, you know, \$21,000 is a fair estimate of what her expense would actually be?

MR. WISNER: It is. Dr. Nabhan has testified at deposition, they queried extensively on it, that it's \$21,000 a month.

MR. ISMAIL: Doctor -- please finish.

MR. MILLER: Thank you.

And he's backed that up with information from his own practice.

THE COURT: Well, I don't doubt that -- you know, we all know that the stated amount for drugs is very high and that what people actually pay is generally something else depending on what their circumstances are. So I'm not suggesting that \$21,000 isn't that published amount by whoever that drug maker is.

I think Mr. Ismail's concern is a different one, I guess, that in realty she's not going to pay that because she is covered by insurance at this point.

So my question was going to be: So is there any evidence -- so what is her insurance situation?

Which is, you know, some people are provided insurance. I will have lifetime insurance because I have the job I have currently. So if anybody is asking what I'm going to pay, well, it would be based on whatever my insurance -- because I'm going to always have insurance. I know that.

MR. MILLER: And you've earned it, Your Honor.

THE COURT: No, no, what I'm saying -- not my point.

My point is simply that some people know they'll have insurance because she receives insurance through Mr. Pilliod's job or he's retired and so their insurance and benefits are set and therefore they know that they're not going to pay the rack rate for Revlimid, if that's what it is.

So I'm just asking: Is all that information at play? Is it not in play?

MR. MILLER: I don't think it's in play. I think she does have health insurance now. Certainly not trying to suggest otherwise. She has charity that is helping out with the difference.

But in the future, that's not what California law says, and we briefed it and I thought the Court ruled. We can go back to it, but I need to get the case out. I didn't know we were going to argue this until

about 10 minutes ago. 1 THE COURT: I just really love the way you 2 3 quys do --MR. MILLER: Maybe tomorrow morning when I 4 have my briefs. 5 THE COURT: Oh, no, we will not be chatting at 6 9:20 tomorrow morning. 7 Let me go back and reread Howell, for one 9 thing. I think it's important to go back and reread Howell. 10 MR. ISMAIL: In fairness, Your Honor, 11 12 Mr. Mills, we got notice yesterday at 9:00 o'clock last 13 night that he's coming on Thursday. So he was not on this week's agenda. So that's why it's coming up today. 14 And, Your Honor, the Howell, of course, is the 15 16 initial case. Corenbaum was the one that applies Howell 17 to future medical expenses. So Howell was retrospective and -- on the facts of that case, but it's been applied 18

And it is -- Mr. Mills was asked at his deposition --

to future.

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THE COURT: Give me the cite because I don't have all those Sargon papers. I have to get them offline because I had to get rid of some of those papers.

1	MR. ISMAIL: Do you want the case cite?
2	THE COURT: Yes, give me the cite.
3	MR. ISMAIL: One moment. So Howell is 52
4	THE COURT: I know what that is. Just the
5	other case.
6	MR. ISMAIL: Just a second. 215 Cal. App. 4th
7	1308.
8	THE COURT: 1308.
9	MR. ISMAIL: Yes.
10	So it remains the case, Your Honor, that we
11	asked Mr. Mills what his factual predicate is for using
12	that amount. He says, "I took it entirely from
13	Dr. Nabhan." He's never spoken to Dr. Nabhan. He just
14	read his report.
15	We take Dr. Nabhan's deposition.
16	We say: Where does this number come from?
17	He says: I went out and I Googled it.
18	We said: Do you have any information as to
19	what Mrs. Pilliod is actually what it's actually cost
20	her?
21	He says: I have no idea.
22	He doesn't know. We've never seen an
23	insurance explanation of benefit. We've never seen a
24	bill that's been submitted on her behalf for Revlimid.
25	We've never seen the price paid for her. So this is

1 literally the rack rate from the Internet which both --2 THE COURT: There must be something based on 3 what's already been paid --(Simultaneous colloguy.) 4 MR. ISMAIL: That's what we asked for and 5 6 we've never been provided. MR. MILLER: Well, that would go to the 7 weight, Your Honor. 9 THE COURT: No, I'm talking about you're 10 asking for both past and future medical expenses. I'm just asking in terms of asking for the past, the 11 12 number, the hard number of what's already been, that 13 information wasn't exchanged? Or was just a number I don't know. 14 given? I'm asking. 15 We stipulated on the past medical MR. MILLER: 16 expenses. We have a stipulation. 17 MR. ISMAIL: We stipulated on the past which doesn't include Revlimid. And they've stipulated --18 THE COURT: It must include Revlimid because 19 20 if she's actually paid -- or she's never paid anything 21 for the Revlimid to date so that's not part of it. 22 MR. MILLER: Exactly. THE COURT: 23 I see. 24 MR. MILLER: And Dr. Nabhan isn't grabbing this off the Internet. He's published with 25

Dr. Rubenstein on Revlimid. He prescribed Revlimid for years and he backed it up by looking at drugs.com.

This is not a wild number. This is a solid number. They're just trying to argue future expenses.

And I think Your Honor hit the nail on the head. If you read those cases, I think the Court will see what we already talked about before, it doesn't apply in the future.

If they want to make some argument about maybe you'll get a charity to pay in the future, that's fine. But they can't argue that future expense isn't legitimate, isn't based on sound science -- isn't based on expert testimony. She needs --

THE COURT: So there seem to be two issues, one is what the actual rack rate is which sounds like it's not really that much of an issue because it's 20-, 21,000. I mean, it's got to be somewhere in that number. I mean, I know it's not 5- and really you're saying 21-. It's a lot of money, it's a whole lot of money.

But the whole lot of money isn't so much in dispute as whether or not the whole lot -- the rack rate is actually the number on which the future should be based.

MR. MILLER: I think two years ago it's

18,000. It's gone to 21,000. We haven't even added -- I don't think we have an inflation figure, I can go back and look.

But, anyway, I think the Court ought to read those two cases, I ought to read those two cases, take five minutes, not 25, tomorrow morning and look at it.

MR. WISNER: I think there's an important legal point here, right, and you'll see in the briefing. Part of her expenses are actually paid for by the drug company. And every year she actually has to -- she'll testify to this before Mr. Mills takes the stand. So the foundation will be laid through her testimony.

And every year she has to reapply, and the drug company says yes or no. And if they say no, then she's in the hole quite a significant amount of money. I don't know the exact number. She'll be able to testify to that.

Whether or not her insurance -- you know, things change with insurance coverage all the time. They could -- she could lose it or whatever. And I think the reason for that is that under California law, when you look into the future you can't assume insurance or charity, you have to assume the full cost.

Now, if her drug company that's giving her or the insurance company wants to say, hey, pay us back

since you got paid for future damages in your thing, they can bring a lien against her judgment. But it's not a point of predicting future damages. That's how it works. So I just wanted to bring that out as sort of the problem.

THE COURT: Let me just go back and look at the cases.

MR. MILLER: Sure.

THE COURT: And we'll talk about it at 8:30.

MR. WISNER: Sounds good.

MR. EVANS: Your Honor, I just want to put one thing on the record before I forget.

We agreed, I think, with counsel that the playing of the treating physician depositions, I think we stipulated that's okay without having to worry about subpoena or not subpoenaing.

I just don't want to be in a position where after we play the treating physicians that the plaintiffs are going to play this week, that when we go try to play some, you know, downstream, there's going to be an issue as to us not having subpoenaed them or whatever.

So I just want to make sure we have on the record that the parties agree that the playing of the treating physicians is going to be okay for both sides.

MR. WISNER: That's correct. And just to be clear, if you're talking about -- I think we're talking about the same thing. If you want to play a treating physician depo that we haven't played in our case in chief in yours --MR. EVANS: Yeah. MR. WISNER: Yeah, that's correct. THE COURT: Thank you. See you tomorrow morning. (Proceedings adjourned at 4:27 p.m.)

1	State of California)
2	County of Alameda)
3	
4	I, Kelly L. Shainline, Court Reporter at the
5	Superior Court of California, County of Alameda, do
6	hereby certify:
7	That I was present at the time of the above
8	proceedings;
9	That I took down in machine shorthand notes all
10	proceedings had and testimony given;
11	That I thereafter transcribed said shorthand notes
12	with the aid of a computer;
13	That the above and foregoing is a full, true, and
14	correct transcription of said shorthand notes, and a
15	full, true and correct transcript of all proceedings had
16	and testimony taken;
17	That I am not a party to the action or related to a
18	party or counsel;
19	That I have no financial or other interest in the
20	outcome of the action.
21	Dated: April 17, 2019
22	^.
23	- Kelly Shainline
24	Kelly L. Shainline, CSR No. 13476