Page 1	Page 3
CIVIL DISTRICT COURT PARISH OF ORLEANS STATE OF LOUISIANA  CAJUN CONTI LLC, CAJUN CUISINE 1 LLC, and CAJUN CASE NO. 2020-02558 CUISINE LLC d/b/a OCEANA GRILL DIVISION M-13 VERSUS CERTAIN UNDERWRITERS AT LLOYD'S, LONDON  * * * * * * * * * * * * * * * * * * *	FOR THE DEFENDANT:  PHELPS DUNBAR LLP (BY: ALLEN C. MILLER, ESQ.) 365 CANAL STREET, SUITE 2000 NEW ORLEANS, LOUISIANA 70130 (504) 566-1311 millera@phelps.com  PHELPS DUNBAR LLP (BY: VIRGINIA Y. DODD, ESQ.) (BY: KEVIN W. WELSH, ESQ.) 400 CONVENTION STREET, SUITE 1100 BATON ROUGE, LOUISIANA 70802 (225) 346-0285 ginger.dodd@phelps.com kevin.welsh@phelps.com  ALSO PRESENT:  TIFFANY THOMAN ALLISON STOCK, PH.D.
Page 2  1	Page 4  INDEX  PAGE  STIPULATION

	Page 5	Page 7
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2	STIPULATION	LEMOLE MOTE, M.D., TH.D.,
3	IT IC CTIDLII ATED AND ACREED I 1	30/1 South Wayne Drive, Chandler, Arizona 83249,
4	IT IS STIPULATED AND AGREED by and among	having been first duty sworn, was examined and
	the parties that this deposition is hereby being	testified as follows.
5	taken for discovery purposes and for any and all	EAAWINATION
6	purposes authorized under the Louisiana Code of	BY MR. MILLER:
7	Civil Procedure.	Q. Good afternoon, Dr is it Moye or
8	All formalities, including the reading	8 "Moye"?
9	and signing of the transcript by the witness, are	<sup>9</sup> A. Moye. Thank you.
10	hereby waived.	Q. That's okay. Thank you. Good afternoon.
11	Except as provided in Article 1455,	Nice to see you.
12	objections are considered reserved until trial or	A. Good to see you, sir.
13	other use of the deposition.	Q. I know you have given a deposition before,
14		right, Doctor?
15		A. Yes, sir.
16		Q. Okay. So I'm going to try to be quick on
17		all the preliminaries, but I'm going to ask you a
18		series of questions. You're going to give me
19		answers. The court reporter is taking everything
20		down. We ask that those answers be verbally so that
21		she can accurately take down your testimony. Is
22		that fair?
23		A. Yes, it is.
24		Q. Okay. Oftentimes, I think my questions are
25		articulate and well-phrased, and they are not. If
		articulate and wen-phrased, and they are not. If
	Page 6	Page 8
1	THE REPORTER: Good afternoon. My	in fact you do not understand what I'm asking,
2	name is Yolanda Pena. I am a nationally	please ask me to rephrase it, and I'll be happy to
3	certified court reporter with Baton Rouge	do so within the best of my abilities.
4	Court Reporters.	However, if you answer, I'm going to assume
5	At this time, I will ask counsel to	that you understood what I was asking. Is that
6	identify yourselves and whom you represent	fair?
7	and agree on the record that there is no	A. It is.
8	objection to this court reporter	Q. Okay. If we ever need to take a break,
9	administering a binding oath to the witness	9 please allow me I imagine we might go a little
10		while because your report is so extensive. We can
11	via remote videoconference.	take a break at any time. I only ask that if we do
12	We'll start with the noticing	take a break at any time. I omy ask that if we do
13	attorney.	take a break and there is a question pending, we
14	MR. MILLER: Allen Miller and	answer that question prior to the break.
15	Kevin Welsh on behalf of the defendants.	A. Tunderstand.
16	And we have no objection.	Q. Okay. Why don't you give me your fun hame
17	MR. ALVENDIA: Rico Alvendia,	and address, please.
	Dan Davillier, Jennifer Kuechmann,	A. Way I ask a question first, just a
18	Jennifer Perez, Matt Sherman on behalf	technical question:
19	of the plaintiff. And we have no	Q. 1es, sir.
20	objection.	A. My screen my screen has been reversed so
21	THE REPORTER: And, Dr. Moye, I'll	that the zoom component is over on the right, and
22	swear you in now.	I'm seeing a looks like a screenshot of a
23	///	computer screen that is Phelps Dunbar LLP.
24	///	Q. Yes, sir. And we can change that right
25	///	now. The throughout the course of the
		I and the second

Page 9 Page 11 1 deposition, that may happen because we'll be sharing Q. And obtaining that -- the Ph.D. was your a screen with you to show you documents. We're not last -- and I know there are multiple educational at that point right now, so I'm going to ask that components to your career. But that was your last Kevin remove that screen so you can then see all of formal schooling? 5 the other participants. Okay. A. Yes, sir. 6 A. And I think --Q. Okay. What was the master's degree? What 7 O. Is that better? discipline was that in? 8 A. There was a question pending. A. Mathematical statistics. 9 Q. Sorry about that. Okay. Q. Tell me what exactly is mathematical 10 10 The question was for you to state your name statistics. 11 11 and address for the record. A. Sure. It's mathematical science of the 12 12 A. Sure. First name is Lemuel, L-e-m-u-e-l, theory underlying the analysis of data, so that 13 13 entails an understanding of probability, I would say last name M-o-y-e. 14 14 completely, but nobody understands probability Q. Now, Doctor, one more preliminary question. 15 Are you on any prescription medication today? 15 completely. So an understanding of probability and 16 16 an understanding of statistical inference. A. No, sir. 17 17 Q. So there's nothing that would prevent you Q. All right. Now, so unfortunately for you, 18 18 Doctor, I was an English major, so... from answering my questions truthfully? 19 19 A. I appreciate -- I appreciate good diction 20 20 Q. Okay, great. Why don't you, just and articulation. 21 21 generally -- and we're going to get to your CV and Q. So we're going to spend quite a bit of time 22 22 kind of walk through that in detail, Doctor. But having you educate me on some of these technical 23 23 why don't you give me your educational background, terms, and I apologize in advance for that today. 24 24 A. Of course. please. 25 25 A. Sure. Where would you like to start? Q. I think I understand probability. But tell Page 10 Page 12 1 Q. Your -- well, start with your undergrad and me statistical inference, what exactly that is. 2 work through your professional degrees. A. Of course. So statistical inference is the 3 A. Sure. So very briefly, I graduated with a 3 process by which one can collect data from a sample 4 BA in mathematical sciences from the Johns Hopkins and attempt to attribute its findings to a larger University in 1974. Graduated from Indiana population from which the sample was obtained. 6 6 University School of Medicine in 1978. Then Q. And I've gone through your CV. So the 7 7 internship in Methodist Hospital in 1978 to '79. import of that is seen heavily throughout all of 8 Q. What was your internship in in '78 and '79? vour clinical trials at work? A. It's a rotating intern. Actually, the A. Yes, sir. 10 10 technical term was flexible internship. It's a Q. Okay. And all of us, I think, are learning 11 11 rotating internship. about clinical trials, particularly with respect to 12 12 Q. Which means that you participated in Pfizer's announcement on vesterday. 13 13 various disciplines in medicine? So when I say "clinical trials," that's the 14 14 A. Essentially, all of the major disciplines practical application of the statistical inference 15 15 in medicine. So general medicine, surgery, where you take a subset of individuals, you test 16 16 endocrinology, cardiology, emergency room of course, them so that you can then extrapolate that to the 17 17 among others. greater population? 18 18 Q. Okay. A. Well, it's -- epidemiologist distinction. 19 19 A. Then I went to Purdue University. Got a I would describe it as the following: A clinical 20 master's degree at Purdue University, and I got the trial is a research environment where --21 21 master's degree I think in 1981. Not really sure Q. I'm sorry. 22 22 about that but approximately 1981. And then went to A. All of the Bradford Hill causality tenets 23 23 University of Texas, where I attained a Ph.D. in are incorporated in the design of the study with the 24 what was called then biometry. But it's really 24 exception of strength of association. Strength of 25 biostatistics, and that was in 1987. 25 association is obtained from the trials data. And

Page 13

the combination of all of those tenets leads to a conclusion as to whether the relationship between the exposure or the treatment and the disease is causal or not.

- Q. And that is what you would describe as a clinical trial?
  - A. Yes, sir.
- Q. Okay. And the Bradford Hill causality, what -- tell me what that is.

A. Sure. It's a collection of guidelines that were established by Sir Austin Bradford Hill. I think they were published in 1965. They essentially describe the thought process a reasonable epidemiologist goes through in determining whether an exposure is -- causes a disease or whether an exposure is merely associated with the disease.

Q. Okay. Let's pull up your curriculum vitae.

MR. MILLER: We're going to mark that as Exhibit 1.

(Exhibit No. 1 was identified.)

MR. MILLER: Kevin? There we go. BY MR. MILLER:

Q. I'm going to walk you through, Doctor, some of the -- your profession experience, certifications, and licenses. And where I have

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Page 16

- A. Sure. So we talked about what mathematical statistics was, and I won't repeat that. Biostatistics is the application of mathematical statistics to health science research.
- Q. And when you say "health science research," what areas does that include?
- A. Actually, that's very -- that's a very broad swath. So health science research includes research in any of the medical fields. So as example, research in cardiology, research in pulmonology, research in orthopedics. So all of the clinical fields, and in addition it includes research in nursing. It includes research in the economics of healthcare delivery. It includes research in quality of life. Any area that involves healthcare delivery is also considered part of healthcare research. So it's very broad.
- Q. Okay. What particular areas have you specialized in within the application of biostatistics?
- A. Well, I have -- to be fair, to answer your question quickly, most of my research has been in cardiology. However, I have worked in other fields as well, including pediatrics, including alternative medicine, as they are distributed through my

Page 14

questions, I'll just -- I'll pose those questions at that time. Is that okay?

A. Yes, sir.

- Q. Okay. Let's look at your professional experience. And the first thing says you're a professor of biostatistics with tenure at the University of Texas. Now, you made a distinction earlier when you said you had a biometry, but it's really called -- it's biostatistics now?
- A. Yes. The language of biostatistics, for better or for worse, evolves. So when I started the school, its program in biostatistics as well as the other programs in the nation were not called biostatistics. They were called biometry.
  - O. Biometry.

A. And without the -- without to -- diverge too much, biometry and biometrics turned into the use of body measurements to indicate something about the person. So the rate of which -- the rate at which irises dilate or -- or constrict, fingerprints, that all became -- lip movements, that all became the science of biometry. And so our field took the name biostatistics.

Q. All right. So tell me exactly what is biostatistics.

publications.

- Q. Okay. The next entry under your professional experience is that you were director of -- director in coordinating -- director at the Coordinating Center For Clinical Trials at the University of Texas?
  - A. That's correct.
  - O. Okay. What exactly was that position?
- A. So the coordinating -- can I tell you first what the Coordinating Center was?
  - O. Yes, sir.
- A. Okay. Sure. So the Coordinating Center was a facility that included faculty members from several different disciplines or departments in public health. It included biostatisticians, epidemiologists, sometimes healthcare specialists, and sometimes, depending on the trial, diabetologist or hypertension specialist or a specialist in a retinal disease or prematurity. We all work together to design, execute, and -- and analyze clinical trial results.
- Q. Okay. And as -- you were the director for -- from 2009 to 2010?
  - A. Just a year, yes.
  - Q. Okay. Did you participate in this group

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### beyond being the director?

A. Sure. Thank you. So I was a member of the Coordinating Center for Clinical Trials the entire duration of my faculty life at the school of public health. It's just this one year I was asked to step in as director.

Q. Okay. The next entry is secondary appointment of epidemiology -- secondary appointment in epidemiology.

A. Yes, sir.

#### O. What exactly does that mean?

A. Well, okay. Let me answer that in a negative -- negative, if I may, first. It does not mean I was a member of the department of epidemiology. I want to be clear about that. I was a member of the department of biostatistics. I have never been and have never said I was a member of department of epidemiology.

Having said that, I have been working closely with epidemiologists since I started in -at the school as a faculty member in 1987, and I was not the only one. In fact, faculty members were encouraged to work with other faculty members in different disciplines then than departments.

Q. Okay. So let me go back to the prior entry

there are simply degree programs.

Q. Do you have a degree in epidemiology?

A. No, sir.

### Q. Okay. There is such a degree as epidemiology, correct?

A. To be clear, there are masters' degrees and Ph.D.s in epidemiology, but I have enough degrees. I don't need another academic degree.

Q. I won't disagree with you on the number of degrees you have. You do have enough.

While you were actively working at the University of Texas, did the -- were there any members of the epidemiologist department that considered you to be an epidemiologist?

A. Oh, for sure, yes.

### Q. Okay. Who were they?

A. I would say Milt Nickerman, Rick Shekelle. These were senior members of epidemiology there. Darwin Labarthe, also a senior member of epidemiology there. Unfortunately, Rick just died, I think, three or four months ago.

O. Okav.

A. We worked on -- I'm sorry. I'll stop

Q. Yeah, I'll get to -- we'll come back to the

Page 18

in your CV. As director of Coordinating Center For Clinical Trials, I imagine there were epidemiologists in the Coordinating Center For **Clinical Trials?** 

A. Yes, sir.

### O. Okav.

A. Yes, sir, just as I served with epidemiologists in their primary work as a biostatistician and a fellow faculty member. There was a lot of cross-pollination.

### Q. But you are not an epidemiologist?

A. No, I wouldn't say that. I would say I am an epidemiologist. I simply am saying to you I was not a member of the epidemiology department, but I am a practicing epidemiologist. I am an applied epidemiologist.

### Q. Is there a certification for that?

A. Is there a certification for applied -- no, I don't believe there is.

### Q. Okay. Are there any specific requirements that the Texas board would require you to have to identify as an epidemiologist?

A. I understand your question. There's no Texas board that certifies biostatisticians or certifies epidemiologists. There are simply --

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work that you did while there.

The next entry -- so again, let me go back to secondary appointment in epidemiology. What does the distinction "secondary appointment" mean?

A. It means that I'm affiliated with and do work in the -- with epidemiologists. However, I am not involved in the administration and the -- well, I'm not involved in the administration of the department. So for example, things I did not do: I did not attend meetings where they talked about what their curriculum was going to be. I did not attend meetings where they talked about when they were going to give their qualifying examination. Those were important matters. Those were administrative

What I was involved in was advising students in epidemiology, helping to teach classes in epidemiology, working with epidemiology students who took my classes, including epidemiologists on my grants and work and being part of the grant work of other epidemiologists. And of course we publish together in epidemiology journals.

Q. Okay. If I went to the University of Texas's description of its faculty members, would they describe you as having a secondary appointment Page 21

### in epidemiology?

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A. I -- I have not looked. I would be surprised if they did because the school has become quite siloed. And at this point, you are part of one department or another. But that is a -- an intellectual movement that flows one direction or the other. Sometimes they move to be more siloed. Sometimes they move to incorporate collaboration. So at this point, I'm going to say I don't know what they would say.

### Q. Okay. Well, then let me ask this: What encouraged you to -- so this term "secondary appointment," was that something you created on your

A. No. No, it was not. It was encouraged that we all -- this is now in the 1980s. It was encouraged that we all be part of other departments and talk about it. And -- and since I took my work in epidemiology seriously, it was appropriate for me to put that on my CV.

### Q. Again, now, it was your decision to put the secondary appointment epidemiology on your CV?

A. Sure. If you want to say "epi," I understand what you mean, rather than going through "epidemiology."

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### Q. It says 1986 to the present. So are you still consulting with Baylor?

Page 23

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A. All right. That's my fault. I should have put retired there as well. So that -- that really ended in 2019.

### Q. Okay. And what -- what's the distinction there, consulting clinical biostatician as it pairs to what -- compared to what you were doing at the **University of Texas?**

A. Sure. So at the University of Texas I worked with senior epidemiologists, senior diabetologists, hypertension experts, both locally and nationally, to design, execute, and analyze large clinical trials. There is a senior level of expertise in that work, and I was pleased to be part of it. The consulting work with -- with -- where are we? With...

### Q. Baylor.

A. Baylor. Thank you. Baylor. So the work with Baylor was local, and it was of a lower caliber in that the investigators I worked with had no real experience with epidemiology and no real experience with biostatistics. They simply knew that they had collected data, and they were interested in having that data appear as an abstract or appear in a

Page 22

But yes, it was. But a couple of things. First, it's my decision to put anything on my CV, and many faculty members would put this as well if in fact they were involved with another department.

### Q. Okay. I'm going down further, and I'm trying not to rehash.

If, for example, you were an associate professor in one of the areas in which we've already discussed, I'm going to skip over those things.

A. Yes, sir.

### Q. Okay. It says that you were consulting clinical biostatician at Baylor. Is that any different from what you were doing at University of Texas?

A. Of a kind, yes. And I'll try and keep this short. University of Texas -- Baylor, University of Texas, Methodist Hospital, they were all a part of the Texas Medical Center. Just as I was encouraged to collaborate with other departments within my school, such as epidemiology, I was also to encourage -- encouraged to work with other institutions in the medical center. And one of them I worked with was Baylor.

### Q. Are you current --

A. All right. I'm sorry. Start, please.

publication and didn't really know how to organize and analyze the data to move forward to a result that might be publishable.

### Q. Understood.

A. I do need to clarify something, if I may. I realize you may want to ask questions about it.

I do not have -- as I have told you, I do not have a formal degree in epidemiology, but I have substantial training in epidemiology. In medical school we get considerable training in epidemiology, to the point where many physicians think they know epidemiology. I'm not there yet, but you learn -we learned -- my first exposure to epidemiology was in medical school, and it was through the second, third, and fourth years of medical school.

In addition, I took courses as a student at the School of Public Health in epidemiology as I took courses in biostatistics. So I have a good deal of formal training in epidemiology. It -- it's important to have experience, but without formal training experience, can sometimes be making the same mistake repeatedly.

### Q. Other than taking courses -- what year did you attend medical school?

A. 1974 to 1978.

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Q. Okay. Other than courses in 1974 to '78 and then the work during your, I guess, Ph.D. in the School of Public Health?

- A. Yes, sir. As a student, yes.
- Q. Other than those two instances as a student, do you have any formal training in epidemiology?
- A. Beyond the -- attending seminars as faculty member, no.
  - O. Okav.
- A. I have a lot of experience working with epidemiologists but no additional formal training.
- Q. Okay. Let me see. It says also that you were the owner of MediClinic Physicians' Association?
  - A. Yes, sir.
  - Q. What is that?
- A. That was a collection of freestanding clinics in Houston that I joined as a physician. I was a licensed physician in Houston, and I joined and worked with them, and then was asked to be the owner of the physicians' association, which is the collection of physicians who were involved in seeing patients across -- I think it was seven clinics.
  - Q. Now, those clinics, were they primary care

Page 27

academic setting. This is the first that I'm hearing of sort of, like, clinical work as an MD.

How -- tell me generally how much experience you have in actually seeing and treating patients.

A. Sure. Be glad to. I started seeing -- I have -- to answer your question tersely, substantial. I had -- when I graduated in 1978, I got and -- and completed my internship in '78, '79. I was licensed, and I began seeing patients shortly thereafter. I was seeing patients at a clinic. I'm trying to see -- yes, Methodist Healthcare Center, which was associated with Methodist Hospital.

I saw patients five days a week, and these patients were -- primarily they were occupational medicine at the time, primarily occupational medicine. The notion of freestanding clinics really didn't take off until the mid-1980s. So I saw many patients there with occupational injuries.

I also worked at the US steel mills in Gary, Indiana. That was for about a year, but it wasn't very often. However, there I really -- I saw the worst clinical situations I'd ever seen in my life. It was really -- it was very educational and sometimes very tragic.

Page 26

### clinics?

- A. Yes, sir.
- Q. Did any of the, say, seven -- estimated seven clinics have any particular type of specialty that they rendered?

A. No. Each of the clinics was open to the public for any -- any healthcare situation or problem patients in the community, subjects in the community might have.

Now, it didn't stop some of our physicians from having some special expertise that they would share. For example, I had a lot of experience in orthopedics. And so when it came to relatively simple things, like bad sprains and simple fractures, I had no problem diagnosing and treating and following patients with these injuries. And I let the other clinics know so that they could have their patient -- if they were not so comfortable seeing those patients, they could come to see me.

So we did have this expertise sharing, but there was no -- there was no -- there was no policy that excluded one type of patient from a clinic in preference to another.

Q. In your -- lots of the information that we've talked about thus far is contained in the Page 28

- O. At -- this was at the steel mills?
- A. Yes, sir.
- Q. When you say you worked at -- were you a physician on site at the mill, or --
  - A. Yes.
  - Q. -- did you work in a hospital?
- A. I took 16-hour shifts onsite at the Gary steel mills.
- Q. So other than when you initially graduated from medical school and became licensed in Indiana and then the job at the steel mills, when was the next job where you were actually seeing patients?

A. Oh, then things really took off. Then the urgent care concept became accepted, and I began to work at urgent care clinics, first in Indiana and then in Houston, when we were talking about the MediClinic Physicians' Association.

There I commonly worked 30 to 40 hours a week as a student -- 30 to 40 hours a week while I was a student in the Ph.D. program, also seeing

- Q. When did your active medical practice seeing patients end?
- A. For all intents and purposes, it ended in 1992. There was the Katrina event that I took --

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took part in. That would have been, I think, 13 years later. But the practice of medicine for me essentially ended 1992.

### Q. And why was that?

A. Well, it was because up to that point in my career, I had my cake and could eat it too. I was doing research and also was seeing patients, but it got to the point where I could not continue to do both satisfactorily. So I had to choose, and I chose to go into research.

### Q. Okay. In 1992 you were already employed with the University of Texas?

A. Yes. At that point in 1992, I was a faculty member.

Q. Okay. All right. I'm going to skip down to page 2 and as -- as decorated as your honors are and impressive as they are, I don't think that I need to go through them, so I'll skip that section.

A. Okay.

Q. Let me ask you about the Journal of Clinical Epidemiology. Well, let's back up. Before we get there, you have a section on your CV that's entitled "Journal Reviewer."

A. Yes, sir.

Q. So tell me exactly what that means.

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A. I think that's fair. As I sit here, I'm not sure which article I submitted to PharmacoEconomics, but I -- but I was asked to be a journal reviewer for that one.

Page 31

### Q. Okay. To become a journal reviewer you have to be invited?

A. Yes. sir.

O. Okav.

A. Well, at that time you did. At this point, the need for reviewers is so great, perhaps the invitation isn't necessary anymore. But when I was active -- when I was active, you had to be invited.

### Q. Now, are you currently a journal reviewer for the list of journals that are on your CV?

A. Now that I've retired, I have put journal reviews behind me. So the -- the question -- the answer to your question is no.

### Q. When did you stop becoming a journal reviewer?

A. When I retired.

O. So in 2019?

A. Yes, sir.

Q. Okay. The Journal of Clinical Epidemiology, which is the first entry there, did

Page 30

A. Of course. Of course. One of the linchpins of medical research is the review process. So a manuscript is submitted to a journal. As many manuscripts are submitted to journals, journals can't publish them all, so journals have to make a determination as to which ones are worthy of publication. So they rely on a collection of authors who publish in their journals, who have already published in their journals, to get their assessment and impression of the article. So that's called peer review.

We are -- as authors and as fellow researchers, we're peers. We review the manuscripts and give the editor of the journal an assessment of whether this manuscript is worthy for consideration for publication.

Q. Okay. And so when you describe yourself as a journal reviewer and then you list these various -- these various journals underneath?

A. Yes, sir.

Q. Okay. And so you have acted as -- well, let me back up.

You have submitted articles then to all of the journals that are listed under your journal reviewer title?

Page 32

you -- did you have occasion to publish a work in the Journal of Clinical Epidemiology?

A. Well, I certainly had occasion to review it. I'd have to look at my CV to see if I actually published in it. I don't think it will take very long, but I can check.

Q. We'll get to that. I guess, in order to kind of short-circuit it, initially when I asked for the definition of a journal reviewer, I think you said the journals would ask a collection of individuals that submitted articles or journals to them to review others.

A. Yes. sir.

### Q. And I just want to ensure that I have the right definition. I don't want to assume that you published in each of these journals.

A. Right. So my definition may be too narrow. While they may or they are -- of course they know the work of the people who submitted to their journal because somebody reviewed it. But I would think that if somebody had articles published in the Journal of the American Medical Association in epidemiology, they might also be asked to be a reviewer for the Journal of Clinical Epidemiology. It's a relatively small universe of editors and

Page 33 Page 35 1 1 reviewers. Q. (Indiscernible.) 2 Q. Okay. Is it fair to say that the journals A. I'm sorry. Again, I stepped on your 3 that you've listed are generally accepted as question. Go ahead. 4 professional and reliable sources of information in Q. (Indiscernible.) 5 the fields in which they publish? A. I would get an occasional request to review 6 A. I would describe them as essentially for Lancet. 7 reliable and high quality. Now, there are -- there O. And what -- some of these are 8 are always exceptions. The good -- a good journal self-explanatory. But what's the subject matter of 9 of medicine will make a mistake and publish a bad Lancet? 10 10 article sometimes, but by and large they are A. Lancet is the European version of the 11 11 reliable and high quality. American -- of the Journal of the American Medical 12 Q. Okay. The second entry underneath "Journal 12 Association. It is their premier general medicine 13 13 Reviewer" is Journal of the American Statistical iournal. 14 14 Association, Biometrics. Is that -- and then Q. Okay. The last one? 15 there's Biometrics behind it. Is that one journal? 15 A. Atherosclerosis? 16 Or is Biometrics a journal in -- in and of itself? 16 Q. Yes, sir. Was it that? 17 A. They are two separate journals. 17 A. That's a specialty journal in cardiology. 18 Q. Okay. So there's a journal entitled simply 18 Cardiology is a broad field. Atherosclerosis is a 19 19 **Biometrics?** pathophysiological process, which has of course 20 20 A. Right. garnered a great deal of an attention over the 21 21 Q. And there is a journal entitled Controlled decades, and they have a journal associated with 22 22 **Clinical Trials?** 23 23 A. Yes, sir. It's a political complexity that Q. Okay. Now, the next entry is a book. 24 2.4 now has turned into two different journals. One is You're -- you're a book reviewer just for that -- is 25 25 contemporary clinical trials. I'm not sure I follow that one book, Springer? Page 34 Page 36 1 the politics of it, but when I was reviewing, it was A. Unless you scroll down and I'm missing -- I 2 2 can't scroll with you, so I have to rely on you. called Controlled Clinical Trials. 3 Q. Okay. Then there on the other side, The Q. If we scroll to the next page, we get to 4 the current -- current funded research. American Journal of Epidemiology, did you publish in A. Okay. So I -- I was just asked to that journal? 6 review -- book review for Springer. A. I'd have to look and see. Q. And who is Springer? What do they publish? Q. Okay. Do you know how long you acted as a A. Oh, Springer is a publisher that focuses reviewer, or how often did vou review things for the on -- well, actually, Springer evolved -- has American Journal --10 10 expanded over the years. So it was primarily A. I'm sorry. I didn't mean to step on your 11 11 mathematics, became statistics and probability, question. Go ahead, please. 12 12 biostatistics, and I presume they continued to Q. How often you reviewed on behalf of the 13 13 American Journal of Epidemiology? 14 14 Q. Okay. Now, there were no dates on any of A. I can't tell you how often I reviewed for 15 your journal reviewers, and nor is there a date on 15 any of these, but I can tell you that it seemed like 16 16 every two or three weeks I was getting the this Springer. Is there a reason why you didn't 17 identify specific dates in articles that you 17 opportunity to review for one or the other. 18 actually reviewed? 18 Q. Okay. And would that be the same for the 19 19 New England Journal of Medicine? A. Well, there's two reasons. Number one, 20 20 it's not the custom in my area to actually put that A. Yes, sir. 21 kind of detail in; and number two, I didn't have the 21 Q. The American Family Physician? 22 time to log that. And if I would have -- yeah, I 22 A. I don't know so much about that. 23 just didn't have the time to do it. So nobody ever 23 O. Okay. What about -- was it -- how do you 24 asked me to do it, so I'd rather put my time 24 pronounce that? Lancet? 25 someplace else. 25 A. Lancet, yes.

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Q. When was the last time you updated your CV?

A. This is dated October, I think, 2020. I think it is.

O. The next section is "Current Funded Research," and as I reviewed it, most of it -- well, there are two areas in the field of cardiovascular and then one in stroke. Would you -- so did you do any funded research in the field of epidemiology?

A. Yes. I would describe all of this -cardiovascular stem cell network, the acute stroke program for sure, Dr. McCormick's program for the Hispanic health research, it's clinical and translational science award, all of that is epidemiology.

Q. (Indiscernible.)

A. Go ahead, please. Go ahead. I'm sorry.

Q. I interrupted you. Go ahead.

A. Okay. And I would say that this is really just the most recent. Bio-funded research goes back to 1987 when I was funded on hypertension programs and the development of statins, and that was all epidemiology and biostatistics.

Q. Let me make sure -- and I'm going to ask you one more time for my clarity, and I apologize if I'm repeating myself.

the individuals that you are compiling data in relation to. So for instance, I'm looking at some of your publications, and there is a publication No. 10.

A. Okay.

Q. "Amino acid composition of preterm and term breast milk during lactation," that is a publication that, as you described it, falls under the umbrella of epidemiology?

A. I would say that's one of the examples where it did not because I know -- even -- let me put it this way. There is the overall research, and there is my role in the research.

The overall research, I would certainly assert that if you look at the overall research, design, and execution, it is all -- all has epidemiology at its core. All of them do. Regardless of my role, all of them do.

Having said that, I was not involved in the overall design of several of my publications. And one of them is the publication you pointed out, No. 10, where I was asked to come in and actually -essentially to analyze the data.

And when I come in and just -- and given a dataset, told to analyze it, and to provide some

Page 38

So cardiovascular, stroke, we'll get to some -- you know what, let me use this.

So cardiovascular, stroke, hypertension issues with respect to low birth rate, lactation, sex bias, cholesterol, you would put all of the work that you did in all of those areas under the umbrella of epidemiology?

A. Almost. It sounded like you were looking at some of my publications.

O. I am.

A. There are a few of my -- there are a few of my publications where I function only and solely as a number-cruncher, and that I can't -- I cannot say is -- in all honesty, is epidemiology.

But when I am involved in the design and the execution and the analysis of a program, then that certainly must involve epidemiology.

Q. So as -- and I just want to make sure I'm clear. Putting the -- putting aside the underlying healthcare issue that you are addressing, the work that you've done throughout your career you consider to be epidemiology?

A. I'm not sure I understand what we put aside. What did we put aside?

Q. We're putting aside the physical ailment of

Page 40

Page 39

result, I don't really consider that epidemiology. It's much too narrow, but somebody must interpret 3 that data, and that interpretation is epidemiology. 4 Does that help?

Q. Yes, very much so.

All right. Well, let's take a look at your publication starting with number -- the first one.

A. Right.

Q. Tell me what you did there.

A. Of course, yeah. This was a -- this is my very first publication ever, and it involved microbiology.

Essentially, I was asked to -- the hypothesis was that we -- we knew that patients with cancer -- this is primarily hematologic cancer, leukemia -- commonly required anticancer chemotherapeutic agents in addition to antibiotics. They required the anticancer chemotherapeutic agents because they had cancer. But the combination of the cancer and the effect on the anticancer chemotherapeutic agents on their bone marrow depressed this immune response. These patients were on the antibiotics. I'm just setting the stage.

So patients are on both types -- both classes of medicine. There have been an observation

Page 41

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that patients who were on certain antibiotics seemed to have a better response to the anticancer chemotherapeutic agent, and I was asked, I guess, at this point -- when was this? This was published in '78. I did this research as an undergraduate back at Johns Hopkins in Baltimore.

I was asked to design a study, an in vitro study. I wasn't a physician yet, so I couldn't see patients. But an in vitro chemistry study to see whether if I expose -- whether the exposure of common pathogens -- pseudomona aeruginosa would be one example.

THE WITNESS: And I will spell that for you, Ms. Reporter -- Ms. Pena. I will spell that for you later.

A. Pseudomona aeruginosa was a very dangerous pathogen. It caused a hemorrhagic pneumonia. It's lethal. And I was asked to see whether the bacterium that caused that pneumonia was more sensitive to antibiotics, I think it was ticarcillin and carbenicillin, when they were exposed to cytosine arabinoside and daunorubin. Those are the only two -- only two anticancer chemotherapeutic agents that I remember.

But we were able to show that in fact some

Page 43

Having said that, I will say I've had an awful lot of physics.

### Q. Okay.

A. I can -- I understand kinematics. I understand Newton's laws of forces. I understand free fall. I understand little about astrophysics, but I would not hold myself out as a physicist.

Q. Okay. So this particular publication was again -- you were asked to perform a specific task within the larger scheme of this trial, I guess, so to speak?

A. Well, I would put -- I'd put it this way. I had a much broader role here. This is different than the publication 10 where somebody just gave me data. In publication 1 I was given the problem, and I had to think through what data I need to collect, what anticancer chemotherapeutic agents do I -- should I study, what antibiotics should I study, what different types of bacteria should I study, and to what degree should this all be replicated.

So I was really given a lot more leeway as an undergraduate student. And even though this was published in '78, this work all happened in '73, '74.

Q. Okay. That's -- that's what I thought I

Page 42

of the antimicrobial agents had more killing power when they were exposed in vitro to anticancer

chemotherapeutic agents, even though the anticancer

chemotherapeutic agents did not kill the microbes themselves.

BY MR. MILLER:

Q. So let me ask this. Because in 1978 you were -- you were getting your undergraduate degrees in mathematical science?

A. In '78 I was -- I was graduating medical school.

Q. I'm sorry. Okay. I'm sorry. In '78 you were graduating medical school.

But -- so you mentioned in this publication, or this task that you were asked to do, you did a chemistry study?

A. Yes, sir.

Q. So tell me -- you're not a chemist, though, right?

A. No, I'm not a chemist. But I have had lots of chemistry.

Q. Yeah, I understand. What about physics? I mean, you're not a physicist or --

A. I've had lots of physics. I would not represent myself as a physicist, I would not.

Page 44

was getting at when you said you were doing this as an undergraduate. Although it was published in '78, the work happened while you were an undergraduate in '74 -- '70 to '74?

A. I think -- I think it was 1973 to 1974. It's over two years.

Q. And the underlying -- this kind of goes back to the point I was attempting not so well to make a while ago. The underlying medical issue for this publication was cancer?

A. Yes, sir.

Q. Do you consider the work done with respect to this publication epidemiology?

A. I do.

Q. Okay. That's all I -- that's what -- I'm trying to learn the definition of epidemiology. Well, let me just ask that.

A. Sure.

Q. And I may have asked it previously, but could you give me what your -- the formal definition of epidemiology?

A. Sure. I would say epidemiology is the process by which the true nature of the exposure/disease relationship is deduced.

Q. And under that definition, epidemiology

Page 45 Page 47 1 1 could include any type of disease? medical school -- correct --2 2 A. Yes, sir, yes. A. Yes, sir. 3 3 Q. Okay. Q. -- at the time of publication? 4 THE WITNESS: I'm going to ask to take A. Sorry. Yes, sir, correct. 5 5 a break in five minutes, just to give you a Q. And so at that time, your -- the most of 6 6 your experience would have been in mathematical 7 MR. MILLER: We can take one right sciences? 8 8 now, actually. I'm ready for a break, if A. Well, I -- I can't disagree. Well, let me 9 9 say this. It's hard to know. I had a math science that's okav. 10 10 THE WITNESS: Okay. Five minutes background, but I'm in the -- I'm at the end of 11 11 medical school now. So medical school has tipped 12 12 the balance some, and I've had epidemiology exposure MR. MILLER: Five minutes will work. 13 13 in medical school, so I don't know what to say. But (Recess taken.) 14 14 most of my background is. BY MR. MILLER: 15 Q. All right. Doctor, we were -- when we 15 Q. I got you. The next publication in 1981 16 left, we were going through some of your 16 involved linen in newborn intensive care units be 17 publications. Let's take a look at the second one, 17 autoclaved. Again, in pediatrics --18 which was published in 1979. 18 A. Autoclaved. 19 19 A. Oh, okay. Q. Autoclaved. And this particular 20 20 Q. The "Environmental contamination of publication, what -- was your role task-oriented? 21 21 continuous drip feedings in pediatrics," was that --A. I would think, again, it was not 22 when you say "drip feedings," was that in relation 22 epidemiology, though I learned epidemiology. It 23 23 was -- it was -- my contribution to the publication to breast feeding? 2.4 24 A. No. I -- I'm going to have to refresh my was through data analysis. 25 25 own memory about this one. Q. Okay. Page 46 Page 48 1 1 O. Okav. A. Essentially, the research had already been 2 2 A. No, it was not -- it was not about breast designed, already been executed, and I was asked to 3 3 feedings; it was about the -- it was supplemental analyze it. And this series of publications are all 4 feeding divided -- provided to infants in the the same genre. Because I was known in medical hospitals. school as someone who knew mathematics and 6 6 Q. Okay. And I only brought up breast statistics, so I was commonly asked at the end of feedings because I noticed that there were a number the day to help analyze. I was not asked at the 8 of breast feedings publications that you had -inception of the project to design the project 9 A. I see. because I had really no pediatric expertise. 10 10 Q. Okay. So in order -- and I'm all about Q. -- throughout your life. 11 11 A. But this -- but this one where the concern efficiency, Doctor. So if in fact there's a 12 12 was that -- this was an antimicrobial study. The publication that you contributed to in the area of 13 13 concern was that there might be some contamination pediatrics --14 14 of the drip feeding apparatus. A. Yes. 15 15 Q. Was your contribution to this particular Q. -- it would have been typically data 16 16 publication task related? analysis? 17 17 A. It was simply -- it was simply data A. I guess I would have said this. By and 18 18 analysis. large, any publication that occurred before -- any 19 19 Q. So this was not -- would not have been of the publications up through 12, most of those 20 20 under your definition of epidemiology -- an exercise publications are data analysis. 21 21 in epidemiology? Q. Okay. 22 22 A. Correct. I learned some epidemiology from A. There are a couple of exceptions. Eight is 23 23 speaking with the senior investigators, but my role an exception, and 1 is an exception, but the other 24 24 was really just as a data analyst. ones through -- up through and including 12 are 25 25 pretty much my work as an advanced medical student Q. Okay. And at that time, you were in

Page 49 Page 51 1 1 who knew some statistics. Q. All right. So what we have in 13 is a 2 Q. Understood. And 8 was -- that entry says, journal concerning stroke and hypertension. 3 "Modeling of pharmacology treatment of hypertension, And hypertension being the disease and clinical math and science." Would that have -- would you trials -- or prevention of stroke and hypertension 5 5 have considered that epidemiology? being the exposure? 6 A. I would consider that epidemiology and A. I'm sorry. The last thing you said was 7 mathematics, yes. what? 8 Q. A combination of the two? O. Exposure. 9 A. Yes, sir. A. Oh, right. So let me be clear. This is a 10 10 O. And you did the modeling for this study which provides our intent -- no, it provides 11 11 particular publication? our design to determine whether the treatment of 12 12 A. Yes. This is my idea, my research, my isolated systolic hypertension could reduce -- could 13 13 design, my analysis. prevent -- excuse me, could prevent the occurrence 14 14 Q. And you -- as evidenced by the title, you of stroke, and it occurred -- this was published in 15 did some type of modeling with respect to 15 an epidemiology journal. 16 16 hypertension? Q. Journal -- I see it. Okay. 17 17 A. Yes, sir, with respect to the pharmacologic But, again, I want to go back to your 18 treatment of hypertension, yes. 18 definition, so I can -- because it'll help us move 19 19 Q. Meaning the types of medicines that would through these a lot quicker if I can, in my liberal 20 20 be prescribed to individuals with hypertension? arts mind, put your definition in two prongs; the 21 21 A. Meaning -- yes, meaning trying to optimize first being the disease, and then the second being 22 22 the best sequence of drugs in treating hypertension. some type of exposure, whether it be treatment or 23 23 You got a universe of drugs. How does one decide something else. 24 2.4 what drug they use first, and if that drug fails and A. I understand. 25 25 the state of high blood pressure persists, what Q. And in this case, the disease or diseases Page 50 Page 52 1 drugs should follow? are hypertension and stroke? 2 2 I did the modeling to follow which drugs A. The disease -- the disease is -- in the 3 3 would be the sequence, the best optimal sequence of two-prong approach, the disease that we want to 4 impact is stroke. Q. And under your definition of epidemiology O. Okav. A. And the exposure is the treatment of which you gave me a second ago, the process by which isolated systolic hypertension. the true nature of the exposed disease is deduced, Q. My science teachers would be so proud of me in this case the exposed disease is hypertension? 9 right now. All right. A. The disease it hypertension, and the 10 10 And so that particular publication, or this exposure now is the treatment. Patients exposed to 11 11 journal under your definition, is under the -- under 12 12 the science of epidemiology? Q. Right. Okay. I'm learning some stuff. I 13 13 A. Yes, sir. appreciate this. 14 14 Q. All right. No. 14 -- and I am going to go A. Well, so am I. 15 15 faster than this, but there are some I'm going to Q. All right. So, now, let's look at entry 16 16 No. 13, which is again another journal with respect spend more time on than others. 17 17 No. 14 is something in particular you were to the hypertension. 18 in charge of. I take it that when I look at your 18 A. Thirteen is clinical epidemiology, if I'm 19 19 publications, if your name is in bold and first, reading right. Thirteen is the SHEP Cooperative 20 20 it's a work that you were primarily responsible for? Research Program? 21 A. Yes, sir. 21 Q. Yes. And if I -- it says "The SHEP 22 Q. Okay. The theory of runs with application 22 Cooperative Research Group rationale and design of a 23 for drought predictions, tell me what that is. 23 randomized clinical trial on prevention of stroke in 24 A. Well, one thing I can tell you, that that 24 isolated systolic hypertension"? 25 is not epidemiology. 25 A. Yes, that's right.

Page 53 Page 55 1 1 Q. Okay. all of these to be epidemiology? 2 2 A. All right. This is based on -- I did -- my A. Yes. I would say, to use our common 3 3 language -- could you scroll back up a little bit? dissertation was on the theory of runs, and runs 4 meaning a sequence of the same kind of event. And Okay. So right there. I would say beginning with 5 5 I've applied that to several different fields. One 15, we have a long run of epidemiology articles. 6 6 Q. Okay. And the disease varying from stroke, of them is hydrology. And so I -- myself, along 7 with Irina, Asha, and Bob, wrote -- wrote an article sex bias, and management of coronary artery disease, 8 8 about the application of my dissertation to cholesterol, disease varying greatly? 9 9 hydrology. But this is -- this is straight math and A. Yes. And --10 10 probability. This is not epidemiology. O. But still epidemiology? 11 Q. Okay. 11 A. Correct. And cardiac arrhythmias. 12 12 A. Same thing with 15, if I get ahead of Q. Repeat that for me and the court reporter. 13 13 A. Sure. I'm sorry. Articles 16 and 17 are mvself. 14 14 Q. No, that's fine. I appreciate it. That is cardiac arrhythmias. 15 math and probabilities in statistics? 15 Q. Okay. 16 16 A. Okay. So cardiac arrhythmias, stroke. A. Well, actually no statistics. I think it's 17 just really probability and math. I have to look at 17 Heart function is measured by left ventricular 18 the paper, but I don't think it's statistics. 18 ejection fraction and cardiac volumes. Sex bias --19 19 Q. Theory of -- let me just go back because I I'm sorry, selection bias, and SHEP and also CARE, 20 20 may have missed it. Theory of runs, what is that? Cholesterol and Recurrent Event trials, begins a 21 21 long run of epidemiology-based work. A. Well, okay. You asked for it. 22 22 O. Well, I want the layperson version of it. Q. Okay. Now, I didn't see it anywhere in 23 23 A. Okay. All right. Okay. We understand -vour -- well, let me -- hold on one second before I 24 24 let's make it very simple. Say we're flipping a make that statement. 25 25 coin, okay, and the coin is fair. We expect that You have never been employed by a Page 54 Page 56 over the next 100 flips, approximately 50 of them government agency in the field of epidemiology, have 2 2 are going to be heads. Fair enough? Fair coin. 3 Q. Yes. 3 A. I have, sir. 4 4 A. Okay. Q. You have? Q. That's the theory of a run? A. I have. 6 6 A. Well, actually no, no. I wish. No, no, Q. Okay. And we're going to get to what I --7 no. The theory of runs talks about -- well, how I see you have community service government on your 8 8 likely is it I'm going to get three runs of six CV. What -- when were you employed by the heads in a row? See the run is a successive -- a 9 government to act as an epidemiologist? 10 10 successive occurrence of the same event. A. If we could -- you have me at a little bit 11 11 O. Understood. of a disadvantage because I can't scroll my CV. But 12 A. Okay. And so my work was on solving that 12 if you go to -- I'm not -- I think it comes after 13 probability. And then -- and then in my CV -- my CV 13 this section. I don't actually remember if it was 14 is punctuated with publications in that area. 14 before or after. If you keep going... 15 Q. Understood. Perhaps we can get more into 15 Q. After this section, you have submitted 16 that after this case is over in a casino, but I 16 editorials, published correspondence, books, book 17 understand. 17 chapters, teaching experience. 18 A. Okay. If they let us in. 18 A. All right. It would be before this. So 19 Q. So 14 and 15 both are just mathematics and 19 let me just tell you. I was a member of a food --20 probabilities? 20 federal food and drug administration advisory --21 A. Yes. 21 advisory council. 22 Q. As I perused through several of these, 22 Q. Okay. 23 17 -- well, not so much 17, but 19 involves stroke, 23 A. And that was from 1996 to 2000 and then 24 20 involves cardiac, 21 involves coronary artery 24 from 2000 to 2002. 25 disease, 24 involving cholesterol. Do you consider 25 Q. Now, where you -- you said you were a

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Page 57

member of an advisory board. Was that a paid position?

A. Yes and no. Yes, I got a check, but no, it was, like, \$150 for three days' work or something like that, so...

### Q. Okay. So tell me what you did for the FDA.

A. Oh, sure. So companies in -- if I'm not getting to the point, just let me know, and I'll shorten it.

Drug companies and the FDA bargain/negotiate/fight over new drug applications should the drug company's new drug application be approved. Most times those decisions are amicable. Occasionally, they are contentious. And when they are contentious, the FDA will take the question to an advisory committee. It's called an advisory committee, not an advisory panel. Let me correct myself. The advisory committee. And the advisory committee reviews the information provided by the FDA and by the drug company and makes determinations.

Now, I don't want to leave a bad taste on this. Occasionally, a drug will be an entirely new class that nobody has any experience with, and the FDA by -- almost by definition will kick that to the Page 59

A. Primarily, yes. I don't remember any kidney drug that we reviewed, so they were primarily, if not exclusively, cardiovascular.

Q. And then the second stint on the committee you said were generics?

A. Yes.

Q. So were there any particular type of diseases that you were focused on from 2000 to 2002?

A. I don't remember.

Q. Okay. But the drugs themselves were generic drugs?

A. Yes, sir.

Q. Going back to -- I think it's page 5 of the CV where we left off.

As I went through the publications, Doctor, most dealt with hypertension, cholesterol, cardiovascular, some smoking, stroke. In all of your publications -- and I'm trying to do this for your own benefit -- are there any publications regarding viruses?

A. There are no publications here focused on viruses. That's correct.

Q. Okay. Have you authored or been a contributing author in any publication that focused on viruses?

Page 58

advisory committee as well.

So it's not always about a fight.

Sometimes it's just very new -- a very new assessment and a very new mechanism of action or very new class of drugs, and that goes to the advisory committee as well.

Q. Okay. And you did that in two separate intervals, in 1996 to 2000 and 2000 to 2002?

A. Yes, sir.

Q. So why do we have -- you said '96 to 2000 and then 2000 to 2002. Is it just '96 to 2002 or --

A. Yes. But I wanted to be clear. It's two different committees. So my first committee was the cardio-renal advisory board, and that was from '96 to 2000. And then 2000, 2002 was -- essentially it's the board that -- the committee that manages generics, generic drugs.

Q. Now, the first one, you said -- it sounded like something to do with the heart, but I'm not sure.

A. I'm sorry. I spoke too fast. Cardio-renal, c-a-r-d-i-o, hyphen, renal.

Q. And would that -- I'm assuming that means that the drugs that you were evaluating were in the area of cardiovascular?

Page 60

A. Well, my publication -- my universe of publications is right here.

Q. Okay.

A. So -- so I'm -- I'm just trying to think about what was the first thing.

No, nothing published. I am working on a book right now that will be published in March 2021 that talks about viruses, but nothing published on my CV, nothing so far.

Q. Now -- so with that I do not have to go through the some 200 plus articles because, in essence, we've talked about some of them.

I understand the scope of what you deem to be the field of epidemiology. And we can agree that none of the articles that you have published to date involve the area of viruses, so we don't need to look at the titles of each one.

A. That is true. The -- the articles are not the universe of my work, but focus only and solely on articles. That is correct.

Q. Okay. No, we'll get to the entire universe, I promise. All right. Let's go to page 20 of 34, please.

Editorials -- the editorial listed here -- what's the difference between an editorial and the

Page 61 Page 63 1 1 journals that we've looked at? statistical analysis. 2 A. The articles -- so far we've looked at 1 I want you to tell me about this 3 through 22 -- are reflective of original research. Experimental Methods in Epidemiology in fall of 4 So one way, shape, or form, original data that 5 5 nobody had seen before had been published along with A. Yes. 6 6 the analysis and the interpretation. Q. Where were you teaching? 7 An editorial is different. An editorial A. University of Texas School of Public 8 8 responds to data that has been published. Commonly 9 these are invited, and sometimes they're Q. Okay. Now, clearly with the statistical 10 10 controversial. But editorials are of a different analysis, the immediate biometry -- I prefer saying 11 11 dope. They are much shorter, and they are biostatistics. 12 12 responding to the data that has appeared in someone A. Okay. 13 13 else's work. Q. You had 100 percent responsibility. So 14 14 Q. Do you recall whether you were invited for tell me the distinction between that and the 15 this particular editorial? 15 25 percent responsibility that you had in 16 16 experimental methods of epidemiology. A. Oh, yes, yes. 17 17 Q. Okay. And you were invited? A. Sure. So this is part of my -- what did we 18 18 say -- a secondary appointment. I was asked to help A. Yes. 19 19 Q. And it involved -- again, in the area of co-teach a class in epidemiology. 20 20 Now, I'm in biostatistics. I as a the heart, cardiac? 21 21 A. Cardiac -- cardiac arrhythmias, yes. biostatistician can't be 100 percent responsible for 22 22 an epidemiology course. The epidemiologists have to O. Okay. And we actually saw earlier a number 23 23 of articles that you published in the same area? take -- have to be responsible for most of the 2.4 24 A. Yes. course, most of the -- most of the administrative 25 25 Q. Okay. Published correspondence, give me burden. So my responsibility was 25 percent. Does Page 62 Page 64 1 the distinction there. that answer your question? 2 2 A. That's just a letter to the journal really. Q. It does. And can you tell from -- and 3 3 That's all that is, is a letter. let's just stick with the fall of 2001. Can you 4 4 tell from -- and I can't, that's why I'm asking --Q. Okay. 5 from your CV, what the underlying disease was? A. One paragraph, two or three paragraphs, 6 6 A. Oh, sure. So here we are talking about the really very short. design of epidemiologic studies, and I believe these Q. Got it. And none of your correspondence were principally clinical trials. dealt with viruses, correct? Q. Okay. A. Could you go to the next page, please. 10 10 A. That -- from an epidemiologic point of Oh, never mind. That -- that's books. So none of 11 11 view. And did that answer your question, or do we my correspondence deals with viruses. 12 12 need to go further? Q. Okay. You have -- and let's go from the 13 13 Q. Well, let me do a follow-up question, and bottom of page 20 to the top of page 21 under 14 14 that may help. "Books." You haven't published any books on 15 15 So you are in collaboration with the viruses, have you? 16 16 A. I've published no textbooks on viruses, no. epidemiologists at University of Texas. They ask 17 17 Q. Go to the next section, which is "Book you to help or to collaborate in the teaching of a 18 18 Chapters." You haven't written any chapters in any class in which you are responsible at 25 percent, 19 19 correct? books on viruses, have you? 20 20 A. Yes, sir. A. No. 21 21 Q. And your contribution to that 25 percent, Q. Okay. Now, the next section is your 22 correct me if I'm wrong, is specifically related to 22 teaching experience, and you have quite an extensive 23 your experience in biostatistics and the management 23 teaching experience dating back to 1987, which I 24 and design of clinical trials? 24 believe is the year before Mr. Davillier graduated 25 A. I think that's fair. 25 high school. Most -- and correct me -- lots of it

Page 65 Page 67 1 O. Okay. In all of these various courses identified as falling under epidemiology, and it's 2 taught beginning in the fall of 2001, they go clear there -- of any of those instances, did the 3 through the fall of 2009, then -- yeah, and then it supervision involve epidemiology with the illness drops down in the fall of 2010 to 15 percent. being a virus? 5 But, nonetheless, is that an accurate A. Oh, boy. I don't know that I can answer 6 description of your role in all of these that for you. I'll look at these. experimental methods in epidemiology where you would O. Well, I'll just tell vou. I went through bring your clinical trial expertise and them. We have lung disease, coronary artery 9 biostatistics expertise to the classroom? disease, maternal weight status, relationship 10 10 A. That is correct. However, it wasn't -- the between prenatal smoking and infant birth. None of 11 11 subject matter was not all cardiology and them that were -- that stood out to me as involving 12 12 hypertension. 13 13 Q. Okay. The subject matter varied? A. Sure. I mean, some of these I don't have 14 14 A. Right. anything. Like, Dillip, one, two -- fourth or fifth 15 Q. Okay. Can you tell from this what the 15 one from the top. 16 subject matters were? 16 Q. Yes. 17 A. Well, from my memory, I could tell you that 17 A. I don't know what Dillip's was in. I don't 18 it was -- there were behavioral trials, trials in 18 know what Wednesday Foster's was in. 19 19 Q. Okay. I'm now looking through your guest cost-effectiveness, vaccine trials that involved 2.0 20 viruses. That's all I remember. lectures. There are several in biostatistics, 21 21 Q. Behavior trials, vaccine trials, several involving clinical trials, several involving 22 22 cost-effectiveness trials? cardiovascular disease and other related cardio-type 23 23 A. Quality of life trials would be another. issues. I did not see a single guest lecture 24 24 Alternative medicine trials would be another. involving viruses. 25 25 Q. And I'm going to try to do a simplified A. I got one for you. Page 66 Page 68 1 version, just for my mind. Q. All right. 2 2 Subject matter notwithstanding, your A. No. 17. 3 3 Q. Seventeen. Let's look at No. 17. "The expertise would be the procedural side of a clinical 4 trouble with" --5 A. As I understand procedural, I would A. Terfenadine. 6 Q. -- "Terfenadine; Seldane, Safeguarding the disagree with you. I would say that my contribution 7 Population, and the Policy Maker's Dilemma,' had to do with the epidemiologic and biostatistical 8 Opportunities in Biostatistics Workshop 1984." component. A. Right. Let me try another way. I'm not in these 10 10 Q. Tell me about that. courses to be a -- to talk about the biostatistic 11 A. Sure. So terfenadine, or the trade name side. I'm there to talk about the epidemiology and 12 12 biostatistics and how they work together. Seldane, is the treatment for rhinitis, and rhinitis 13 13 can appear from viral infections as well as seasonal O. Okav. 14 14 allergies. And the problem with terfenadine was A. And I pointed out earlier that -- I won't 15 15 that it caused a very rare cardiac arrhythmia, rare repeat this, that my definition of a clinical trial 16 16 involving the Bradford Hill causality -but lethal. And there were discussions about -- and 17 17 I lead discussions about what is the best thing to 18 18 do with a drug that is effective for most of the A. Right. Well, the question is, well, how 19 19 do -- how does one do that? I mean, it sounds nice, population but lethal for a few. 20 20 Q. All right. So this drug -- so this affluent, but how does one actually do that? I 21 21 particular guest lecture focused on a drug, that the would talk about that. 22 drug was intended to treat a virus? 22 O. Okav. That's fair. 23 23 A. Yes. I mean --So let's look at student supervisions. Of 24 Q. That's virus being a viral infection? What 24 all of the instances where you served in a 25

supervisory role of students, many of which you have

type of viral infection?

Page 69 Page 71 1 1 A. A mere -- a common cold. So a rhinovirus, A. -- as being about viruses. 2 2 Q. Okay. And so what about basic adenovirus, picornavirus, coronavirus, these kind of 3 biostatistics is -- what is NIH? What organization viruses that cause common cold. Now, it wasn't used 4 4 only for that. It was also used for hayfever. is that? 5 5 A. National Institute of Health. Q. Now, you just used a buzz word, 6 6 coronavirus. Q. Okay. K to R0 meeting in Bethesda? 7 7 A. Yes. A. Oh, I'm sorry about the jargon. K to R01 8 Я is really a breadth of grants that grant -- that Q. And in the same context, the common cold. 9 9 grantees request. A K award is an award for junior 10 10 Q. You're not talking about COVID-19 when investigator. An R01 award used to be for junior; 11 11 you -- just now, when you used the phrase now it's for more senior. 12 12 Q. Both 90 and 91, is it fair to say that the 'coronavirus''? 13 13 primary purpose of those lectures were A. I would say I am not talking about 14 14 biostatistics? SARS-CoV-2. 15 15 A. Oh, sure. I mean, we gave examples using Q. Right. Which causes COVID-19? 16 16 viruses, but yes, they're titled biostatistics. A. Correct. The other coronaviruses that have 17 17 Q. Okay. Certainly, Doctor, I went through been around for many years that cause the common 18 18 all of your community service government and your 19 19 community service nongovernment. And although there Q. But the actual lecture was not about the 20 20 are a lot of biostatistics, there's a lot of virus; it was about the drug? 21 21 cardiovascular work, lung, blood institute. I did A. Well, the actual lecture was about viruses 22 22 not see anything that was specifically related to and what they do, as well as about the arrythmia. 23 23 I mean. I have to lecture about the diseases the 24 2.4 A. Can we scroll through those, please? I drug treats and also the side effects of the 25 want to make sure we don't have any trojan horses disease. So that was an important component of the Page 70 Page 72 1 1 lecture. And then the question is, what advice do here. 2 2 you give the community about the use of the drug? Q. Sure. 3 3 Q. Okay. You can see how someone with a A. Let's see. Well, the sickle cell 4 program -- let's look at 7. Is that right, 7? Let layperson mind like me, when I read the -- this 5 me scroll down to -- where am I -- 21 and 22 and 24, title, it wasn't obvious to me that this involved 6 6 and if we go below that, 30, and below, 35, okay, treatment of the common cold. 7 36, 41, 51, 61. So are there any other guest lectures that 8 8 Q. Let's stop at 61 for a second. you would like to point out to me that involve 9 A. All right. viruses? 10 10 A. Can we move past -- I think we've done 1 Q. Scroll back to the first page, page 27 of 11 11 34. So all of the ones that you've identified thus through 25. 12 12 Q. All right. Let's go to the next page, 26 far deal with the -- sickle cell? 13 13 A. Correct, yes. through 58. 14 14 A. So let's see. Yeah, I think some of these Q. All right. 15 15 do, but we don't talk about a particular virus. A. Right. 16 16 For example, Sample Size Guidelines Invited Q. And why don't you tell me your reasoning as 17 17 Lectures, No. 28. I think I gave examples involving to why you identified these as involving viruses. 18 18 A. Sure. These all are part of a program to vaccine trials. Forty-one, I gave examples of 19 19 vaccine trials. But most of these are not about test the effect of a therapy in sickle cell disease 20 20 viruses. in children. And one of the measures that we had to 21 21 Q. Okay. Let's go to the next page, 59 keep focused on were infections, including viral 22 22 infections. It wasn't exclusively viral infections. through 97. 23 23 A. Okay. I think 90 and 91, we talked about These poor kids get pneumonias as well, bacterial 24 24 viruses, but those are the only ones -pneumonias, which are very bad. 25 25 But viral infections. And so my role was Q. (Indiscernible.)

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Page 73

to review the data, including the data on viruses, to determine whether the study should be allowed -- allowed to proceed or not.

# Q. Okay. So let's just take them one by one. The -- this entire program was designed to help sickle cell patients with sickle cell anemia?

A. Correct. Right, the hypothesis -- the clinical hypothesis was that there was an alternative therapy to the standard therapy for these children with sickle cell disease. There is now a therapy for their sickle cell disease --

O. Okav.

A. -- that would help them. That was the clinical hypothesis.

## Q. And your role was -- first of all, this was community service, right? You didn't do this for pay?

A. I actually don't remember if it was pay. It wasn't very much, but I think it was for nothing. It might have been \$150 a year, but...

Q. Okay. No. 7, National Heart, Lung, and Blood Institute SCOR Site Visit. So what exactly did you do in May of 1987 that warrants putting this on your CV?

A. Oh, we reviewed the applications of -- yes,

Page 75

### Q. In 1987, what epidemiology background did you have?

A. Oh, in '87, again, I was -- I've been a physician, and now I've been a physician for almost ten years. So as a physician, I bring epidemiology background because I have training in epidemiology and also at -- in medical school. I also have clinical acumen, and I also had training -- excuse me, training in my Ph.D. program in epidemiology. And that was all behind me as of May 1987.

## Q. Got it. And so the primary purpose of this particular organization was to address alternative therapies for sickle cell, right?

A. Well, I put it this way. The purpose of this group -- the specific purpose of this meeting was to identify the candidate universities who had sufficient research experience and expertise that they could design a good program.

# Q. Okay. And your testimony to me today is, within that, those applicants had to identify what they had done or their research with respect to viruses?

A. Well, actually no, sir. It is that these were-- these institutes had to show their ability to manage and treat infections in sickle cell children

Page 74

the applications of a host of different national -- a host of university research institutes that wanted to conduct this program. And in that review they had to provide data showing their research prowess in general and their ability to successfully treat children. And in that data included data about bacterial and viral infections.

### Q. Okay. Were you on the board of the National Heart, Lung, and Blood Institute?

A. I don't know what a board is for the National Heart, Lung, and Blood Institute. Did I say that somewhere, board?

# Q. No, no, no. I'm -- so how did you come to be one of, I assume, a number of physicians that would review these applications?

A. Oh, that's a fair question, especially since I wouldn't get my Ph.D. for another month. I didn't graduate until June of 1987. But they were interested in people who had -- who were physicians and had an epidemiology background, who could provide some insight into whether these candidates -- which of these candidate sites was worthy for funding.

### Q. Okay. But you didn't --

A. So I was invited to be part of it.

Page 76

with diligence. So it wasn't their research prowess with viruses. It was their clinical prowess.

# Q. Okay. Got it. So is it fair to say in each of these that we have identified thus far involving sickle cell, your role was the same?

A. Yes.

# Q. Okay. Other than your work with this group involving sickle cell, are there any other governmental community service roles that you played that would have involved viruses?

A. I don't know. How many of these do we have?

### Q. They stop at 112. I'm now on 61.

A. Okay. I'll try to be quick. Let me look at these real quickly. Okay. We can scroll to the next page. 102, but that's the same genre we've been talking about. I think No. 112, the Innovation in Regenerative Medicine Symposium.

### Q. Okay. What makes you think that?

A. Well, that was a collection of lectures that were given by experts in the field of regenerative medicine, and some of them discussed the role of virology in regenerative medicine.

Q. Would you have given a lecture on the role of virology?

Page 77 Page 79 1 1 A. No, I would not have. all done within the -- the sphere of drugs. 2 Q. Okay. So in December of 2017, you didn't Q. Okay. So, now, I'm getting some clarity. 3 lecture anyone on the role of virology, did you? Berlex is a pharmaceutical company? A. I did not. A. Yes, sir. 5 Q. What about your community service in Q. And they are conducting clinical trials 6 6 nongovernment? Do you have any recollection of with respect to one or more of the drugs that 7 lecturing or participating in any type of seminars they're creating? 8 or meetings involving viruses? A. Yes, sir. 9 A. I'm up to 26, and so far the answer is no. Q. And they ask you to be a part of the data 10 10 Oh, 52, so far the answer is no. Oh, goodness. safety and monitoring board for them? 11 11 Seventy-four and 75 definitely. All of the Berlex A. Yes, sir. 12 12 Pharmaceutical meetings were virology. Also, the Q. Okay. In 2003, do you recall the drugs 13 13 sickle cell disease DSMB meetings were virology, that you were asked to evaluate? 14 14 A. I would say in general, yes. I don't discussed viruses. 15 Q. Let's talk about 74. 15 recall the specific species, but in general, yes. 16 16 Q. Okay. But in essence -- and is it fair to A. Sure. 17 O. What is DSMB? 17 say that because of your experience in handling 18 A. DSMB is a data safety and monitoring board. 18 clinical trials or being involved in clinical 19 19 It -- I'll stop there and then... trials, you received the invitation? 20 20 Q. Now, these are non -- you described them as A. I don't think so. I think it's the fact 21 21 nongovernmental community service. So did you serve that I was a physician as well and that I had 22 22 on the data safety and monitoring board? epidemiology skills. 23 23 Q. Epidemiology, meaning the process by which A. Yes. 24 24 Q. All right. What role? the true nature of the exposed disease is deducted? 25 A. I was -- I was an epidemiology and A. Yes, sir. Page 78 Page 80 1 1 biostatistics member of the board. Q. Okay. And do you have any recollection, in 2 2 either 2003 or 2004, what the underlying disease was Q. So you were just a member of the board? 3 A. Well, everybody was just a member of the 3 that Berlex was attempting to find a cure for, for 4 4 board. Yes, right. lack of a better phrase? 5 Q. I didn't mean that disparagingly. You were A. I would only say it's a cardiovascular 6 6 a member of the board. You brought whatever skill disease. I don't remember the details of it. 7 set you had, but you were a member of the board? Q. Okay. Was that a paid position on that 8 A. Right. The clinical epi and biostat skill board? 9 set, I brought to the board. A. I think it was. 10 10 Q. Okay. And you were on that board for how Q. So the viral or virology that you just 11 11 long? described would have been in the context of 12 A. I guess, you know, this list will tell us, 12 cardiology? 13 but I think a couple of years at least. 13 A. Yes, sir. 14 Q. Okay. And what is -- what was the primary 14 Q. Okay. All right. We stopped at 74, 75. 15 mission of that board? 15 Why don't you take a look --16 A. The board was to oversee the conduct of a 16 A. Well, I'm sorry. I do have to clarify one 17 clinical trial. We would look at all of the data 17 18 and determine whether there was sufficient reason to 18 Q. Sure. 19 stop the study because of an early benefit or to 19 A. The -- remember in the exposure/disease stop the study due to harm or to continue the study. 20 relationship, the disease was a cardiovascular 21 Q. Now, this says nongovernmental. Is this 21 disease. Here the exposure was the virus. The 22 board, data safety monitoring board, a private 22 exposure is the virus. 23 institution? 23 O. What virus? 24 A. Berlex Pharmaceuticals funded the study, 24 A. I would have to look it up. It was never 25 and they asked us to oversee the trials. So it was 25 approved. I -- for all I know, it's still

proprietary. I don't remember. But I can tell you that the -- the treatment was to deliver about a trillion viruses to people to see if it had a beneficial cardiovascular effect. Q. The treatment was to deliver viruses to see if it had a cardiovascular effect? A. Right. A beneficial cardiovascular effect. Q. So what role in that analysis did the drug play that you were evaluating? A. The drug was the virus. 11 Q. Okay. So the drug, meaning the 12 pharmaceutical that Berlex was attempting to create, 13 was the virus? 14 A. Yes, sir. That -- the exposure was the 15 virus. 16 Q. Okay. So you defined -- you can then 17 define virus to include a drug that would actually 18 have a beneficial effect on a patient? 19 A. Well, it -- the drug itself was the virus. 20 It was nothing else but the virus. So these 21 patients were deliberately infected with the virus. 22 23

Q. Which was a pharmaceutical drug that Berlex

A. Right. If we define drug as something that they created, yes, that's right. It was the drug

Page 81 Page 83 1

A. Well, it's not that I changed the definition of a virus. I think what we've done is expand the definition of a drug to include virus.

### Q. Got it. Okay.

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A. But, you know, it -- it was a very novel program, and I will tell you, quite honestly, it was very scary because now in most times in medicine, you want to avoid exposure to viruses. We don't think of exposure to viruses as being a good thing. Maybe a hundred years from now, we'll know better, but we don't think of it as a good thing.

Here, patients are exposed not to a handful of viruses, but to a trillion viruses. And it's our responsibility to see if these patients are being managed well and that they are doing well and to stop this if in fact it looked like they were not doing well. So this was a very, very sensitive research effort.

### Q. (Indiscernible.)

A. Otherwise, we wouldn't know much about it. Yes?

### Q. How many members would be on this data safety monitoring board?

A. It depends. I would say, in general, between three and seven.

Page 82

that they created.

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### Q. And why then do you call the drug a virus?

A. Because the drug was the virus. You know, we think of drugs as molecules, statins for example, ACE inhibitors, aspirin. Well, there was no such molecule in this drug. It was only viral particles.

### Q. Okay. You said -- what drug was it?

A. I don't -- for me to answer that question, I would have to have remembered the exact genus and species of virus they delivered, and I don't remember that.

Q. Okay. So the data safety monitoring board hired by Berlex to monitor their application of a particular drug that they created to patients that had cardiovascular disease?

A. I would say that we were to monitor the effect of the virus they injected on its role in producing or preventing cardiovascular disease.

Q. Right. And the virus being what they hoped would be something that would help cardiovascular disease?

A. Yes, sir.

Q. Okay. I wonder how many other doctors define virus in the manner in which you just defined it, but it's very interesting to me.

Q. Okay. And of the three to seven individuals, what disciplines do you recall -- let's say just in 2003. When you served on the monitoring board for Berlex Pharmaceuticals in 2003, how many

Page 84

A. I don't remember, but I would say approximately five or six. I just don't remember.

individuals were on that board?

### O. Okav. And of the disciplines of the five and six, what backgrounds would the individuals

A. Well, if the end point is cardiology, then you'd have to -- you need a cardiologist. You would also need to have a specialist in the exposure. So in this case, there would be the physicians or the -- or the scientists who governed the exposure, who determined what right -- what the best dose was, what the virus should be.

So what do we have? We have a cardiologist so far; the scientist who developed the exposure -that's two -- an epi biostat person. That's me. That's three. You could have the chair of the steering committee. And I don't want to make it -get too complicated here.

### Q. Yeah.

A. But clinical -- clinical centers that

Page 85 Page 87 1 1 conduct the research, they have their own group. picornaviruses. That group has a chair called the chair of the Q. Other than respiratory tract viral 3 3 steering committee. They can be on this -- on the infection, what about type of viral --A. Oh, I would include early flu as well. It DSMB as well, so that's four, I think. And then 5 5 there would be anesthetists. was September, so it wasn't really flu season, but 6 6 Q. And in 2003, when Berlex Pharmacy was early flu. intentionally injecting viruses into individuals, O. Anything else? 8 8 how many individuals participated in that trial? A. In terms of what we -- what patients came 9 9 A. I don't remember. It certainly -- I would to the facility with, I would say no. However, they 10 10 say this, it certainly was not thousands. It was did develop viral diseases while they were there. 11 11 smaller than that, but I just don't remember. Q. And the work that you're describing would 12 12 Q. Would this have been a trial that would have been work -- volunteer work that you did as a 13 13 have been FDA approved? medical doctor to assist individuals that had 14 14 illnesses when they evacuated to Houston? A. Absolutely. 15 Q. Okay. Where could I find literature on a 15 A. Yes. But it turns out it was, 16 trial like this? 16 unfortunately, a little bit more than that. 17 17 Q. Sure. Trust me --A. I don't know how to answer that. 18 18 A. It was a --Q. Okay. 19 19 A. I mean, but -- but I don't mean to be Q. Go ahead. 20 20 facetious. I would think something like therapeutic A. I -- I simply mean to say that we had a new 21 21 epidemic that developed while we were there, that I 22 22 was involved in the identification of and treatment O. When you served, you weren't serving in the 23 23 capacity as virologist, were you? 24 2.4 A. I was not; just as a physician, Q. Why don't you describe that epidemic to me. 25 25 epidemiologist, and biostatistician. I'm expected A. Sure. It was a rotavirus epidemic that led Page 86 Page 88 1 to know virology, but I wasn't serving as a formal to --2 2 Q. A what virus? virologist. 3 Q. Other than 74 and 75 --A. R-o -- r-o-t-o, rotavirus, or r-o-t-a 4 A. It would be everything. I'm sorry to cut maybe, rotavirus, epidemic in children. 5 you off. Everything that had Berlex Pharmaceutical Q. All right. Tell me about it. 6 at this point, would be the same thing. So 76, same A. Sure. We noticed that the number of children with new and persistent diarrhea were -thing. was increasing. And we had to put our heads O. Okav. 9 together to help identify the cause until we got A. Of course 86. 10 10 viral types back. O. 86? 11 11 Now, this was really clinical epidemiology A. Yeah. Can we scroll down, please? Oh, 12 12 okay. Near the end here. thinking because we didn't have sophisticated 13 13 technology to help us in the first few days, and we Q. So tell me how your work during 14 14 decided that this was a new -- new for the Katrina Hurricane Katrina had all -- had anything to do with 15 15 evacuees, diarrheal viral infection, and began to viruses? 16 16 isolate these children. And at that point, we got A. Of course. So as a treating physician 17 17 tires back which suggested that it was rotavirus. there, many patients had viral diseases with 18 18 Q. You got what back? complications. They had viral diseases plus rampant 19 19 hypertension or viral diseases and the -- in the A. I'm sorry. We got viral -- viral chemical 20 20 results back by -- viral chemical essays back that presence of poorly treated diabetes or viral 21 21 told us it was rotavirus. diseases in cancer. 22 22 Q. How many individuals does it take to be Q. Give me an example of some of the viral 23 23 infected before something is determined to be an diseases you're describing. 24 24 epidemic? A. Sure. Upper respiratory tract viral 25 25 A. That's a good question. It really depends infections caused by rhinoviruses, adenoviruses,

Page 89 Page 91 1 1 on the infection. It depends on the -- the -physician -- the only physician healthcare team were 2 2 what's causing the disease. volunteers, at least in the early phase. 3 3 For example, I'll give you tuberculosis. Q. Was the group organized with a particular 4 4 It takes more than one or two if these people are name or structure? 5 5 spaced far apart, for example. The bacteria -- the A. No. 6 6 Q. Okay. tuberculosis bacteria spreads relatively slowly unless people are really close together. MR. MILLER: Why don't we -- we've 8 8 Q. How many people -- oh, I'm sorry. Go been going for a minute. Why don't we take 9 9 another five-minute break. ahead. 10 10 A. On the other hand, to go with the other THE WITNESS: Fine with me. 11 extreme, pneumonic plague spreads like wild fire 11 MR. ALVENDIA: Yeah, it's fine. 12 12 through a population, where it's plague -- plague (Recess taken.) 13 13 Yersinia pestis, plaque bacteria that is spread not MR. MILLER: All right, Doctor. 14 14 from broken abscesses but through exhaled droplets. Back on the record. Thank you. 15 That spreads very rapidly. 15 BY MR. MILLER: 16 Q. How many people does it take to be infected 16 Q. What -- who engaged you in this matter? 17 for -- with a rotavirus for it to be determined to 17 A. Dave Matthews out of Houston. 18 18 Q. Dave Matthews? be an epidemic? 19 19 A. Okay. So I -- so I'm a physician so I'm A. Yes. 20 20 Q. Who is Dave Matthews? going to have to make a physician's distinction 21 21 here. It doesn't take much to get infected at all. A. Dave Matthews is an attorney in Houston. 22 22 Infected simply means that -- in this case the virus O. And who does Mr. Matthews work with? 23 23 is on or in the other person. That's infection. A. I couldn't tell you. If you know, maybe --24 2.4 maybe you can ask me a different way. I can be more Illness is the impact of the infection on the 25 25 individual. helpful. Page 90 Page 92 Q. Which one -- which one triggers an Q. Is he with a law firm that you -- do you 2 2 epidemic, illness or infection? know the law firm that he's with? I'm sorry. 3 3 A. Actually both do. A. Oh, I -- I -- I think it's just Dave 4 4 Matthews LLC, but he does work with other lawyers Q. All right. So --5 A. Virus leads to illness, leads to spreading whose names I don't recall right now. I'll probably 6 6 an epidemic. get in trouble for that. 7 Q. How many individuals does it take to have Q. No, that's okay. And what did Mr. Matthews 8 infection and illness in a rotavirus for it to be engage you to do? determined to be an epidemic? A. Well, he put together a phone call with 10 10 Mr. Houghtaling. I think -- I think Rico was on A. I would -- I would have to look and see. 11 11 the -- yeah, it was a Zoom call. I think Rico was I can't remember -- don't remember that. 12 12 on the phone. There might have been one or two O. Okav. Who determined that the work that 13 13 you were doing in Houston with children, during other people on the phone. And --14 Hurricane Katrina, that contracted rotavirus that 14 MR. ALVENDIA: Allen, just --15 15 Hold on a second, Doc. became infected and ill, was an epidemic? 16 16 A. It was our decision as physicians and Just to clarify, this is not an 17 17 objection. Dave Matthews introduced the epidemiologists. 18 Q. Who's "our"? You and who else? 18 doctor to John Houghtaling and our team. 19 19 A. Oh, and the other physicians I was working We obviously have retained him, have paid 20 20 with. him, and so forth. 21 21 MR. MILLER: All right. Q. Was there an organized group of physicians, 22 22 or was this just volunteer work? Go ahead, Doctor. You can finish. 23 23 MR. ALVENDIA: That's the A. Well, it was all volunteer work. 24 24 Q. Okay. But what -clarification. 25 25 THE WITNESS: Thank you for that. You A. We were the only healthcare -- only

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	Page 93		Page 95
1	said that more eloquently than I did.	1	A. Actually, no. I was just asked some
2	A. We had the first Zoom call where we talked	2	questions about my opinion based on my background
3	about the impact of coronavirus on businesses,	3	and expertise about what would happen with viruses
4	particularly restaurants and particularly the Oceana	4	in restaurants in general and specifically would
5	restaurant, and just what does a virus do in a	5	there would there not be physical damage.
6	restaurant.	6	Q. So you were asked specifically would there
7	BY MR. MILLER:	7	or would there not be physical damage?
8	Q. Now, prior to that call, had you worked	8	A. From the virus, from the SARS-CoV-2 virus,
9	with Mr. Matthews before?	9	
10	A. Yeah. Dave and I have worked in the past,	10	yes. Q. And what was your response?
11		11	•
12	a long time ago.	12	A. My response was there would be damage.
13	Q. What type of work did you-all do together?	13	Q. And at the time you had this initial call
14	A. It was a Big Pharma litigation.	14	in 2020, and I think you just testified that you had
15	Q. Do are you working with Mr. Matthews on	15	never done any studies with respect to SARS-CoV-2 in
16	any other coronavirus cases?	16	restaurants, how did you come to the conclusion that
	A. No.		there would be damage?
17 18	Q. Okay. And so what you said you had a	17	A. Because I understand viruses. I understand
	Zoom call and you talked about the impact of	18	physics. I understand chemistry.
19	coronaviruses on restaurants?	19	Q. Did someone give you a definition of what
20	A. Yes.	20	damage was?
21	Q. Okay. Had you done any types of studies	21	A. No.
22	with respect to coronavirus on restaurants prior to	22	Q. You used a general your general
23	the Zoom call with Mr. Matthews and Mr. Houghtaling?	23	understanding of what the word "damage" means?
24	A. No, I had not.	24	A. Yes, sir.
25	Q. Okay. When was that call?	25	Q. Okay. And what is that?
	Page 94		Page 96
1	A. Oh, I would say well, this is the 10th	1	A. Physical transformation that leads to loss
2	of November. Maybe three week three and a half	2	of use.
3	weeks ago.	3	Q. And where did you get that definition from?
4	Q. So your first call with Mr. Houghtaling and	4	A. I don't know. I think it's just my
5	Mr. Matthews was in October?	5	experience with the use of the English language.
6	A. Yes, sir.	6	I didn't look it up anywhere. It's just my
7	Q. Had you spoken to I'm going to do it in	7	impression of what damage is and does.
8	pieces. Had you spoken with Mr. Matthews about the	8	Q. All right. And so I just want to make sure
9	effect of coronaviruses on restaurants prior to	9	I'm right. The first call that you had regarding
10	October of 2020?	10	being involved in this lawsuit for Oceana Grill was
11	A. I had not.	11	about three weeks ago?
12	Q. Had you in your call of October 2020	12	A. Between three and four weeks ago, yes.
13	where it was both Mr. Matthews and Mr. Houghtaling,	13	Q. Okay. So that would have been I'll
14	was that the first time you had spoken with	14	give let's say four weeks. That would have been
15	Mr. Houghtaling?	15	October 10th?
16	A. Yes, sir, it was.	16	A. Fair enough.
17	Q. Was anyone else on that October 2020 call?	17	Q. Somewhere between October 10th and
18	A. Rico was on the call.	18	October 17th was the very first call with
19	Q. That's right. You did say him.	19	Mr. Matthews, Mr. Houghtaling, Mr. Alvendia, and
20	A. And there might have been one or two	20	maybe a couple other individuals on the phone?
21	others, but I do not remember their names.	21	A. Yes, sir.
22	Q. Okay. And what was were you given any	22	Q. All right. After they asked you the
23	instructions during that call as to what your role	23	questions about whether you believed SARS-COVID
24	would be with respect to this litigation involving	24	SARS-2, COVID-19 would result in damage, and you
25	Oceana?	25	said yes, did they then give you any other
	Occalia:		said yes, did dien dien give you any other

Page 97 Page 99 1 1 instructions? vour work consist of? 2 A. Not yet, no. A. Well, the work consisted of identifying the 3 Q. All right. When were you given additional information that I thought would be -- that was a -my scientific foundation for my argument. instructions? 5 5 A. I think after I told them why. Let me be Q. Let me make sure. The call that happened 6 6 clear. After I told them the basis of my opinion on -- somewhere between October 10th and and we talked about that for a few minutes, they October 17th, you gave the lawyers in this case a 8 8 asked me would I be willing to be involved in this conclusion about physical damage, correct? 9 9 case as an expert. Let me back up for a second. A. I gave the lawyers in this case my opinion 10 10 They then gave me some background on the about the --11 11 case about the -- about the restaurant, about the MR. ALVENDIA: Wait, wait, Doctor. 12 12 insurance companies, the argument the insurance Objection to form. 13 13 Go ahead and you can finish your companies were making, and then they asked me would 14 I be willing to be an expert, serve as an expert in 14 answer. 15 the case. 15 THE WITNESS: Okay. 16 Q. And then what? 16 A. I gave the lawyers in this case my opinion 17 17 about what CoV -- SARS-CoV-2 -- essentially what A. Then I said yes. 18 18 human viruses or viruses that infect humans in Q. All right. Then what? 19 19 A. Then they told me that I had to have a general do. Yes, I did do that. 20 20 report, and I said, well, that's fine. And so I BY MR. MILLER: 21 21 began to work on a report. Q. Okay. And when you say "viruses that 22 22 O. Okay. Have you ever been inside of Oceana infect humans," you don't just mean -- so let's 23 23 affectionally call SARS-CoV-2 and then the ultimate restaurant? 2.4 2.4 A. I have not, sir. sickness -- is it fair scientifically, for the 25 25 Q. So somewhere around October 10th, purposes of this argument, to call it all COVID-19? Page 98 Page 100 1 October 17th -- was that one call where you A. Actually, it's not fair. 2 2 ultimately gave them your opinions, gave them your Q. All right. Well, then --3 rationale, discussed the contents of the lawsuit and 3 A. I would be -- I would be willing to accept 4 the insurance company's position, asked to be an if you said CoV-2 as the virus -- COVID-19. I think 5 expert, agreed to be an expert, started working on a that's shorthand. If it's helpful to you, it's 6 report? Was that -- that sequence of events was all unambiguous to me. one phone call? Q. So CoV-2 as SARS, can I say that -- or no. 8 A. Well, I didn't work on the report on the So SARS and CoV-2? 9 phone call. But up to that point, absolutely. A. SARS -- we can call the virus either SARS 10 Q. Fair enough. Okay. 10 or CoV-2, and the infection is COVID-19. 11 A. My recollection of the sequence. 11 O. All right. Let's call it CoV-2, because 12 Q. And then you hung up, and then you started 12 then I think that would be more familiar to the 13 working on the report, presumably, at some point? 13 trier of fact that's going to ultimately see all of 14 A. I ate my Wheaties first, and then I started 14 this. So we're going to say the virus CoV-2. 15 15 A. Yes. 16 Q. Okay. All right. And at no time before 16 Q. Okay. So what you gave to the lawyers on 17 you started working on the report did you go inside 17 the -- somewhere between October 10th and 18 of Oceana Grill? 18 October 17th was your conclusion regarding what you 19 A. No, sir. 19 believe a virus similar to SARS or similar to CoV-2 20 Q. And in fact as we sit here today, you still 20 would do to property? 21 have never gone inside of Oceana Grill? 21 A. Yes, sir. 22 A. That is correct. 22 Q. Okay. That it would cause a physical 23 Q. All right. What did your work -- to begin 23 transformation that would lead to loss of use and 24 to draft your report, mind you, after you had 24 also damage? 25 already made a conclusion about damage, what did 25 A. Well, damage was the physical

Page 101

transformation and loss of use, yes.

- Q. Okay. Now, you said viruses in general separately, a second ago. So there are other viruses that would also fall within the definition of a transformation, physical loss of use, which is ultimately damage?
  - A. Yes, sir.
  - Q. Would the flu be one of those?
  - A. Yes, sir.
- Q. Okay. What other -- would the common cold be one of those?
  - A. It depends on the virus.
- Q. But we can just use the flu. The flu would cause a physical transformation, loss of use, and then therefore damage to property?
  - A. Yes, sir.
- Q. Okay. So once you hang up the phone, you've now had the opportunity to discuss the chemical, physical, and epidemiological effect of a virus on property with these lawyers, though then -- and they asked you then to write a report?
- A. I think we have to clarify a couple of nings.
- Q. Okay. That -- that's fair. I want to be --

the body, understand something about the surfaces it would impact, and then write my conclusion.

Page 103

- Q. Okay.
- A. Now, I will say this, I had to -- I'm sorry. Let me stop for a second.
  - Q. Go ahead, please.
- A. I had to learn about things. I had to learn what the mayor and the governor said in March. I had to learn that. I had to learn where Oceana was in New Orleans. I had to come to a conclusion about the -- about the density of people with the virus within a certain distance of the restaurant. I had to do those things as well.
- Q. Why -- if we're going to go through all of these, why was it important to learn what the mayor and the governor said?
- A. Because the governor and the mayor's statement led to the decision at Oceana as to what level of customer support they should sustain, number one; and number two, the -- now, this is the epidemiologic part. This is not the physics and chemistry; it's the epidemiologic part. The governor and mayor gave their assessment based on epidemiology as to what was going on in New Orleans and what the best public safety measures should be

Page 102

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- A. Epidemiology -- I'm sorry. Go ahead.
   Q. No, I said I want to be accurate, so please.
- A. Epidemiology has a role in this case, in my understanding. Epidemiology is not, I say again, not part of the determination of physical damage.
  - Q. Okay.
- A. That is the realm of physics and chemistry, not the realm epidemiology. So our conversation about physical damage had nothing to do with epidemiology. I want to be very clear about that.
  - O. Physics and chemistry only?
  - A. Physics and chemistry, yes, sir.
  - Q. Okay. So you begin to draft your report?
- A. Begin to do the research, getting the literature I know that's there that supports my opinion, yes.
  - Q. Tell me specifically what you did.
- A. Sure. Well, it was an opportunity to review physics, review chemistry, then learn about what happened at Oceana, I mean, how -- I dare to say the characteristics of Oceana -- how big it is, how many people ingress and egress, then examine the studies that talk about the -- let's call it the durability of the virus, how long it exists outside

Page 104

undertaken to protect people.

### Q. Why is that helpful to your report?

A. Well, because I have to understand, in fact, whether it is likely or not that people with -- people infected with, what did we say, CoV-2 -- people infected with SARS-CoV-2 would be in the vicinity of the restaurant.

If they're not in the vicinity -- if the probability they're -- if the probability they're in the vicinity of the restaurant is zero, then I say the probability is also zero, or close to zero, that you'll get any kind of spread within the restaurant, and therefore there won't be any damage. There has to be virus to cause damage. So I have to assure myself that, in fact, the virus is there.

Q. So if -- in a hypothetical similar to the one you just gave, if the mayor or the governor's orders included a complete shutdown and none of the individuals that went into the Oceana restaurant had the virus, the virus would not be present?

A. Well, I would say I would agree with you; however, how one determines somebody walks into a restaurant doesn't have a virus is an important question for that hypothetical.

But accepting your hypothetical, if I can't

Page 105 Page 107 1 demonstrate that there's virus, then there's no Q. You have never -- you have not seen a study 2 point in invoking laws of physics and chemistry to or a test done by some other third party that would 3 understand what happens with viral interaction. confirm to you that CoV-2 is actually physically 4 Q. And I would get to this later on. You present in the restaurant? 5 know, generally, I would ask an expert like A. I have not seen such an activity. 6 yourself, tell me what assumptions you made to come Q. Okay. to your conclusion. MR. ALVENDIA: I'll object. So I'm 8 And so on October 10th through 17th, when not continuing to interrupt you here -- I 9 you had this phone call, and the scenario was was going to object a second ago. 10 10 presented to you that COVID-19 is present in Oceana The legal standard here is more likely 11 restaurant -- I'm sorry, I take that back -- CoV-2 11 than not. The doctor has issued in his 12 12 was present in Oceana's restaurant, what's the report that the virus is present more 13 13 effect? Your conclusion, that there would be a likely than not. I just want to make sure 14 transformation of property resulting in loss of use 14 that we're following those terms here; 15 and therefore damage, there's an assumption there 15 we're not talking in absolutes. 16 that the virus is actually in the restaurant. 16 MR. MILLER: I hear you. I think 17 MR. ALVENDIA: Objection to form. 17 that's your position, and I accept that. 18 You can answer the question. 18 We can argue the legal standard to the 19 19 judge. But yes, I understand that that's A. Correct, yes. 20 20 BY MR. MILLER: what his report says, and I know that's 21 21 Q. The entirety of your analysis, which is a your position. 22 22 chemistry/physics analysis, assumes that there is MR. ALVENDIA: Okay. 23 23 CoV-2 inside of the actual facility at 739 Conti MR. MILLER: All right. 24 24 MR. ALVENDIA: Because you're using Street? 25 25 words like "assumptions," "assume," and so A. I would only disagree with that in saying Page 106 Page 108 1 that my report is more than chemistry and physics. forth. I just want to make sure you're not 2 2 But the chemistry and physics component is mischaracterizing his report, which 3 3 predicated on the assumption that the virus is clearly -- I'm looking at the conclusions 4 right now -- all say "more likely than Q. And the -- not to argue. I accept that not." I just want to make sure we stick to 6 6 your report has more than chemistry and physics. that. BY MR. MILLER: The portion of your report that is necessary for Q. Well, I'll use Mr. Avendia's scenario for there to -- well, one of the major portions of your 9 report that is necessary for there to be you, Doctor, just so we can have some clarity. 10 10 transformation of property, loss of use, therefore In your report when you say "more likely 11 11 damage, is the chemistry, right, the chemistry and than not," that assumes that CoV-2 is present in the 12 12 physics? I mean, the transformation of the property facility, correct? 13 13 is the chemistry and physics? A. Well, one of my -- actually, no. 14 14 MR. ALVENDIA: Wait, wait. Hold on. A. Correct. 15 15 Let me make an objection to form. You're Q. Okay. And if you do not have CoV-2 in the 16 16 using the word "assumes" interchangeably facility, you don't have the physics, correct? 17 17 A. Correct. There's nothing -- right, there's with his conclusion of more likely than 18 18 not, which is based on much more than no agent of transformation. 19 19 Q. And you have never been in 739 Conti Street assumption. It's based ---20 20 in New Orleans -- correct -- Oceana Grill's MR. MILLER: We'll clarify, but go 21 21 restaurant? ahead. 22 22 MR. ALVENDIA: Hold on. It's based on A. I have not. 23 23 Q. You have never done an actual physical test statistics, science, he just said, his 24 24 of the surfaces in Oceana's restaurant? medical experience and so forth. So 25 25 A. That is correct. just -- I'm going to object that you're

Page 109 Page 111 1 1 BY MR. MILLER: mischaracterizing his conclusion. It's not 2 just an assumption. Q. Let's assume the restaurant is locked. 3 But, Doctor, if you understand the No one, not the owners, no person comes into the 4 question, please answer it. restaurant. Is it fair to say that CoV-2 would not 5 5 A. My -- the conclusions in my report are all be present? 6 6 based on "more likely than not," so I don't assume A. If the scenario is -- this is outside the in my report. I don't take for granted, in my bounds of my report, a hypothetical. 8 report, that there's COVID-19 in the vicinity of the O. Yes. 9 9 restaurant. I actually think, go through a thought A. If the assumption is that there is 10 10 process, that leads me to the conclusion that there absolutely no organism that comes near or in this 11 11 is. It's more likely than not it's there. That's restaurant with CoV-2 -- no humans, no bats, nothing 12 12 what I do. And the same -- and the same -- I follow that has CoV-2 -- if CoV-2 is not in the restaurant, 13 13 then you can have no damage from CoV-2 because the that cascade right on through. 14 BY MR. MILLER: 14 agent isn't there. But again, it's outside the 15 Q. All right. And we were talking in 15 bounds of my report. 16 hypotheticals. So I accept that your report walks 16 Q. And I'm going to get to your report in just 17 through a list of scenarios that lead you to "more 17 a second. We're going to talk about well within the 18 likely than not." I don't argue about that at all. 18 bounds of your report, I promise you. 19 19 And I understand that your conclusions are "more Similarly, when you wrote your report, you 20 20 likely than not." and no one else had done any testing of the surfaces 21 21 inside of Oceana Grill, correct? What I'm asking of you, similar to the 22 22 scenarios that you have walked through, that -- if A. Well, I know I didn't do any testing. 23 23 there was another scenario, i.e., like the one we Q. And you didn't rely upon any testing in 24 24 doing your report, correct? talked about a minute ago where, if there was a 25 25 A. Correct. complete shutdown of the restaurant and no Page 110 Page 112 1 Q. Okay. What did you -- and, in fact, let's individual that had CoV-2 and/or the ultimate 2 2 go back. I want to make sure. sickness, COVID-19, was present, the virus would not 3 3 be present. Is that --Following your telephone conference, when 4 MR. ALVENDIA: Objection to form. you started with the lawyers and you started to do 5 When you say the word "shutdown," you your -- to draft the report, you learned about the 6 characteristics of Oceana Grill? mean shut down to the public or shut down 7 to the employees, the staff, the vendors, A. Yes, sir. 8 Q. You reviewed studies about the durability and so forth? I just want clarification on 9 of the virus? that. 10 10 A. Yes, sir. MR. MILLER: Okay. 11 11 BY MR. MILLER: Q. Are all of the studies regarding the 12 12 durability of the virus contained -- that you relied O. No one is in the restaurant --13 13 upon, contained within your report? MR. ALVENDIA: No human --14 14 A. Are all of the studies -- all of the BY MR. MILLER: 15 15 Q. -- that has -- no one is in the studies that I have relied upon in writing the 16 16 report are in the report. restaurant --17 17 Q. That was my question. MR. MILLER: Let me ask the question 18 18 A. Okay. to the witness, Rico. You can put your 19 19 Q. There aren't any studies outside of what objection on the record, and then if the 20 20 vou have identified for us -- there are no studies witness doesn't understand, he can tell me. 21 21 outside of this report that you're going to come MR. ALVENDIA: You understand the 22 back later on and say, you know what, I also relied 22 question, Doctor? 23 23 upon this? THE WITNESS: I need to hear it again. 24 24 A. Well, that's a little different, so let me MR. MILLER: I'm happy to repeat it 25 25 just be absolutely clear. Again, all of the -- my for you.

Page 113 Page 115 1 1 report is based on, and only on, the studies that Q. And that -- okay. 2 are referenced. A. And also, I assume we're not going over the 3 universe of -- the universe of what I reviewed was O. Okav. 4 A. Since then, I have read other studies that in the -- is in the references, but I also read the 5 I think buttress my findings, but those are not interrogatories where I learned that there were 6 referenced in my report and the report isn't based actually individuals who were in Oceana who had 7 tested positive for CoV-2. 8 Q. Are any of the other studies that you've MR. MILLER: All right. Let's go 9 read -- do any of those studies drive you to a ahead and pull up Dr. Moye's report. 10 10 different conclusion that is in your report. (Exhibit No. 2 was identified.) 11 A. They do not, sir. 11 BY MR. MILLER: 12 12 Q. So then all of the studies that you have Q. Before we get into the -- the nuts and 13 13 read since you authored your report lend you to the bolts of the report, Dr. Moye, did you speak with 14 same exact conclusion that we have from your report 14 any of the representatives from Oceana, not the 15 that has been submitted in this case? 15 lawyers but the owners of the restaurant? 16 16 A. That is correct. A. This is -- I'm sorry. Ask me the -- the 17 17 Q. Okay. And those studies generally dealt first part of your question again. 18 with the durability of CoV-2? 18 Q. Sure. Did you speak with any of the 19 19 owners -- have you ever spoken with any of the A. Actually, not. 20 20 Q. Okay. owners from the restaurant? 21 21 A. Those studies provided information about A. No, I have not. I did listen to the 22 22 the interaction of this virus with other organic deposition of one of them last week. I think it was 23 23 compounds, so that's on the physics/organic last Wednesday. I don't remember for sure, but I 24 chemistry section. 24 haven't directly spoken with any of them. 25 25 Q. All right. Q. And in preparing your report, which was Page 114 Page 116 1 1 A. And other studies, one other study in clearly before the deposition last week, you did not 2 2 particular, informed me about the -- let's call speak with any of the -- any representative, other 3 3 it -- well -- well informed me about the presence of than the lawyers, in preparing your report? 4 the virus in the vicinity of Oceana. A. No, sir, I did not. 5 Q. Okay. I know you were provided with the O. What study was that? 6 6 A. That, I think, is -- I think that it's interrogatory responses. Was that before you 7 Haverly. I can -- I can look to be sure, but I drafted your report or after? 8 8 think it's Haverly. A. I would say early in the process of me 9 Q. And that study is not referenced in your working on the report. 10 10 Q. Okay. Were you provided with anything report? 11 11 A. Correct. else? 12 12 Q. Okay. But it dealt primarily with A. Well, I was provided with the arguments for 13 13 geographic location? summary motion made by all parties. 14 14 A. It -- it dealt with the density of cases --Q. Summary judgment? 15 15 Q. Okay. A. I beg you pardon. I'm -- I'm not using the 16 16 right word. Thank you. A. -- in New Orleans. 17 17 Q. In the New Orleans area. All right. O. Did that --18 18 And in preparing yourself to write your A. Go ahead. 19 19 report, you looked at these studies, you considered Q. How did that influence your report at all? 20 20 surface impact, you learned what the mayor and the A. Well, I don't think it influenced my 21 21 governor said and how that would impact your report. It just gave me a more complete 22 22 understanding of the issues before the Court. conclusion? 23 23 Q. Okay. Anything else? A. Yes, sir. 24 24 O. Looked at Oceana's location? A. I was provided a map, which actually is in 25 25 A. Yes, sir. the report, of the location of Oceana Grill in

his testimony.

Page 117 Page 119 1 1 New Orleans. You can answer doctor. 2 2 Q. Anything else? A. Actually, I think what I said was you can 3 3 A. I don't think so. They may have provided divide my report into two components. The first 4 an article or -- one or two articles for me, but component is epidemiology, and that's how we get to 5 5 most of the articles I identified on my own. the number of cases around Oceana. Epidemiology 6 6 Q. Okay. Now you've never taught a chemistry gets us to the number of cases. Epidemiology gets 7 course before, have you? COVID through the door into Oceana. Physics and 8 8 A. Have I ever taught a chemistry course? organic chemistry get the virus to the surfaces. 9 9 I've never formally taught a chemistry course. I've BY MR. MILLER: 10 10 certainly tutored many, many students in chemistry, Q. Physics and organic chemistry get to 11 11 but I've never formally taught a chemistry course, damage? 12 12 A. Fair, yes. 13 13 Q. Okay. Epidemiology -- okay. That --Q. And -- and you've never taught a physics 14 14 that's fair. course, have you? 15 A. That would be the same answer. No, but I 15 Epidemiology gets CoV-2 into the 16 have tutored many, many students in physics. So 16 restaurant? 17 17 A. Yes, sir. 18 18 Q. Because of all of the reasons that we're Q. And I know I asked you --19 19 A. I just tutored a student in physics about going to go through in your report? 20 20 two months before I heard about this case. A. Yes, sir. 21 21 O. I know I asked you earlier and you said you Q. Physics and chemistry create the damage --22 22 didn't consider vourself a physicist. You also the loss of use -- the transformation, then loss of 23 23 use, and therefore damage? wouldn't consider yourself a chemist -- do you? 24 2.4 A. Yeah. Damage, which is transformation and A. I would -- I would answer this way. I 25 25 certainly do not hold myself out as an expert in loss of use, yes. Page 118 Page 120 1 physics or an expert in chemistry. I certainly Q. Okay. I understand where we are. 2 2 don't do that, however, I have studied physics and Okay. Let's take a look down at No. 6 of 3 3 chemistry for years, taken exams in physics and your report. It discusses research experience, and 4 chemistry, use physics and chemistry as the basis of two-thirds down in No. 6, you indicate that -- and my understanding of the human body. So I would say we've talked about this, quote, [As read]: "I 6 that, while I am not a -- I'm not an expert, I supervised a collection of adverse events that were 7 understand it. reported to both sponsor -- to the sponsor and to 8 8 Q. What specific discipline are you being the F -- the Federal Food and Drug Administration, 9 9 offered as an expert in in this case? FDA." 10 10 A. Well, I can tell you what I think I'm being Do you see that? 11 11 offered as an expert in. And I'm being offered as A. Yes, sir. 12 12 an expert in medicine, in epidemiology. Q. That FDA work is what we discussed earlier 13 Biostatistics, I'm not -- I don't really see plays a 13 where you were -- and just refresh me. I don't want 14 role here, but my -- and my background in chemistry 14 to mischaracterize what you did. 15 and physics. 15 A. Okay. So this is when I was at -- in my 16 Q. Well, now if you're saying you're being 16 work as a member of the Data Coordinating Center. 17 offered as an expert in epidemiology, when I asked 17 Q. Right. 18 you about your report, you made clear to me that 18 A. We collect safety information and per 19 your report had no basis in epidemiology. 19 regulations report that safety information to the 20 A. Actually --20 sponsor of the study, which may be private or it may 21 MR. ALVENDIA: Objection --- objection 21 be NIH, and also the FDA. 22 to form. 22 Q. All right. In a calendar year, when you 23 THE WITNESS: Actually --23 were doing that work, how much time did vou spend 24 MR. ALVENDIA: Mischaracterization of 24 doing this work for the FDA? 25 25

A. Okay. Just so I can be clear. I -- I am

Page 121 Page 123 1 1 working at the Data Coordinating Center meeting my A. Well, let's see. No. There is a -- I 2 responsibility to the FDA (inaudible). think the first one -- I actually can't recognize it 3 3 O. Uh-huh. from this, from the abbreviations, but I think the 4 4 A. Okay. So now your question to me is what? first one was a drug company on drug company issue. 5 5 Q. Well, how much of that work specifically I think it was a patent issue, so both were drug 6 6 related to any reporting to the FDA? companies. 7 A. Oh. Maybe 15 percent of my time. O. Okav. So what --8 8 Q. In a calendar year? A. But the others were --9 9 A. Just an -- just an approximation, yes, Q. Let me ask you, what type of opinion were 10 10 15 percent, 20 percent. you asked to render in the case entitled Eli Lilly 11 11 Q. Were you given a per -- per diem for that and Company versus ICOS Corporation? 12 12 A. Well, if that's the -- the case I remember, work? 13 13 A. No. it was a case involving patents, and I was asked to 14 14 Q. That -- that was part of your salary with review the content and the data analysis of a 15 15 particular patent and render an opinion. I don't what organization? 16 A. Yeah. So that was part of my salary with 16 remember the details about it. 17 17 Q. All right. But it was a data analysis the University of Texas. 18 18 engagement? Q. Okay. 19 19 A. However, I -- however, when I receive a A. Yes, sir. 20 20 large grant, the -- the organization paying for the Q. Okay. And the -- the next case, Abbott 21 21 Laboratories case? grant gives money to the University of Texas to pay 22 22 my salary. So it's all salary support, but the A. Yes. 23 23 University gets money from the paying --Q. Do you recall the type of opinion you were 24 24 Q. Organization. asked to give? 25 A. Organization, correct. A. I believe that's an Actos, the antidiabetic Page 122 Page 124 1 1 Q. Okay. All right. Let's take a look at drug Actos and the relationship with bladder cancer. 2 2 paragraph 18. Fees. Does -- does paragraph 18 Q. And did you represent the plaintiff there? 3 3 accurately reflect the fees that you are being A. Yes. 4 paid -- paid by the lawyers in this case? Q. And the next one is products liability A. It does. litigation? 6 6 Q. Okay. Is there any variation of fees? Is A. Yes. Oh, that was Mirena, a IUD, and there any bonus for outcome? Are any of your fees pseudo -- pseudotumor cerebri, which is a form of intercranial hypertension. outcome related? A. They are not, however, I do reduce my fees Q. What type of opinion were you asked to 10 10 in some circumstances. For example, if I'm render? 11 11 listening -- if I'm listening to a deposition, I A. Oh, I was -- both in Abbott and in Mirena, 12 12 cannot in good conscience charge somebody \$400 an I was asked the epidemiology question, is -- is 13 13 there a relationship between the exposure, in this hour to listen to it, so I discount my fees. 14 14 case the medicine, and the disease? One case it was Q. That's very nice of you. 15 15 A. Well, I -- I -- I appreciate the value of bladder cancer, the other court case it was a 16 16 pseudotumor cerebri. Was it an association or was money. 17 17 it causation? And I reasoned in each time it was Q. In No. 19 you list a number of cases that 18 18 you have been involved in; one, two, three, four, causation. 19 19 five -- six cases. Is that the universe of Q. Got it. What about the next case, 20 litigation that you've served in an expert capacity? Daniels-Feasel versus Forest Pharmaceuticals, Inc.? 21 21 A. No. It's only the -- for the past four A. I have been trying so hard the past five 22 22 years. That's all that I believe I've done in the minutes to remember this one, and I don't. I'm 23 23 last four years. 24 24 Q. In any of those cases that are listed in Q. Okay. That's fine. 25 25 No. 19, did you act on behalf of the defendants? A. If my memory gets refreshed later on, I'll

Page 125 Page 127 1 1 let you know. But right now, I don't. and solid surfaces within Oceana. 2 2 Q. What's this Pradaxa, October 18, 2019? Q. Okay. And, again, in the next sentence you 3 3 A. Oh, Pradaxa. That was a -- yeah, sorry -referenced -- you referenced the things that you 4 that was a -- I'm sorry for the incomplete relied upon in coming to your conclusion. 5 5 A. Where are we in my report? descriptor. 6 6 But that was a case we were looking at the Q. Second -- second sentence in paragraph 20. 7 7 anticoagulation effect of Pradaxa and its It says, "To draw my conclusions, I relied upon my 8 8 relationship to cardiovascular (inaudible). training in medicine, my experience seeing and 9 9 Q. And you represented the plaintiff there? treating physicians, my advanced training in 10 10 epidemiology and biostatistics, my public health A. Yes. 11 11 experience, and my clinical research experience as Q. And I apologize for butchering that word. 12 12 applied to my review of the literature." And lastly this -- the Johnson and Johnson 13 13 case? A. Yes. 14 14 Q. Do you see that? A. Yes, sir. That involves the relationship 15 between Motrin, ibuprofen, and Stevens-Johnson 15 A. Yes. 16 Syndrome; and I concluded that there was a causal 16 Q. Okay. And not to quibble with words, but 17 17 association between Motrin and SJS. at the time you started drafting your report, you 18 18 hadn't reviewed any -- at the time you came to your Q. Is there any specific reason why you only 19 19 listed -- list the last four years of cases? conclusion, you hadn't reviewed any literature, 20 20 A. Well, that's what I thought -- well, okay. right? 21 21 I guess a couple of reasons. One is that's what I A. At the time I came to my conclusion? 22 MR. ALVENDIA: Objection. thought the -- the custom and culture were, and also 23 23 I'm not sure I could remember all the cases prior to BY MR. MILLER: 2.4 24 the past four years. Q. Yeah. Yeah. December 10th to 25 Q. Okay. All right. Have you worked December 17th, when you had your telephone Page 126 previously for any of the lawyers that are involved 2 2

Page 128

in this case?

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A. The only lawyer I worked with before was Dave Matthews, who I think I mentioned earlier. But not with -- I have not worked with anybody from the Houghtaling group or Rico.

Q. Okay. In preparing your report, did you ask for any information that was not provided to vou?

A. Well, I did not -- I don't think I asked the attorneys for any additional information. I don't think I did. I used my own resources to find what manuscripts I could. I may have asked the attorneys one time to direct me to the location in the interrogatories that discussed the number of people who entered the Oceana facilities with CoV-2, but I -- I -- as I sit here, that's all I remember.

Q. Okay. Let's talk about paragraph 20. Now, you used the word or the phrase "environmental impact" a few times throughout the report, or the word "environment."

Could you tell me within the context of this report, what environment are you talking about?

A. Sure. I'm talking about -- specifically, I'm talking about air and surfaces. The atmosphere conference with Mr. Houghtaling and Mr. Alvendia, you communicated to them your conclusion about whether or not CoV-2 created a physical transformation loss of use and therefore damage?

A. Correct.

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MR. ALVENDIA: Wait. Wait. Objection to form; asked and answered. Now, he's -- he's said what -- what he relied upon in his conclusion was everything he just said and his years of experience, and I'm trying to understand -objection to form.

MR. MILLER: Okay.

BY MR. MILLER:

Q. So, Doctor, let me get back to paragraph 20 in the report. You relied upon all of your experience "and my clinical research expertise as applied to my review of the literature" to draw your conclusion.

That last sentence, "as applied to my review of the literature," you had yet to review the literature when you came to your conclusion?

A. I will -- I would say a couple of things about that. I had not reviewed any of the literature on -- on SARS-CoV-2, and it's -- in

32 (Pages 125 to 128)

relation to Oceana, I had not reviewed any of the literature.

However, before they asked me the question,
I did understand SARS-CoV-2, because I had read a lot of literature up to then. I also rely on my experience in the physics and organic chemistry, which I should have said here but I did not.

But all of that was in play, all of that was my background --

O. And --

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A. -- when the question was posed to me about damage.

Q. And, in fact, you really did not need to read literature specific to CoV -- CoV-2 to formulate your conclusion. You could have read literature related to generally any virus, even the flu. I think we talked about that earlier.

A. I would -- I would say that -MR. ALVENDIA: Wait. Wait. Wait.
Let me object.

Objection to form.

And, Doctor -- you can answer the question, Doctor.

A. I -- I would say this: To some extent, viruses are viruses. They have common

some of the difference. There has certainly been a lot more attention given, of course, to the pandemic causing CoV-2 than there has been to the flu, but in general it's a little longer.

Q. Okay. But putting -- putting durability aside, exposure to a physical substance, all things created equal, meaning temperature, humidity, all things created equal and the initial exposure by both viruses creates physical change that results in physical loss of use and then damage?

A. We're talking about H1N -- I'm sorry. We're talking about the flu virus and --

O. Versus --

A. COVID-2.

Q. Yes, sir.

A. Right. Right. Essentially damage is both in -- both make physical transformations -- how deep the physical transformation is in the substance depends on the size of virus. The virus is of different sizes, then the extent of the damage will be different.

Q. Got it. Okay. All right. Let's get back to your report. The last sentence of paragraph 20, and the words are in quotes -- I'll just read the entire sentence. "Determining whether SARS-CoV-2

Page 130

characteristics across the -- well, the billions of different species, they have common characteristics, so yes.

However, there are individual characteristics that have to be taken into account, and one of them is how long the virus survives outside the body. Now, that answer may be different for some viruses than other viruses. It turns out it is, not by very much.

BY MR. MILLER:

Q. Okay. So -- and just generally speaking, unrelated to your report, and I know you've done extensive research, so I'm asking for -- really for my own edification.

What is -- is the durability of COVID-2 in comparison to the durability of what we affectionately call the flu?

A. Okay. Well, the -- the durability of COVID-2 is a little longer than the durability of the flu -- a little longer.

Q. Okay. A little by -- by what number?

A. By days. By days. A little longer. It depends on the -- the experiments that look at the durability of the flu versus the experiments with CoV-2 are different in design. That could explain

Page 132

Page 131

impacts its surrounding environments, meaning air and surface, is an exercise in critical thinking using the weight of the evidence approach."

All right. And you have that in quotes, to me, meaning it is a term of art?

A. Fair.

Q. All right. So tell me exactly what the weight of the evidence approach is from your perspective?

A. Actually, I was just going to read paragraph 21 back to you. That's what it is.

O. All right.

A. Okay. You --

Q. Go ahead and read it to me.

A. Go ahead and read it to you or not?

O. Yes, sir.

A. Okay. "The weight of the evidence approach is the process by which a body of evidence is examined piece by piece, each component being sifted and assessed using a transparent and standard methodology. In this case, the goal is to assess the relationship between SARS-CoV-2 and its environment."

Q. All right. The body of evidence that we're talking about here are the things that we kind of

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generally outlined earlier, but we're going to see in more detail if all of the factors you considered in your -- in your, quote, "weight of the evidence," which would include all of the -- the articles that you reviewed and your experience and so forth?

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O. Okav. Is there any evidence that was weighed to come to your conclusion in formulating this report that is not evident, other than the articles we talked about earlier?

I know you said since you've drafted the report you've reviewed a couple of articles. I'm not talking about those.

A. All right. I understand.

Q. I just want to confirm that the weight of the evidence to support your conclusions is within the corners of this document?

MR. ALVENDIA: Now, objection to form. Once again, he did testify earlier, Allen, that there are several articles that's he's done research on that he relies upon and will rely upon at trial.

A. I would agree with you with the -- we need to be sure that we include physics and organic chemistry.

Page 135

- A. Yes. But I don't remember -- I just remember I read them, saw they weren't peer reviewed and then discarded them.
- Q. Okay. Did any of those papers disagree with your conclusions that are evidenced in this report?
- A. I -- I don't know. Once I saw they weren't peer reviewed -- I mean, for some of them I maybe didn't even finish the paper. Once I saw it wasn't peer reviewed, I just rejected it out of hand. It may have supported me, it may not have. I just rejected it.
- Q. The Lancet, that's a -- those papers in the Lancet are peer reviewed. Is that correct or not?
  - A. My understanding, yes, that's right.
- Q. Okay. Because, I mean, you actually were one of the reviewers?
  - A. For Lancet papers, yes.
- Q. Yes. All right. 23, "Following this process, I identified the lines of evidence created in each study assessing the methodology used by the research effort. Since the value of a research result is tightly circumscribed by the methodology used to generate it, research methodology is critical. Each manuscript is assessed on its

Page 134

BY MR. MILLER:

- Q. Of course. Yeah, because without those there's no damage?
  - A. Right, correct. Correct, yes.
- Q. All right. In paragraph 22 you indicate that you identify -- you identify the collection -a collection of peer reviewed published papers reflecting the universe of useful information about the state of the relationship between SARS-CoV-2 and its environment.

When you say the universe of useful information, you're not contending that the articles that you have relied upon in your report are the only articles related to SARS-CoV-2?

- A. No, I'm not. But I am rejecting that, for example, what's happened with CARS -- SARS-CoV-2 research is that main -- in the rush to get information out there about this virus, there are papers that are not peer reviewed that are now appearing in literature, and I had to be careful to make sure not to include those.
- Q. Okay. Is there any specific papers that vou've read, because clearly this insinuates that you read some papers that were not peer reviewed, that you specifically excluded from your analysis?

Page 136

merits. It is rare -- it is the rare manuscript that provides contributions in all aspects of SARS-CoV-2/environment relationship."

Tell me what you're telling me right there.

A. Oh, I wrote that so eloquently too.

Q. I don't disagree.

A. Here we go. Here we go. Okay. It is one of the biggest mistakes in research in general, medicine in particular, to be persuade by the results of a study without absorbing the methodology, because results can be very misleading and one has no idea whether they are being misled or not, unless they understand the methodology.

The methodology is the prism to which we under -- review results.

O. Okav.

A. Some of the biggest embarrassments in medicine have been findings that were not based on solid methodology, but were attractive and the field ran with them.

This paragraph right here actually is the -- it is or should be or better be the creed of epidemiology. You have to understand the methodology before you can draw a conclusion.

Q. That makes sense.

Page 137

A. I'm glad, because some people want to throw me out of the room when I say that.

- Q. And so in 23 the statements are less about the methodology that you utilized in coming to your conclusions, but in assessing the -- the works that you consulted to determine how they affected your conclusion?
- A. Yes. I asked myself -- it's -- it's my -- it is -- the metric I used to assess the contribution of a manuscript to my point of view or not
- Q. Notwithstanding your own -- notwithstanding your own methodology?
- A. Correct. Correct. I mean, I'm describing my own methodology here. This is the metric I use. So if people disagree with this metric, then they're not going to accept my results.
- Q. But, Doctor, isn't it fair to say that before you employed your methodology -- I'm sorry -- yes.

Before you employed your methodology of meticulously evaluating the works that you would consult, you had already made you conclusion in October -- between October 10th and October 17th about damage?

Page 139

Page 140

understand kinematics. I already understand the appropriate physics. I understand the organic chemistry. I also understand viruses. So I can certainly opine about that because I have already assimilated the appropriate (inaudible). BY MR. MILLER:

- Q. Okay. I understand. Okay. The works that you rely upon then and that are contained within the report, and we'll get to them in a second, each of those had a methodology that you found to be sound as it applied to Oceana Grill restaurant?
- A. Yes. I mean, some stronger than others, but yes.
- Q. Okay. All right. We're going to skip 24, because I think what 24 is saying is that in your experience in epidemiology, the methodology in which you gather evidence to support your conclusions is sound.
  - A. Yes.
- Q. All right. Virology, we'll skip that. In 26, talking about viruses you state that they're -- they're not self-propelled but travel through the air or water, or access solid objects to rest on the surface, waiting for and in order to be absorbed into a cell?

Page 138

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## Q. Why don't you, in lay -- in layman's terms, tell me what that means?

A. Oh, of course. Okay. So viruses are not alive. They're on the verge of life, but -- and therefore not being alive they don't swim, they don't fly. They are -- they are the victims of physics and chemistry, which means they land someplace based on physics and chemistry.

Now, the only thing viruses do when they land is attach and -- I don't want to use the word "try," because try suggests intent -- but they open up and their RNA is to pour out of them into a cell.

Now, most viruses aren't so lucky. Most viruses -- as I'm sitting at a wooden table, there are viruses landing on that table, opening up, letting their RNA -- their DNA get in the table. It's not going to infect anything, they're not going to reproduce. They don't know better. That's all they do.

So viruses are the -- their movement and their attachment is not purposeful, and it's very easy, of course, when people are dying to anthropomorphize them. But they are simply victim -- they are simply -- not taken advantage of,

MR. ALVENDIA: Objection --

A. It is not fair --

MR. ALVENDIA: Wait. Wait, Doctor. Objection --

THE WITNESS: Okay.

MR. ALVENDIA: Objection. He's already gone through the entire process of how he reached his final conclusion in his report. The fact that he gave his initial opinion in the initial meeting does not mean he reached the final conclusions in his report.

He's got several conclusions in this report by the way, several opinions. So objection to form.

A. The question that was asked of me was the lawyers -- about lawyers, was not a question about whether CoV-2 was in the vicinity, though I had to ultimately address that. It wasn't a question about whether Oceana [sic] was in the building. I had to address that. The question I was asked was given the assumption that CoV-2 is in Oceana or in the building, will it cause damage.

Now, I already have a background in that. I already have a background in that. I already

but caught up in laws of physics and laws of organic -- organic chemistry, and -- and air movement -- and, of course, air movement. Q. What is the difference between RNA and DNA? A. Sure. They are not nucleic acids. They are essential for the reproduction of cells. Classically, and this has changed the past couple of years, but classically DNA is called

I'm going to try to do this from memory, but I think if you took, Mr. Phelps, one of your cells or one of my cells and took the DNA out and unraveled it, it would be five feet long. That's how tightly coiled it is. That reproduces itself as the cell reproduces, and so that's -- that's what

deoxyribonucleic acid. DNA contains the content of

all of our genetic information. Every cell has its

own copy of DNA. It is tightly coiled. It's in the

### Q. Okay.

DNA is.

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A. And the information -- the information in the DNA has to be turned into something, right? It's got to be used. So we make -- you and I make proteins. We make polysaccharides. We make hormones, enzymes. That information is in the DNA.

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### Q. So, Doctor, do you have a hard copy of your report in front of you?

A. I can get an electronic copy in front of me, yes.

BY MR. MILLER:

Q. The only reason I ask is because, as I go through the remainder of the report, I'm going to talk about some of your findings, and I'm also going to reference the footnotes and the articles you use as support. And so those are found at the end. It would just be easier if you had a copy in front of

A. Okay. Just one second. It won't take long.

### Q. And, Doctor, if you would just go to the references at the end of your report, then we can keep on the screen the actual paragraph that we're talking about.

A. So I have -- so I have an electronic version of my report right here, and I am in the

Q. All right. So that's -- paragraph 33 of the report says SARS-CoV-2 virus is 125 nanometers in diameter.

A. Yes.

Page 142

It has to get out.

I'll trying to keep it short. RNA helps to get the information out where that information can be converted to the molecules you and I need to live. That doesn't help, I'll try it another way.

### Q. No, that's fine.

### Does a virus contain both RNA and DNA?

A. To my knowledge, no. But I -- I will tell you, viruses surprise us, so sure as -- and there are, as I said, hundreds and thousands and millions of species, so maybe there's one that does. But by and large, they have only DNA or only RNA.

### Q. Okay. What does CoV-2 have?

A. CoV-2 is RNA virus. Probably one of the largest RNA viruses known.

> MR. MILLER: Does anybody need to take a break?

THE WITNESS: Let's see. We've been going for 55 minutes --

MR. MILLER: About an hour. THE WITNESS: -- or so. Yeah, I'll take a break for five.

MR. MILLER: All right. Let's take five minutes.

(Recess taken.)

Page 144

Page 143

### O. Where does that -- where does that information come from?

A. Actually, there are multiple sources. I kind of thought that was just common knowledge and didn't need to provide that, but the -- it's a -it's a virus that ranges in size from 60 to 120, 125 nanometers.

### Q. Okay. So it's not a virus is 125 nanometers; it varies in size?

A. Yes, sir. Every species of virus has a different size range.

### O. Okav.

A. The size range for SARS-CoV-2 is 60 to 125.

### Q. Take a look at paragraph 35. Paragraph 35 generally talks about the effect of COVID-19 infection. What are -- what is the purpose of this with respect to your report?

A. I'm just providing the signs -- for the -for the Court, I'm providing the signs and symptoms of COVID-19. Why is it a disease? Why are we so concerned about it? So I'm just providing this really as background.

### Q. Okay. Let's look at paragraph 37. Why is this necessary for your report?

A. Well, this is necessary because it

Page 145 Page 147 1 demonstrates, number one, the seriousness -- how A. I do. Q. If the mayor's proclamation didn't affect serious the elected officials, the governor and --3 of Louisiana, the mayor of New Orleans -- of your conclusions at all, why was it necessary to put 4 New Orleans took the problem, took the COVID-19 it in this report at the end in your conclusions? 5 infection, number one; how they were beginning to A. It was necessary for me to reverse --6 track the COVID-19 infection, the COVID-19 illness, reverse the given here. What I'm doing is saying, 7 number two; and then with the driving force behind based on my understanding of contamination of 8 the decision to close the bars and restaurants, to surfaces and based on my understanding of causing 9 go through this closure and period of property loss and damage, there was scientific 10 10 staged-to-phase openings. support. 11 11 Q. Paragraph 39 references the March 16th, Q. Okay. 12 12 2020, proclamation from the mayor. And you quoted A. I -- that -- that -- that was my reasoning, 13 13 in your report, and it reads, "There's reason to and therefore since they came to the same conclusion 14 14 I did, then that's the basis for my consideration. believe that COVID-19 may be spread amongst the 15 15 Q. Do you know whether or not Mayor Cantrell's population by various means of exposure, including a 16 16 propensity to spread person to person and a proclamation on March 16, 2020, had any scientific 17 17 propensity to attach to surfaces for prolonged basis whatsoever? 18 18 periods of time, thereby spreading from surface to A. I don't know one way or the other. 19 19 person and causing property loss and damage in Q. Okay. Let's go back to paragraph 44. 20 20 certain circumstances." MR. MILLER: Allison, you're not on 21 21 How did this proclamation by Mayor Cantrell mute. Allison, you're off -- you're not on 22 mute. Allison, you're not on mute. affect your opinion? 23 23 A. Well, this proclamation -- it affected my DR. STOCK: I should be -- okay. I'm 24 2.4 opinion in the following way. Number one, that trying to get back on. I thought I was. 25 25 COVID-19 was -- was a disease that was becoming Sorry. Page 146 Page 148 1 increasingly prevalent given it's propensity to MR. MILLER: Okay. 2 MR. ALVENDIA: Allen, are we waiting spread. Secondly, she described to a T the epidemiologic point of view about how it spread 3 on something? 4 person to person and also surface to person. MR. MILLER: Yeah, my thought process. 5 Q. What about the -- the phrase "and causing MR. ALVENDIA: Okay. All right. I'm property loss and damage in certain circumstances"? sorry. I thought you were waiting for her A. Let me just find that. Oh, yeah, "causing to get back on. property loss and damage." MR. MILLER: Let's look at 9 That really didn't affect my opinion. It's paragraph 44. 10 10 for me to determine, based on science, whether BY MR. MILLER: 11 11 there's loss and damage. I respect her point of Q. And generally you talk about some of the 12 12 view, but it really is -- I mean, my understanding orders. The last sentence states, "The mayor stated 13 13 of my role in this case is not to be influenced by that restrictions were particularly aimed at Bourbon 14 14 what her perspective is, but to generate my own Street and other areas of the city where 15 15 perspective. alcohol-fueled gatherings have," quote, "gotten out 16 16 of control," closed quote. Q. Turn to the very last person of -- last 17 17 We -- I think we established earlier that page of his report, the conclusions. 18 18 Conclusion E -- and we'll come back to the Oceana Grill is on Conti, but it's close -- it's in 19 19 rest of it, but I just want to talk about this since the French Quarter, right, and -- and relatively 20 20 we're on this topic. "It is more likely than not close to Bourbon Street? 21 21 that the mayor and governor's reason that SARS-CoV-2 A. Yes, sir. 22 22 attach to surfaces, contaminate surfaces, and causes O. You're aware of that? 23 23 property loss and damage was scientifically A. I am, sir. 24 24 supported." Q. Are you aware --25 25 Do you see that? MR. ALVENDIA: And actually, Allen,

Page 149 Page 151 1 1 objection to form. Just -- just so we know read this way, and my apologies because it's my here, it's on the corner of Conti -fault it doesn't -- "Based on notice of the change 3 MR. MILLER: On the corner, yeah. in the property condition from safe to dangerous, 4 portions of Oceana Grill were closed for 24 hours." MR. ALVENDIA: Touches it, right. 5 5 It's on Bourbon. But go ahead. Q. Okay. "The dates of these actions on or 6 6 MR. MILLER: I don't spend much time about April 3rd, April 10th, June 3rd, June 27th, on Bourbon, so I don't know. But I -- I'll and August 1, 2020." 8 8 A. Right. It should be "The dates of these take your word for it. 9 9 MR. ALVENDIA: Oceana -- and I've been actions were on or about." 10 10 to Oceana before, and I can tell you it's O. Okav. 11 11 on the corner. A. Sorry. 12 12 MR. MILLER: All right. Q. Now, the change in the property condition 13 13 BY MR. MILLER: from safe to dangerous is the fact that individuals 14 Q. Do you know what -- whether or not Oceana 14 who contracted COVID-19 -- that you were informed 15 followed the mayor's orders? 15 that individuals that contracted COVID-19 were in 16 16 A. I believe I do. the property? 17 17 A. Yes. Q. All right. 18 A. They followed the phase -- the shutdown and 18 Q. The statement "change in property and 19 19 the phased reopening. condition from safe to dangerous" is not the result 20 20 Q. Do you know if Oceana was ever in violation of testing of the property condition at Oceana 21 21 of the mayor's orders and had to be reprimanded by Grill? 22 22 the city for failing to file -- failing to follow A. That is correct. 23 23 her phased shutdown orders? O. Now, these -- the areas of Oceana Grill 24 24 A. I -- I don't know. I -- I imagine were closed for 24 hours. Do you know what Oceana 25 25 initially, like in Phoenix where I am and other did for -- during those 24 hours to the property Page 150 Page 152 1 cities, there was confusion the first couple of days condition? 2 2 about when do orders go into effect, who's involved A. I would have to check later in my report, with that, and what should they do. Are they 3 because I do talk about remediation. I mean --4 shutting down completely? I imagine there was some excuse me -- cleaning and disinfecting later in my confusion about that. Other than that, I don't report. But I don't remember right now whether they 6 were closed and they went through cleaning or not. 7 Q. Okay. Let's look at paragraph 47. I just don't know. MR. ALVENDIA: And, Allen, do me a Paragraph 47 reads, [As read]: "The Oceana Grill 9 had patrons who had SARS-COVID-2 infections, favor -- look, could we break this down 1 0 10 specifically the owner tested positive for COVID-19. into time phases or time periods? Because 11 11 In addition, there were four other instances where clearly, in the different phases -- I'm 12 12 individuals with property access reported a positive just trying figure out a way here to --13 13 to --SARS-COVID-2 positive test. These two were -- these 14 14 MR. MILLER: I'm just reading his were two office employees, a maintenance worker, and 15 15 report. a prospective employee to be -- interviewing for a 16 16 position. Based on" -- I think that's -- you meant MR. ALVENDIA: The problem with it --17 17 the problem with it is it's a compound to say, "Based on notices of the -- based notice of 18 18 question though. I mean, you're asking him the change in property condition from safe to 19 19 what they did in response to a dangerous dangerous, portions of Oceana Grill were closed for 20 20 condition. I just want to be sure we're 24 hours." 21 21 talking about a time frame, because they Tell me exactly what you meant there, 22 22 went through different restrictions at because it looks like a typo. 23 23 different times. A. Good point. Let me look at this. This is 24 24 MR. MILLER: Okay. Well, his report not the best sentence I ever put together. 25 25 says that "Based upon the change in Oh, okay. I think it reads -- it should

Page 153 Page 155 1 1 property condition from safe to dangerous, Q. Okay. That's fair, because they could have 2 2 had -- if it was in a complete shutdown -they closed for 24 hours." 3 3 MR. ALVENDIA: Right. Then it goes on A. Right. 4 to say the dates of these actions, and then Q. Right. Okay. June 3rd, we weren't in a 5 5 complete shutdown in June 3rd. We were at least in he gives different dates, because it's in 6 6 different phases is what I'm saying, so... Phase 1? 7 MR. MILLER: Closing for 24 -- all A. Okay. 8 Я right. I'll -- I'll ask another question. Q. They would have closed -- the -- your 9 9 MR. ALVENDIA: Okay. Go to the next understanding from what -- what you were told and 10 10 sentence. It talks about the different thus transferred into your report was that they 11 11 closed a section of the restaurant for -- for phases. I'm just trying to make sure we're 12 12 not conflating that they were closed for 24 hours on June 3rd after being notified that 13 13 24 hours all the time. They went to someone had contracted COVID-19? 14 14 different restrictions on those dates that A. Yes. 15 15 Q. Had been in the building? he says in the next sentence. 16 16 But go ahead. I just want to make sure A. Yes. 17 17 we're not mischaracterizing his testimony Q. Same for June 27th, and same for 18 18 August 1st? to mean that's how they were the whole 19 19 time. But go ahead. A. Yes. 20 2.0 MR. MILLER: All right. Let me --Q. Okay. And you would expect that on these 21 21 MR. ALVENDIA: Doctor, do you dates, June 3rd, April 10th, April 3rd, June 27th, 22 understand his question? and August 1, that during the 24-hour period where 23 23 MR. MILLER: Let me ask the question. this area was closed, that Oceana Grill followed the 24 24 CDC guidelines and disinfected the area where this BY MR. MILLER: 25 Q. So, Doctor, what I think you're saying in person had been? Page 154 Page 156 paragraph 47, and correct me if I'm wrong, that A. I mean, I would like to think that. But 2 2 someone -- you were informed that someone notified sitting here now, I can't say that I know that 3 that's the fact. Oceana on April 3rd that they had contracted 4 Q. Okay. But you -- you would hope that's the COVID-19 and had been in the facility, and on 5 April 3rd Oceana closed the area where that case? 6 individual was for 24 hours? A. I would hope, yes. 7 A. Yeah. On or about April 3rd, yes. Q. All right. And the next paragraph talks 8 O. Right. And on or about April 10th, Oceana about the number -- where did you get this 9 was notified that an individual that had been in the information that's contained in paragraph 49? 10 10 facility was present, and they closed that area A. Actually, I'm trying to remember. I don't 11 11 where the individual was for a total of 24 hours? recall. I mean, it's not -- it's not in any 12 A. Yes. 12 published literature. That may also have come from 13 13 O. On or about June 3rd, Oceana was notified the interrogatory. 14 14 that an individual that had contracted COVID-19 had Q. All right. But you don't give any cite in 15 15 been in the facility, and on June 3rd they closed your report, so --16 16 A. Correct. the facility for 24 hours? 17 17 Q. -- do you know? Do you know whether -- do 18 18 vou know for sure where this information came from? Q. And in each instance after the 24 hours, 19 19 A. I don't know for sure, but I -- I believe it's your understanding that they reopened that area 20 20 of the building? it was from the interrogatory. 21 21 A. The degree to which they reopened that Q. Okay. Let's look at paragraph 50. 22 22 really depends on what phase the -- what phase of [As read]: "Given the established infected --23 23 infectivity of SARS-COVID-2 and the ongoing customer reopening or what stage of reopening they were in. 24 24 and patronage level of Oceana Grill, the degree of But there was some resumption of activity after the 25 25 environment exposure, air and surface at the 24 hours.

Page 157 Page 159 1 1 restaurant rose to dangerous levels." A. I believe it does, yes. How do you know that? 2 Q. Okay. Despite the fact that Oceana 3 3 A. I know that because patients came to CO --Grill -- despite the fact that they disinfected the 4 restaurant once they had knowledge in -- in 5 5 Infected patients came to -- patrons -accordance with CDC guidelines? 6 6 infected patrons came to Oceana Grill. A. I --7 O. Okav. MR. ALVENDIA: And you're asking --8 A. Infected patrons sneeze, infected patients again, your question -- your question, cough, infected patients speak and they spew out Allen, for the legal standard for this 10 10 viruses. They changes the environmental -- that expert is within a medical degree of 11 changes the environment to a dangerous level. 11 certainty more likely than not, correct? 12 Q. Was Oceana Grill, in your opinion, a 12 That's -- that's what you're asking him. 13 13 restaurant that had environmental exposure that rose MR. MILLER: No, that is not what I'm 14 14 to a dangerous level on April 5th of 2020? asking him. 15 A. Well, I'd have to look back to April --MR. ALVENDIA: Well, then ask --16 well, I can't. But April 5th is one of the days? 16 MR. MILLER: I asked him --17 Q. April 3rd is one of the days where an 17 MR. ALVENDIA: If you ask --18 18 individual was infected. MR. MILLER: You can -- you can argue 19 19 A. Okay. that to the judge about whether or not his 20 Q. That was in -- that had COVID-19 that came 20 answers fit within the legal standard, but 21 21 into the building. I'm not worried about the legal standard. 22 2.2 A. Okay. I would -- I would say this. Okay. MR. ALVENDIA: Objection --23 2.3 I don't know where that individual was in the MR. MILLER: I want to ask him 24 24 building, but where he was, his presence changed the questions. 25 environment to dangerous. MR. ALVENDIA: Objection. If you're Page 158 Page 160 1 1 O. Even after -- let's assume for sake of this -- objection to form. It's an improper 2 2 conversation that Oceana Grill applied all CDC question. 3 recommended disinfectants on April 3rd once they MR. MILLER: All right. And then you learned of this individual's presence in the can -- you can have that question stricken building. Okay? when we get to trial. 6 6 A. Okay. So -- so help me with the sequence MR. ALVENDIA: You can move on. now. I can answer your question. I need to know --MR. MILLER: All right. Can we go 8 make sure I know the sequence. back to my last question and the Doctor's A patient -- I mean, the subject comes in, answer, Madam Court Reporter --10 10 THE REPORTER: Yes, sir. they are infected. Okay. Now -- now, tell me what 11 11 MR. MILLER: -- so that I can finish happens. 12 12 this line of questioning? O. Oceana Grill sections off the portion of 13 13 the building where that individual was present. (Record read back by reporter.) 14 14 A. So my answer is pending. A. Okay. 15 15 BY MR. MILLER: Q. And performs all disinfectants recommended 16 16 by the Center for Disease Control for -- in that O. Yes, sir. 17 17 A. Okay. My answer is, yes. And it is no 18 18 A. Okay. disparagement of the cleaners. It's no 19 19 disparagement of the CDC. It's just that if their Q. On that -- on the -- on that day and the 20 following day, April 3rd through April 4th. It is guidelines are not sufficient to get rid of all of 21 21 the virus. So the area is still infected, so that's now April 5th. 22 22 A. Yes. why my answer is yes. 23 23 Q. And your answer would be the same if I gave Q. Does the restaurant still have a degree of 24 24 environmental exposure at the restaurant that is you the same scenario for April 10th, June 3rd, 25 25 June 27th, and August 1st, which are all instances considered a dangerous level?

Page 161 Page 163 1 in your report that you became aware that Objection. He's already answered that individuals with COVID-19 were present in the Oceana question. More likely than not is the **Grill restaurant?** legal terminology requirement. "The A. Yes. minimum" is what he said, and that's true. 5 Q. Okay. You go on to say, "The restaurant's Minimal standard -- the burden of proof in environment was transformed into a" --6 a civil case is more likely than not. He's 7 A. Deleterious. saving it's much more beyond that. He's 8 Q. -- "deleterious condition as the virus already answered this question. Objection. 9 physically transformed the air and the restaurant MR. MILLER: Thank you. 10 10 contents from one of safety to one of infectivity BY MR. MILLER: 11 and illness." 11 Q. You can answer my question, Doctor. 12 12 Tell me what that means. A. Well, I mean, it's the same thing. It's 13 13 A. It means that the presence of the virus in certain, therefore it most certainly is more likely 14 the air of the restaurant and on the surfaces of the 14 than not. 15 restaurant make this location unsafe, because people 15 Q. So were you instructed to include more 16 can get infected and ill from it. 16 likely than not into your report by the lawyers? 17 Q. Okay. "This transformation changes the 17 A. I was simply instructed to -- I -- I was 18 structure of the surface of the restaurant contents 18 instructed a number of times to make sure I 19 19 by a process predicated by physical law." understood what the legal standard was. 20 20 What does that mean? Now, I exceed -- so I exceeded the legal 21 A. "By a process predicted." 21 standard, in my estimation, using science and facts 22 22 Q. Predicted. I'm sorry, predicted. as I know how to use them. But I wanted to also be 23 23 A. Right. Right. I mean, the transformation sure that the language was consistent, as long -- as 24 24 long as it was consistent with my scientific takes place in the restaurant contents by organic --25 25 by chemical processes that are -- that are conducted opinion, I wanted to also be sure that it was Page 162 Page 164 1 by well-established physical law. consistent with a legal interpretation. 2 2 Q. So that's chemistry and physics that we're As long as there was no conflict, I have no 3 3 problem using more likely than not. But I think talking about? 4 A. Yes, sir. You have principally chemistry, we've exceeded that. Q. All right. So I get it, the lawyers yes. 6 6 authored this last statement to include the words Q. Now, "The change in the structure is the damage, so the transformation more likely than not "more likely than not"? leads to physical damage." A. Sir --Why more likely than not? MR. ALVENDIA: Objection. 10 10 A. Well, because the transformation is A. -- the only person who authored this is me. 11 11 damaging. The transformation happens. That MR. ALVENDIA: Objection to form. 12 12 transformation is the damage. So given the A. They're the only -- I'm the only one. 13 13 transformation happens, there is damage. Nobody touched a single letter on this report than 14 14 Q. So then why more likely than not? me. 15 15 A. Well, because, I mean, more likely than not BY MR. MILLER: 16 16 is a weak -- it's a weak metric here. I'm saying O. Okav. Now --17 17 A. They've asked me questions about it, we've it's certain. 18 18 debated about it, but I am the author. I chose to Q. Right. So why -- if you're certain and you 19 19 say it's certain that a transformation happens and put in more likely than not, even though, in fact, 20 that's damage, why then in your report do you it weakens my argument. But I have no problem using 21 21 indicate the transformation more likely than not it, because it's consistent with my argument. 22 22 leads to physical damage? Q. Okay. Paragraph 51 talks about how the 23 23 virus is spread -- the virus is primarily spread A. Because --24 24 MR. ALVENDIA: Objection -- wait. through the air. And the first sentence reads, "The 25 25 Wait, wait. airborne dissemination of the virus was" -- yeah,

November 10, 2020 name? A. Buonanno. before you did this research? A. No, sir. data, right? A. For this study, yes. 11 Q. Why? 12 13 14 15 16 17 18 19 paragraph 50. 20 21 a little bit, Kevin? 22 BY MR. MILLER: 23 24 Buonanno. 25 A. Yes, correct.

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by -- what's this gentleman's name, or person's

- Q. Buonanno. Were you familiar with Buonanno
- O. Okav. If you look on -- at your references, you utilize Buonanno for this scientific
- A. Because I believe that the methodology that they used was sound, and I believed -- and I understood what their conclusions were and that therefore it's -- and -- and it provides data that supports my hypothesis that this virus does spread.
- Q. Okay. In paragraph 50 -- you don't have a reference for the analysis that you use in

MR. MILLER: Could you scroll back up

Q. Paragraph 51, you've got some data from

A. Yes.

# Q. Do you have any other peer-reviewed article that would say that?

A. Yes. They -- they -- let's back up for a second. Can we go up one?

Paragraph 49 talks about the number of patrons entering the restaurant. It also describes what happens when somebody sneezes. Now, if you actually need a reference that says there are as many as two billion viral particles that can fall on an individual after a sneeze, I can get that.

But that is -- it is well understood that when somebody sneezes, they don't just sneeze out one or two. They sneeze out hundreds of millions and billions. That --

# Q. Let's go back --

A. That in and of -- that -- that -- that in and of itself, if one understands what viruses do, make the environment -- let -- let me just use my own words here -- change the environment from one of -- an environment from one of safety to one of infectivity and illness.

#### O. Let's --

A. -- (indiscernible) virus in the air leads to that conclusion.

Page 166

O. Paragraph 50, you don't have any reference for the number of statements where you talk about the physical change in the restaurant condition.

A. So let me say a couple of things about that. The ref -- no. First of all, I agree. I have got no reference number, no question about it.

#### Q. Okay.

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A. Okay. However, a predicate for the first sentence is the previous discussion, level of environment -- "level of environmental exposure, air and surface rose to dangerous levels." That's previous discussion.

"The restaurant's environment was transformed into a harmful condition as the virus physically transformed the air and the restaurant contents from one of safety to one of infectivity and illness."

To me, at this point, that's self-evident.

Q. Self-evident, so -- so do you have any other peer-reviewed article that supports the statement that "A restaurant was transformed into deleterious condition as the virus physically transformed the air and the restaurant's contents from one of safety to one of infectious and illness"?

Page 168

Page 167

- Q. Let's stay right there. This next sentence in paragraph 50, [As read]: "This transformation changes the structure of the surface of the restaurant contents by a process predicated by physical law."
  - A. Right.
  - Q. You don't have any citations there either.
- A. Oh, that's -- actually that's coming. I just pre-stage what I was going to get to, but that's coming.

### Q. Oh, it's coming from whom?

- A. I mean -- I beg your pardon. That appears -- the justification appears later in the report. That's what I mean.
- Q. Oh, your -- your physics -- your physics versus -- your physics and chemistry analysis?
  - A. Organic chemistry, yes.
- Q. Okay. But you don't have any peer-reviewed article sources that would support that statement?
- A. Right. "By a process predicted by physical law"? Is that the statement you're concerned about?
- Q. The transformation that you're describing. That -- this trans -- the -- that "the virus physically transformed the air and the restaurant contents."

Page 169

The virus physically transformed the restaurant contents?

A. Yes.

- Q. Point me in a direction of one person, other than yourself, that says the virus physically transforms the restaurant's contents.
- A. Well, I will point to you several places. I will point you to text in virology. I will point you to text in organic chemistry.
- Q. Okay. That are peer-reviewed -- why don't you cite those in your -- why don't you cite those in your report?
- A. I mean, I'm happy to, but those are very basic chemistry books, very basic virology books. I mean, I'm happy to do that, but it's -- perhaps not self-evident to everybody, but this is what physical law is.
- Q. Okay. The last sentence, "The change in the structure is the damage, so the transformation more likely than not leads to physical damage."

Point to me one individual, one article, one report, one peer-reviewed article, even a paper, where somebody says that the change in the structure is the damage, so the transformation more likely than not leads to physical damage --

Page 171

- A. I have no idea.
- Q. Okay. You haven't seen that anywhere else?
- A. I don't know if I haven't seen it anywhere else, but when I'm asked what damage is, this is my definition.

the way you do in the contents of this report?

O. Okav.

A. Now -- now, over the years maybe I have absorbed an understanding of damage from different people, that's likely. That is -- forgive me -- more likely than not, but this is my definition of damage.

- Q. Okay. Paragraph 53 talks about the risk of infection in ventilated areas.
  - A. Yes.
- Q. Can you describe to me the ventilation system that's in Oceana?
  - A. I cannot.
- Q. You have no idea what type of HVAC system that they utilize when patrons are present?
  - A. I don't know what that would be.
- Q. Do you have any idea of how they aerate the building, whether they leave doors and windows open during this COVID situation that we've been going through?

Page 170

A. Well, first of all.

- Q. -- because COVID-19 is on a surface?
- A. They're my words and not anybody else's yords.
- Q. Okay.
- A. All right. So I defined -- clearly in this report I define damage as change in the structure. I defined it that way.
  - Q. Anyone -- does anyone else --
  - A. My definition --
  - Q. I'm sorry. Go ahead.
- A. My definition -- my definition, and I stand by that definition.
  - Q. Okay.
- A. Now, if -- now -- okay. So the change in the structure is produced by physical law, so therefore the transformation is more -- it's -- the transformation -- it's total logic.

The transformation leads to physical damage. The transformation is the physical damage. And the transformation occurs by physical law. And that physical law is based in the fundamental tenet of organic chemistry.

Q. Can you point me to one article, peer-reviewed or not, person, that defines damage

Page 172

- A. Well, having been in New Orleans, I hope they don't open doors and windows. But I would say beyond that, I don't know.
- Q. Okay. When determining whether or not the restaurant's environment is transformed, would the ventilation have any bearing upon your analysis as to whether or not there's property damage?
  - A. Ask me that question again, please.
- Q. Sure. In determining whether or not the inside of Oceana Grill has been transformed into a dangerous condition or a dangerous environment, would the type of ventilation system affect your opinion at all?
  - A. I think it would.
- Q. Okay. But you don't know what the ventilation system is at the restaurant?
- A. Right. And just to be clear, I'm not saying I would reverse my conclusion. I think it would be impacted by the kind of ventilation.
- Q. Okay. In paragraph 54 you talk about -you say in the second sentence, [As read]: "So the virus is spread by other means as well. And, for example, highly sensitive laser light scattering observations have revealed that loud speech can emit thousands of oral fluids."

Page 173

A. Droplets.

# Q. "Oral fluid droplets per second." And then you cite to number 25. What is that source?

A. I have to see. Well, it's an article. I can't pronounce the last name. Stadnytskyi, I would say. The title, "The airborne lifetime of small speech droplets and their potential importance in SARS-CoV-2 transmission."

Does that answer you question?

# Q. And why -- why did you utilize that source?

A. Well, I utilized that source because I thought their methodology was sound, and it provided yet one more vector of spread of the virus.

Q. Okay. Let's look at paragraph 55. This is your analysis with respect to the virus and surfaces. And I'm just going to go down to the surface portion of this paragraph where it says, "The virus also alters the surface of solid objects," you list a number, "through the process of molecular cohesion and adhesion."

#### Tell me what that means.

A. Of course. I'll try to say this in English. Viruses, being very small, are affected by movements of electrons, and movements of electrons determine what -- whether a molecule cleaves to Page 175

Q. Now, the scientific makeup of -- would the scientific makeup of the wax that you reference in here have any bearing on cohesion and adhesion?

A. Well, my understanding, wax is essentially paraffin, and there are different types of waxes. They are by and large all alkane products, all organic products, and organic compounds interact with each other.

Q. Do you know what type of wax Oceana Grill uses on its furniture?

A. I do not.

Q. Okay. When we were talking about this study in paragraph 51 -- I'm sorry. Go back to paragraph 51 -- and you relied upon -- again, I will butcher that name if I tried.

A. Let's just say Buonanno.

Q. Buonanno. And I asked you why, because you said the data would seem to be reliable, for lack of a better --

A. The methodology was reliable.

Q. Methodology was reliable. And he's talking about -- so that study was done in China, correct?

A. Yes.

Q. Okay. And do you think it's fair to compare a restaurant or building in China to a

Page 174

something or does not.

Should I go on?

# Q. Yes, I'm listening.

A. All right. All right. All right.

Therefore, the virus by it -- by its very size is maneuvered into positions in the surface, and those maneuvers lead to it being attached to other molecules in the surface. In this particular example, I'm talking about wood wax and virus -- not amalgam, but the wood wax virus environment.

### Q. Does the --

A. I should say -- excuse me. I'm sorry.

The use of the word "insinuation." I think one of your experts objected to that. I certainly don't mean to give the virus any intent. I mean it just as I described it. It is maneuvered into this position by forces it doesn't control, physical forces it does not control.

Q. So let's get to that. The next paragraph said, "This insinuation of virus into and through the surface of solid objects alters the surface to a wood wax virus hybrid that cannot be disassembled, separating the virus from the original source. This is the physical damage to the surface."

A. Okay. Yes.

Page 176

#### restaurant in the United States?

A. Well, I would say this to you -- and just so we're very clear, because we didn't talk about this when the Buonanno study --

Their study results were simulations. You can't do an experiment like this on people. These are simulations, and these are simulations based on modeling. So do I think the Chinese modeling is relevant to U.S. restaurant modeling? Yes, I think so.

Q. Earlier, we talked a little bit about, you know, the RNA that's contained in the CoV-2 virus.

Does RNA on a surface necessarily mean that there's a live virus that can infect someone?

A. Probably not. To infect an individual, the infective particle has to be moved from the surface -- and I assume we're talking about an inanimate surface here.

#### Q. Yes.

A. -- to basal epithelial; and if we just have raw RNA, raw RNA doesn't get into -- I mean, at least the state of the art virology tells us that raw RNA doesn't get into the cell and cause infection.

Q. Okay. Can an individual contract COVID-19

Page 177 Page 179 1 from CoV-2 viral fragments? A. In my analysis --2 A. CoV-2 viral fragments, the best I can tell Q. In your analysis, it can't -- what was the 3 you, it depends on what type of fragment we're word you used? I'm sorry. I said killed; you talking about. called it? 5 If it's only the enveloping coat, if that's A. Denatured, I hope. 6 being part of the enveloping coat and no RNA, then Q. Denatured? 7 no, they would not get -- they would not get A. Right. 8 Q. So your opinion is that a tabletop cannot 9 be denatured by disinfectants as the CDC says it can Q. And so let me ask this. Does the existence 10 10 of CoV-2 RNA on the surface necessarily mean that 11 11 there's a live virus that can infect someone? A. What I am saying is this -- close. 12 12 A. That's a question I don't know how to What I'm saying is this. The tabletop is 13 13 answer. I mean, the presence -- certainly, as you so irregular that the virus can find itself in nooks 14 14 pointed out, the raw RNA is not infected. I already and crannies in that table surface, that the 15 said that. 15 cleaning and the disinfectant can't reach. 16 Now, does the presence of raw RNA not 16 Q. Okay. So when the CDC says the virus can 17 indicate that if there's raw RNA that, in fact, 17 be killed by using appropriate disinfectant on 18 there are -- there might be a viable virus nearby? 18 surfaces, they're wrong? 19 Perhaps it does. Viable viruses, I should say. 19 A. It's making assumptions that are not true. 2.0 20 Perhaps it does. O. Okav. 21 21 Q. Okay. Let's go to 58. And the first two A. So it's misleading. However, it is simply 22 22 the best that they can do. I'm not here to sentences of 58 talk about human infection, and 23 23 criticize the CDC. They have a tough job; they do we'll skip over that. 24 24 The third sentence, it says, [As read]: the best they can. But unfortunately, the size of 25 25 "When the SARS-CoV-2 virus is driven down by gravity the particles mitigates against them. Page 178 Page 180 1 and micro air currents to a bar surface, the S1S2 Q. All right. Let's go to -- let's go to 2 2 projections adhere to the surface molecules of the paragraph 65. All right. 3 3 What does it take to restore the unhealthy bar, attempting the chemistry to infect the bar. 4 However, since infection fails, since there is no environment? And you rely upon the van Doremalen? 5 membrane for the virus to break through, the virus A. Yes, sir. 6 simply stays adhered to the bar surface." Q. How do you pronounce that? 7 Do you see that? A. Yes, sir. Just like you did, van 8 8 A. Yes. 9 Q. And that is true through and unless the Q. Why did you rely upon this van Doremalen 10 10 virus is killed on the surface, correct? 11 11 A. Okay. If we are assuming that we're A. I believe that the van Doremalen report 12 working with an absolutely flat and smooth surface, 12 used very good methodology. 13 then you're right. In that circumstance, there's no 13 Q. Okay. And the methodology used in -- all 14 place for the virus -- for the virus to go, except 14 right. Well, let's just -- let's talk about -- you 15 to stay on the surface, and at that point it can be 15 say van Doremalen evaluated the stability of 16 denatured. 16 SARS-COVID-2 and SARS-COVID-1 in aerosols and on 17 You obviously don't kill it because it's 17 various surfaces, estimating their decay rates. The 18 alive, right? We denature it. Okay. That is not 18 research consisted of ten experimental conditions 19 the reality of surfaces, which is what my report 19 involving two viruses, SARS-CoV-2 and SARS-CoV-1, in goes into it. In fact -- I'm sorry, go ahead. I'm 20 five environmental conditions: aerosol, plastic, 21 21 stainless steel, copper, and cardboard. In fact, 22 Q. So this -- my scenario doesn't apply to a 22 the SARS-CoV-2 remained viable in aerosols 23 restaurant table --23 throughout the duration of our experiment, three 24 A. Correct. 24 hours. And this reduction was similar to that 25 Q. -- in your analysis. 25 observed with SARS-COVID-1.

Page 181 Page 183 1 1 And you agreed with the methodology used people who are in the restaurant and infected and 2 in -- by van Doremalen -sneezing. 3 A. Yes. You're talking about billions of viruses 4 Q. -- to support these principles? here. So I think the concentration is perhaps not 5 5 so irrelevant, and I don't know what the temperature A. Yes. 6 6 at Oceana is. I don't know what the humidity is. I Q. And you thought that these principles are 7 7 applicable to your analysis with respect to Oceana assume that it's there to make the people 8 8 Grill? comfortable, right? 9 9 A. I thought they were applicable, yes. Were they that different than the 10 10 Q. Okay. In this report, one portion of environmental conditions here? I'm not sure. 11 11 methodology was that it was -- I'm sorry, in this Q. So your testimony is that lab conditions 12 12 study was that it was under lab conditions, correct? for this study you think are comparable to -- let's 13 13 A. Yes. just pick a month. Let's pick the height of the 14 14 Q. All right. And so they utilized controlled pandemic. Let's say the summer, right? 15 temperatures? 15 Is there a chance that there's 40 percent 16 16 A. Yes. humidity in New Orleans during the summer months? 17 17 A. Well, I would say this. If you leave the Q. 23 degrees Celsius or 73.4 degrees 18 18 doors and windows open and it's 40 percent humidity, Fahrenheit, correct? 19 19 I'm coming. I'm going there. A. Yes. Yes, sir. 20 20 Q. They used controlled humidity, correct? My experience, the humidity is a lot 21 21 higher, but as you pointed out or reminded us, I am A. Yes. 22 22 not a specialist in HVAC. I don't know what the O. And I think the humidity was 40 percent, 23 23 humidity HVAC produces at Oceana. Maybe it produces right? 24 24 A. I don't remember. But, yes, they 40 percent. I just don't know. 25 25 controlled humidity. Q. Right. And the fact that you don't know Page 182 Page 184 1 Q. And they also used a distinct concentration would render your reliance upon this study under lab 2 2 conditions irrelevant to the conditions at Oceana? of the virus in the report to do the test in the 3 3 labs, right? MR. ALVENDIA: Objection to form. 4 4 He's already answered this question. A. Yes. 5 Q. And that concentration would be much higher But, Doctor, you can go ahead and answer it 6 than, say, if an individual sneezed, right? A. I actually don't know. I'd have to look. A. I would just say this. I mean, I do debate 8 O. Okav. But taking that prong out, or if you and continue to debate with you the notion about how 9 just indulge me for a second that concentration was much virus is there because I think it's easy to 10 10 in, fact, higher than what you would expect from underestimate virus in an environment where there 11 11 either an individual speaking or a sneeze -are multiple people spewing it out. 12 12 Having said that, having said that, the A. Right. 13 13 conclusion that it remained viable in aerosols I Q. -- there were lab conditions, controlled 14 14 temperatures at 73.4 degrees Farenheit, controlled think is relevant. 15 15 humidity at 40 percent, and then a concentration of Now, we can debate minutiae in terms of how 16 16 long it's viable in aerosols, in Oceana versus this the virus. 17 17 None of those factors are present ever at experimental condition. But the point is, it's 18 18 Oceana Grill, correct? viable in aerosols. 19 19 A. Actually, I can't say that. BY MR. MILLER: 20 20 Q. You can't say --Q. Okay. Let's look at paragraph 67. 21 21 A. The one thing I could say is that while I A. Okay. 22 22 agree that the concentration of virus might be high, Q. There you're relied upon Chin report, 23 23 I'm not so sure I agree with you that it would be right? 24 24 higher than the concentration seen in Oceana when A. Yes. 25 25 you've got -- let's not say hundreds -- say tens of Q. And similar to the Doremalen report, the

Page 185 Page 187 1 1 Chin report was done under lab conditions, correct? there's no UV light in the dark; and I know that it A. Yeah. I do have to -- I do have a gets dark in New Orleans. I don't have to go there 3 3 criticism of the Chin report in that it is very to know that. And I, therefore, know that the 4 brief. If I remember right, it was a difference is how much gets in during the day. 5 5 correspondence. It wasn't a full paper. I'm trying And I'm now conceding that there's more UV 6 6 to go by memory here. It wasn't a full paper. So light getting in during the day, but I really don't 7 it's hard to know exactly what was going on in all know how much more is getting in. 8 8 the detail we would like. Q. Okay. Do you know what the current state 9 9 Q. But it too had controlled temperatures and of literature with respect to the survivability of 10 10 controlled humidity, right? CoV-2 in non-lab conditions is? 11 11 A. Yes. What's the current scientific hypothesis 12 12 about the survivability of CoV-2 today in non-lab Q. And it was also done in lab conditions, 13 13 correct? conditions? 14 14 A. Yes. A. Yeah. So my focus -- thank you for 15 15 Q. Do you know what the concentration of the clarifying that. 16 viruses that was used in the Chin report --16 My focus has been lab conditions, so I 17 A. I don't recall. 17 don't know that -- I'm trying to understand what the 18 Q. Okay. Let's go to paragraph 70. This, 18 value would be of studies that don't look at it in 19 19 again, is another source you rely upon, the Riddle unstandardized condition. I don't know what -- what 20 20 does that really mean? report? 21 21 A. Yes, sir. Q. Well, perhaps because Oceana Grill is not 22 22 O. And similar to others, this was under lab in a lab, there may be some utility to them to know 23 23 what the survivability is of CoV-2 outside of a lab conditions, right? 24 2.4 A. Yes. This is quite a bit more detailed condition. 25 25 than Chin was. A. Fair enough. Fair enough. However, when Page 186 Page 188 1 Q. And, in fact, in Riddle, they actually you're outside of -- when you're outside of 2 2 maintained the virus in the dark, right? standardized conditions, lab or not, there's 3 3 A. Yes, to avoid the effects of UV light. variability day to day; UV light changes day to day; 4 4 temperature changes day to day; and so viability 5 Q. Which would do what? changes day to day. 6 6 A. UV light is high frequency radiation. It So it's hard to know what the -- what the 7 real measurement actually means since the conditions can ionize, which means it can break, break 8 continue to change. Put another way, the mean may electrons off from atoms, destabilize molecules, and 9 be different, but the variability confuses the denature the virus. That was the short version. 10 10 issue. Q. And so it's fair to say that Oceana Grill 11 11 has much more UV light with its doors, windows, and O. Let's look at 73. 12 12 openings than keeping a virus in the dark in this A. Okay. Can I suggest we take a break for a 13 13 few minutes before we go on? Riddle report, correct? 14 14 MR. MILLER: Well, assuming your --A. I would say there was more UV light. 15 MR. ALVENDIA: Yeah, let's do that. 15 However, if it's much more, I don't know. But there 16 16 was more UV light. But, again, let's do a time check here 17 17 Allen. We're now at 5:30. We've been Now, of course, UV light is only present 18 18 going for five-and-a-half hours. What are during the day, not present at night. It's present 19 19 we looking at here? on cloudy days, but not present at night. So there 20 20 MR. MILLER: I mean, hopefully close is a difference; but how much the difference is, I 21 21 to 6:00. If it's past 6:00, it won't be don't know. 22 much past 6:00. 22 Q. And part of the reason is because you don't 23 MR. ALVENDIA: All right. Let's take 23 know how much UV light comes into Oceana Grill 24 24 a five-minute break. because vou've never been there? 25 25 (Recess taken.) A. No. Part of the reason is that I know

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- BY MR. MILLER:
  - Q. Doctor, take a look at paragraph 73.
  - A. Okay.

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Q. And it is a discussion of Oceana Grill's remediation efforts, and the last sentence basically says -- oh, let's go with the first sentence.

[As read]: "Oceana has attempted to restore the pre-COVID-19 environment by maneuvers, e.g., arising out of -- arising out the property, generally -- airing out the property, generally cleaning the surfaces with bleach-based cleaners, and closing off rooms where individuals who reported COVID-19 were located for 24 hours.

"While this effort is understandable, it's inadequate because, one, the effort was undone by the continued arrival of SARS-COVID-2 positive patrons and employees; and two, the recent research of Riddle" -- or "Riddle" -- I don't know --"reveals the virus is viable for up to 28 days and perhaps longer, invalidating the process of cordoning off a room for 24 hours."

The 28-day number that you cite there is the -- where the virus was kept in a dark area -darkness in a laboratory under controlled temperatures and humidity, correct?

Page 191

kills the virus. And I can show you exactly from their website where they say it kill the virus.

They're wrong when they say that?

MR. ALVENDIA: Objection. You're mischaracterizing his testimony. He's not saying -- he's not saying if the cleaner doesn't touch the virus itself, it kills it. What he's saying is if it can't touch it, it doesn't kill it. He said that about three times now. He's not saying they're wrong.

MR. MILLER: We could be here until 7:00 o'clock, if you want, because I'm going to ask the questions. So let me ask the questions. You can make your objection and then let me ask the question.

MR. ALVENDIA: My objection is you're mischaracterizing his testimony. Please

Doctor, you can answer, if you understand his question.

A. In order for the virus to be denatured, not killed, but denatured, it has to be touched by the cleaning agent. If it can be touched by the cleaning agent, then it will be denatured.

Page 190

A. Yes.

# Q. So that condition doesn't normally exist at Oceana Grill, does it?

A. That is true. It does not ordinarily exist.

# Q. Okay. Are you familiar with --

A. I'm sorry. I do need to also tell you that -- so I've given you two reasons that I think it was inadequate.

But the third reason that I didn't say here but I want to say on the record so that you understand and you can examine me on it, is that, in fact, the bleach-based cleaners simply can't reach the virus when it is embedded in the surface. That's a third reason bleach can't reach the virus.

Q. So again, the recommendations that are given by the Centers for Disease Control are simply not accurate?

A. They -- I don't want to say -- I wouldn't say it that way. Again, I'm not going (indiscernible) their effort. They do the very best they can. But the size of virus mitigates against their belief.

Q. So they are wrong when they say that if you take CDC-approved cleaners and wipe the surface, it Page 192

I don't disagree with them. I'm simply saying that many virus are embedded so deeply that the cleaning agents can't reach them. BY MR. MILLER:

O. And let's talk about Oceana Grill, because that's really the only reason we're here, and the surfaces that are in a restaurant.

In your conclusion, your opinion, is that the guidance given by the Centers For Disease Control, that restaurants can denature the virus to the point where it is safe for patrons to eat in a restaurant, is inaccurate because the surfaces in Oceana Grill will not allow you to denature the virus?

A. I am simply saying that the CDC is correct in that the sequence of cleaning and disinfectants can denature viruses when they reach them, and they reach many viruses.

But they do not reach all viruses because these viruses are embedded, as I've said; and so those viruses continue to be a threat. That's what I'm saving.

Q. Let's talk about Oceana Grill, because you have authored a report that says, "The viruses have reached the surface at Oceana Grill, and it creates

Page 193 Page 195 1 1 a dangerous environment." publicly? 2 Am I incorrect about that? 2 Q. Yes. 3 3 A. I'm sorry. You said it was in press. That A. Correct. 4 Q. Okay. The Center For Disease Control has means it hasn't appeared publicly. 5 5 said that establishments like Oceana Grill can Q. Okay. I'll tell you what, what about 6 6 create an environment safe for its patrons and, in the -- what I'm going to refer to is the Goldman 7 their words, kill the virus -- but we're going to report from the Lancet, which is one of your 8 8 use denature -- if you use the cleaning agents that reviewed articles. 9 9 they suggest to wipe down the surfaces. Are you familiar with Goldman's report in 10 10 And because you're here as an expert and the Lancet? 11 authored a report with respect to Oceana Grill, my 11 A. Is it one of my reviewed articles? 12 12 question to you is, the surfaces that are at Oceana Q. No, it's not one of your reviewed -- well, 13 13 Grill, the viruses that you say are at Oceana Grill, Lancet is one of your reviewed articles, yes? 14 14 A. The Lancet is a journal I did peer reviews your opinion is they cannot be denatured? 15 MR. ALVENDIA: His opinion is more 15 for. 16 16 likely than not. Is that what you're Q. Okay. This is a report by Goldman in the 17 17 asking, Allen? Once again, it's important Lancet, which is the publication you did peer 18 18 to use -reviews for. 19 19 MR. MILLER: I'm asking him what his A. Just to be clear, the Lancet publishes 20 20 opinion is. I don't want to ask -- I'm not thousands of articles. 21 21 interested in what the lawyers' opinions O. Sure. 22 22 are. I just want to know what the A. This sounds like an article I have not 23 23 witness's opinion is. read. 2.4 24 MR. ALVENDIA: I'm asking you what Q. Okay. 25 25 level are you asking him. Are you asking A. Okay. All right. Page 194 Page 196 1 1 more likely than not? beyond a reasonable Q. Tell me what fomites are. 2 2 doubt? It's an improper question. A. Fomites are the vehicles that spread 3 3 BY MR. MILLER: infection. 4 4 Q. You can answer my question, assuming you Q. All right. And how do fomites apply in 5 understand it, Doctor. this case? 6 A. That level of cleaning is insufficient. A. Well, I think -- the way I think about 7 It's more likely than not insufficient. fomites is that they are the droplets that spread 8 8 Q. To denature the virus? the virus. A. Correct. MR. MILLER: Okay. Can you pull up 10 Q. Okay. Despite the recommendations of the 10 the Lancet report concerning exaggerated 11 11 **Centers for Disease Control?** risk of transmission of COVID by fomites? 12 A. Centers for Disease Control does the very 12 A. I don't know. If it's not --13 best they can using the cleaning agents. But 13 BY MR. MILLER: 14 unfortunately, as I said before, the virus mitigates 14 Q. Not you. I'm talking -- I'm talking to my 15 against that. The virus's size mitigates against 15 colleague. 16 that. 16 A. I apologize. Okay. 17 Q. Are you familiar with a Clinical 17 Q. This is an article out of the Lancet that, 18 Microbiology in Infection publication? 18 which again, is the publication where you've done 19 A. I'm not sure what -- you're talking about a 19 peer-review articles. 20 book? a journal? an article? 20 Have you ever seen this article before? 21 I'm not sure what we're talking about. 21 A. I have not. 22 Q. CMI, it's a -- articles in press. 22 Q. Okay. In your work on this case -- I know 23 What about a gentleman by the name of 23 we talked about a number of articles and a number of 24 **Ben-Shmuel?** 24 things that you did use, all of which were, I would 25 A. No. The article, does that appear 25 say, in the spring of 2020, correct? Early on in

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the COVID analysis.

A. I'm sorry. What was your question again? I was just trying to read this.

Q. Yeah. Sure. Take your time. Why don't you go ahead and --

A. Can we go to the next page?

MR. ALVENDIA: Yeah, I was going to say -- I was going to say, Allen, if you're going to question him on the articles --

MR. MILLER: Yeah, I'll let him look at it first.

MR. ALVENDIA: Let him look at the whole article, please.

A. Can we go back to the previous page, please? Okay. Thank you. BY MR. MILLER:

# Q. Do you have any reason to disagree with this article?

A. I have a reason to question its results.

Q. Why?

A. First of all, I don't know if its results are correct or not. By that I mean I don't know if they reflect the true state of nature or not because there's not very much information about methodology here. This is really a brief. This is not a

Page 199

events we just talked about, individual comes in, droplets go onto the table, immediately after droplets are on the table -- we're talking about just inanimate surfaces right now, not in the air or person to person.

The virus is on the inanimate surface, person then has to touch the inanimate surface, likely with a hand, then that hand has -- prior being -- the virus being compromised, go, then, to that person's either face or eyes. The concentration on your hand has to be sufficient enough to get through your individual membranes and into your cell to then infect the cells and you then become infected.

Because of that, this gentleman says that -- let's see. "Although periodic" -- "I believe that fomites that have not been in contact with an infected carrier for many hours do not pose a measurable risk of transmission." And in this case, he did it in non-hospital settings.

Do you agree or disagree with that?

A. I would say, based on what I have read here and what I know, I would disagree. And what is missing here is the methodology he used to come to his conclusion.

Page 198

full-length manuscript. This is really a very brief report.

If this is pre-staging a full-length manuscript, then I'd be happy to read that. But I don't get much of anything from this.

Q. Okay. Go to the second page. Let me -okay.

So in this, this gentleman opines -- and this is Mr. Goldman -- that in his opinion the chance of transmission through inanimate surfaces is very small, and only in instances where an infected person coughs or sneezes on the surfaces, and someone else touches that surface soon after the cough or sneeze.

And then I will go a step further and say, then that individual touches their nose with a sufficient concentration of the virus on their hand that can then penetrate your personal membranes into your cells and then you catch -- then you would be infected.

So although he doesn't go that far, I'll take it two steps further, and you would probably agree with me on those last two steps.

- A. The pathophysiology is sound.
- Q. Okay. So as a result of the sequence of

Page 200

Q. And let me ask you one question, then, as a follow up.

In all of your studies, in all of your readings, and in preparation of your report, can you identify one instance anywhere in the world where an individual has been confirmed to have contracted COVID-19 from an inanimate object?

MR. ALVENDIA: Once again, using the standard of more likely than not?

MR. MILLER: No. I'm not asking about an opinion. I'm asking about a fact. I'm asking for him to identify to me one fact, that being a confirmed case of an individual somewhere on this globe that contracted COVID-19 from a inanimate object.

A. My answer to you is that I have not done anywhere near a sufficient survey to be able to answer that question with a surety.

To answer you directly, no. But I haven't done a study of it. The fact that I can't think of an example doesn't mean an example doesn't exist. It just means that I haven't contacted the right or wrong people, depending on your point of view. BY MR. MILLER:

Page 201 Page 203 1 1 O. Okav. of COVID-19 by Fomites" as Exhibit 3, I 2 A. I don't have an informative answer. think. 3 (Exhibit No. 3 was identified.) Q. So the answer is, no, you do not have one 4 fact, meaning one person worldwide that has ever THE WITNESS: If you're waiting for 5 confirmed contracting COVID-19 from an inanimate me, I've read this. BY MR. MILLER: 7 A. My answer is, no, I do not have an O. Okav. You've read this. 8 informative answer. And do you disagree with this article? 9 Q. Okay. Despite not having that fact, not A. I will say that there is a -- yes. And I 10 10 having researched whether or not a person has would say there's a substantial body of literature 11 contracted COVID-19 from an inanimate object, not 11 that -- that is much more detailed than this that 12 12 person to person, but on the surface; despite not contradicts the finding. 13 13 having surveyed, not having looked for articles, you O. Okav. 14 have opined that the restaurant, because of, in your 14 A. I'm sorry. And maybe -- this is in Italy. 15 assertion, CoV-2 is present on the surface, is a 15 And actually, they refer to studies 4 and 5, which 16 16 are Colaneri -- actually, they're both by Colaneri, dangerous -- creates a dangerous condition? 17 17 and I haven't read those. So I would need to read MR. ALVENDIA: Objection to form. 18 18 those to fully opine on this. MR. MILLER: That was really poorly 19 19 But I would say that I disagree with this, 20 20 MR. ALVENDIA: So objection to form. based on its face, what they presented. 21 21 MR. MILLER: But I'll say it again. Q. All right. Their findings -- one of their 22 22 BY MR. MILLER: finding -- and I'll just read a sentence. 23 23 Q. Doctor, did you understand my question? It says, "Our findings suggest that 24 24 A. The first 40 percent. environmental contamination leading to SARS-CoV-2 25 25 Q. Fair enough. All right. transmission is unlikely to occur in real life Page 202 Page 204 1 You have just told me that you have not conditions, provided that standard cleaning 2 2 taken any steps to determine whether or not an procedures and precautions are enforced." 3 3 individual has ever contracted COVID-19 from an Do you see that? 4 A. I do. Now, I would use -- I'm sorry. Go inanimate object. Is that fair? A. That's fair. 6 Q. That's also the same guidance that's given Q. And so you do not have an informed answer by the Center for Disease Control, isn't it? That about whether or not there is one person worldwide standard cleaning procedures or precautions would that has, in fact, contracted COVID-19 from an create an environment that prevents contamination inanimate object. 10 10 from SARS-CoV-2 transmission? MR. ALVENDIA: Objection to form. 11 11 A. This is Italy. I don't know that Italy You can answer. 12 12 A. That's fair. follows CDC, I just don't. 13 13 Q. I'm not asking if they follow CDC. I'm BY MR. MILLER: 14 14 asking whether or not this finding is consistent Q. Okay. Established -- now that we've 15 15 with the guidance that the CDC gives us. established that, that you have not done that 16 16 A. Standard cleaning procedures, I can't tell research, you still in this case opine that the 17 17 from this. inanimate objects in Oceana Grill create a dangerous 18 18 Q. Okay. Does the CDC not recommend to us environment, correct? 19 19 that in order to denature the virus -- they use the A. I do, yes. 20 20 word "kill" -- standard cleaning is what a Q. Okay. That's it. I think we labored 21 21 restaurant like Oceana would need to do? through that series of questions. 22 A. Actually, it's standard cleaning and 22 MR. MILLER: Can you pull up the 23 23 disinfectants, right? Mondelli Lancet article, September 29th? 24 24 Q. Standard cleaning and disinfectants, yes, For the court reporter, I'm going to 25 25 sir. make the "Exaggerated Risk of Transmission

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Page 205 1 A. I think there's a difference here. I think 2 there may be a difference in what they do and what 3 Oceana did. 4 MR. MILLER: Okay. I'll mark this as 5 Exhibit 4. 6 (Exhibit No. 4 was identified.) 7 BY MR. MILLER: 8 Q. All right. I'm going to try to be quick 9 with these questions, Doctor, if you can. 10 Ionic bonding, could you describe that to 11 me? 12 A. Ionic bonding is a process by which two 13 atoms interact with one atom, taking the electron of 14 another. 15

Q. Okay. And covalent bonding?

A. Covalent bonding is a process by which two atoms share electrons.

Q. Can you have molecular adhesion covalent bonding with a particle ion character?

A. I will say, yes, you can. Nature is not black and white, even though we make -- we make definitions of black and white.

Covalent bonds are not always exclusively covalent. The electron does not always share the ionic bonding. The theft of the electron is not

Page 207

Page 208

- A. Van der Waals forces and London forces.
- O. Are different from ionic and covalent bonds, right?
  - A. Correct. Yes.

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# Q. Okay. So what is a reaction mechanism?

A. I don't know what you mean. I mean, I remember old definitions where reaction mechanisms were defined at SN1 mechanisms and SN2 mechanisms based on the availability of -- based on the availability of electrons, whether there are lone pairs of electrons available or not. That's my understanding of reaction mechanism. That may be a little dated.

Q. Now, let's go to -- all right. Let's go back to paragraph 55 in the report. So -- and we talked about this paragraph for quite some time, but I'm just going to focus on one particular phrase where you indicate that -- and what sentence is it

It's really the second-to-last sentence. "This insinuation of the virus into and through the surface of solid objects alters the surface to a wood wax surface hybrid that cannot be disassembled" --

A. Right.

Page 206

always complete, so there is overlap.

Q. Can you give me an example, something that I could recognize?

A. I don't know. Let me think. Hydrogen atoms are -- maybe something like carbon tetrachloride.

- Q. And how would that manifest itself in the real world, carbon tetrachloride? What --
  - A. Oh, what is it used for?
  - Q. Yeah.

A. The chemistry reagent I want to say is also a cleaning solution. I'm not really sure about that. It's been a long time.

Q. Could you spell it?

A. Oh, of course. Carbon tetra, t-e-t-r-a, chloride, c-h-l-o-r-i-d-e.

- Q. Okay. And that is an example of molecular adhesion covalent bonding with a partial ionic character?
  - A. I believe so, yes.
- Q. Okay. Now, you -- you mentioned this in your report, and I'm not trying to get into all the science of it, but I just want to ask you to make sure I have clarity. Van der W-a-a-l-s forces and London forces are different?

Q. -- "separating the virus from the original" -- "separating the virus from the original

surface." So the hybrid that cannot be disassembled, how then -- explain to me how then if it -- if the virus hybrid can't be disassembled, how then can an

A. Sure. So really I should say cannot be disassembled easily. I should really say that. I mean, I don't want to say it could never be disassembled. It cannot be disassembled easily.

O. Okav. So...

individual contract it?

A. So, but the answer -- go ahead. Go ahead, please.

Q. No, you talk. I -- go ahead.

A. So this is the insinuation sentence. So let me -- so if the virus inserts itself into this wood and wax and it bonds, now these bonds are typically weak bonds. They are weak covalent bonds. There maybe be some Van der Waals and London forces, but these are essentially weak bonds. Okay?

Can they be dislodged? Yes, of course they can be dislodged. They can be dislodged by wind, they can be dislodged by -- remember the surface is not smooth. So if the surface is not smooth, if

Page 209

it's rough, that rough surface can change. And that change in the rough surface by somebody -- by molecules running across it can expose viruses that otherwise were not available to be exposed. So these changes in the surface top would allow it to be released in a microcurrent of air.

- Q. So -- and look, it's getting late. I'm trying to wrap up. My brain is probably not working as good as it was six hours ago, but if you're telling me that the virus hybrid cannot easily be disassembled, in order for an individual to contract it from the inanimate object, they have to disassemble it, correct?
  - A. Correct. Yes.

- Q. All right. So just hear -- hear me out for a second.
  - A. Sure. Of course.
- Q. If the individual's hand can disassemble the -- disassemble the virus hybrid that ultimately would through a series of events cause that person to contract COVID-19 from touching the surface --
  - A. Yes.
- Q. -- in the same veracity that the individual could disassemble the virus and contract COVID-19, the cleaning of the surface in the same percentage

Page 211

- Q. Got it. But it also -- but it would mean that the same viruses that are embedded could not contract the person's hand, because if you couldn't get them with -- with cleaning, how are you going to get them by just sitting at a table?
- A. Oh, okay. So I -- I -- I understand your point, and I would say this. There are some viruses that will not -- will never be released from the table, right? They are too deeply embedded in the table, they are too tightly bonded with the table. So they're not going to infect anybody. They're not going to be cleaned out. They're not going to infect anybody.

So there are three conditions here, three classes. Class 1, viruses get on the table. They can be cleaned. No question. They're not going to infect anybody. They can be denatured. All right.

Class 3, viruses that are so far deep they don't get cleaned, but they don't get exposed to anybody.

It's the Class 2 that is of concern. Too deep to be cleaned out, but not really bonded too tight and ultimately they do get out.

Q. Okay.

A. Okay. Three mutually exclusive conditions.

Page 210

would allow for the virus to be --

- A. Denatured.
- Q. -- your word -- denatured.
- A. Denatured.
- O. Correct?
- A. I understand. Let me explain this to you, okay.

#### O. Please.

A. When the -- viruses settle on the surface many -- viruses settling on the surface of a table is essentially like -- for -- to help you with scale, is like dropping a million marbles into the Grand Canyon. Many of them stay on the surface, many of them can be cleaned, many of them fall deeper and cleaning can't get to them. Okay?

So the viruses that remain behind are the viruses that either are loose but can't get cleaned because they are so far embedded or they are covalently bonded. So the cleaning stops, the viruses that the cleaning could not get to can be rereleased, and the viruses that are bonded can be -- and they don't all have to be -- but they can be -- have that bond broken, because the bond is fairly loose. So cleaning will kill many viruses, but it doesn't mean the table is not infected.

Page 212

- Q. And in your work in this case or your work, period, have you, in any scenario, created those three conditions to see how the virus reacted with respect to the infection of individuals?
- A. If your question to me is, have I ever demonstrated that viruses in Class 2 infect people, I've never done that experiment. Nobody has ever done that experiment.
- Q. Okay. Let's talk about Class 2. The ones that can't be cleaned --
  - A. Right.
- Q. -- the ones that -- those are the -- for purposes of your analysis for physical loss of use resulting in damage, correct me if I'm wrong, are we -- is it fair to say that it's those Class 2 viruses that are at issue?
- A. I think those are the major ones. Ultimately Class 3 can become Class 2, but that's more complicated and way down the line. So we can focus on Class 2.
- Q. All right. Will a Class 2 virus corrode a bar or a table top?
  - A. What do you mean --
  - Q. Corrode it?
  - A. What do you mean by "corrode"?

Page 213 Page 215 1 1 O. I mean cause rust. damage be repaired or will there be property loss? 2 2 A. They don't have iron. No, they wouldn't Will the -- the item in question be unusable? 3 Q. And when you say "unusable," do you mean -cause rust. 4 Q. Okay. Would they cause any physical change let's just use a table for instance. 5 5 A. Right. to the naked eye? 6 6 A. To the naked eye? Q. Unusable meaning the table cannot be used 7 O. Yes. in the manner in which the manufacturer intended or 8 A. No. I don't see how that's relevant but, it cannot be used because it should not be used? 9 no, they're too small. MR. ALVENDIA: Let me -- let me just 10 10 O. Okav. Would any -- let's see. I want lodge an objection here. You're asking him 11 to -- I want to be clear so we can go through it 11 for legal conclusion, definition of 12 12 physical loss. I think he's already given quickly. 13 13 Would any measurable amount of CoV-2 or any that answer from his perspective, but 14 14 objection to form of the question. amount of concentrated CoV-2 evidence to the naked 15 eye any type of physical change on any inanimate 15 BY MR. MILLER: 16 16 Q. I only want you to tell me how you, in 17 17 A. Ask me that again, please. I think it drafting your report, are using the phrase "property 18 makes sense. I just need to hear it again. 18 loss"? 19 19 Q. Okay. Would any -- any level of A. And I'm using the phrase "property loss" as 20 20 concentration of CoV-2 placed on an inanimate object meaning the damaged entity really cannot be used 21 21 evidence a physical change to the naked eye? safely anymore. 22 22 A. I -- I guess I don't know how to answer Q. Okay. Give me about two minutes to look at 23 23 that. I mean, if somebody -- if you're asking me my notes and then I'll wrap up. 24 24 whether you could put down a heptillion viruses on a (Recess taken.) 25 25 surface --BY MR. MILLER: Page 214 Page 216 O. Fair enough. Fair enough. Fair enough. Q. All right. Doctor, just a few more 2 2 All right. Would any amount of CoV-2 that questions. Are you a member of the American College 3 would cause an individual to become ill, so the 3 of Epidemiology? 4 4 smallest amount of CoV-2 that would cause an A. No. I am not, no. 5 individual to become ill, on an inanimate object Q. Are you a member of the Public Health 6 6 evidence a physical change in the object to the Association -- American Public Health Association? 7 naked eye? A. I think I was at one point, but no. 8 8 A. I don't think so. And that, sir, is the O. All right. In formulating your 9 problem. That's the problem. You don't see it with conclusions, did you take any consideration into the 10 10 the naked eye, but nevertheless it causes illness. mandate in the city of New Orleans that required --11 11 Q. And I'm looking at the conclusions in your requires individuals in a restaurant like Oceana to 12 12 report, and you can go to B. And I'm -- I'm merely wear masks? 13 focused on the term "property loss." I know we've 13 A. No, I did not. 14 14 talked about damage. Q. But you are aware that since the initial --15 A. Yes. sir. 15 since the opening -- the reopening after the 16 Q. Do you have a specific definition of 16 shutdown in the city of New Orleans, that there was 17 property loss that you're using when you use that 17 a mandate by the mayor and her staff that 18 phrase throughout your report? 18 individuals inside of a building were required to 19 A. The -- the definition I'm using is the loss 19 wear masks? 20 of property due to damage. Does that help? 20 A. I under --21 There's damage and --21 MR. ALVENDIA: Objection -- wait. 22 Q. Which I know how -- I know how you define 22 Objection to form. That's 23 23 mischaracterizing it. If they're -- if 24 A. Right. And then the damages -- the 24 they're sitting at the table eating and 25 damages -- I mean, then it comes down to can the 25 drinking, they don't have to wear a mask,

Page 217	Page 219
Allen. We know this.  Given the full given the full restriction, given the full mandate.  MR. MILLER: Yes.  MR. ALVENDIA: It's not constantly wearing a mask inside a building.  MR. MILLER: All right. That's fair. MR. ALVENDIA: That's it. That was my objection.  MR. MILLER: Yeah.  BY MR. MILLER:  Q. So let me just ask you, Doctor. So I think you answered the question that the analysis that you came up with did not take into consideration whether or not individuals were wearing masks inside of Oceana restaurant?  A. Correct. Correct, because when you're in an environment where there's eating and drinking, if you're eating and drinking, you don't wear masks.  Q. Okay. Or did did your analysis take into consideration whether the staff members inside the restaurant were wearing masks?  A. It did not, no.  Q. Did your analysis take into consideration whether or not Oceana Grill used temperature checks	expert. You have to say for the purposes of this deposition, if you assumed this, would it would you have taken that into consideration. You're anyway. Look, it's getting late. Objection to form. MR. MILLER: All right.  BY MR. MILLER: Q. Dr. Moye, did your analysis consider whether or not temperature checks were made at the restaurant?  A. It did not. It didn't it really didn't need that.  MR. MILLER: Okay. All right. I think that is all I have. MR. ALVENDIA: All right. Thank you, Dr. Moye. (This proceeding was concluded at 6:25 p.m. on November 10, 2020.)
for individuals before they allowed them into the restaurant?  MR. ALVENDIA: Objection to form. MR. MILLER: What's wrong with that question?  MR. ALVENDIA: Assumes facts not in not in the record. Are you telling him that's what they do to every person MR. MILLER: No. I'm asking him if he took that into consideration.  MR. ALVENDIA: I know, but you're asking MR. MILLER: That was a fact MR. ALVENDIA: Are you saying that's what happens at Oceana? Because that's MR. MILLER: I don't know what Oceana does. That's not the question.  MR. ALVENDIA: Okay. Well, so you're MR. MILLER: My question is whether or not that fact MR. ALVENDIA: Right. MR. MILLER: whether or not MR. ALVENDIA: Then it's a it's objection to form, because you he's an	REPORTER'S CERTIFICATE  1, YOLANDA J. PENA, Certified Court Reporter in and for the State of Louisiana, Registered  Professional Reporter, and as the officer before whom this testimony was taken, do hereby certify that LEMUEL MOYE, M.D., PH.D., after having been duly swom by me upon authority of R.S. 37:2554, did testify as set forth in the foregoing 219 pages.  I further certify that said testimony was reported by me in the Stenotype reporting method, was prepared and transcribed by me or under my direction and supervision, and is a true and correct transcript to the best of my ability and understanding.  I further certify that the transcript has been prepared in compliance with transcript format guidelines required by statute or by rules of the board and that I have been informed about the complete arrangement, financial or otherwise, with the person or entity making arrangements for deposition services.  I further certify that I have acted in compliance with the prohibition on contractual relationships, as defined by Louisiana Code of Civil Procedure Article 1434, and in rules and advisory opinions of the board.  I further certify that I am not an attorney or counsel for any of the parties, that I am neither related to nor employed by any attorney or counsel connected with this action, and that I have no financial interest in the outcome of this matter.  This certificate is valid only for this transcript, accompanied by my original signature and original raised seal on this page.  Baton Rouge, Louisiana, this 11th day of November, 2020.

#### A

**Abbott** 123:20 124:11 abbreviations 123:3 abilities 8:3 ability 74:5 75:24 220:8 able 41:25 200:18 abscesses 89:14 absolutely 85:14 98:9 111:10 112:25 178:12 absolutes 107:15 **absorbed** 139:25 171:9 absorbing 136:10 abstract 23:25 academic 19:8 27:1 accept 100:3 106:5 107:17 109:16 137:17 accepted 28:14 33:3 accepting 104:25 access 139:23 150:12 accompanied 220:18 account 130:5 accurate 65:5 102:2 190:18 **accurately** 7:21 122:3 **ACE** 82:5 acid 39:6 141:9 acids 141:5 act 56:9 122:25 acted 30:21 34:7 220:12 action 58:4 220:16 actions 151:5.9 153:4 active 28:22 31:12,12 actively 19:11 activity 107:5 154:24 Actos 123:25 124:1 actual 69:19,21 105:23 106:23 143:17 acumen 75:8 **acute** 37:10 addition 15:12 24:16 40:17 150:11 additional 25:12 97:3 126:11 address 8:16 9:11 75:12 138:19 138:21 addressing 38:20

adenovirus 69:2

adenoviruses 86:25 **adhere** 178:2 adhered 178:6 adhesion 173:20 175:3 205:18 206:18 administering 6:9 **administration** 20:7,8 56:20 120:8 **administrative** 20:14 63:24 **advance** 11:23 advanced 48:25 127:9 advantage 140:25 adverse 120:6 **advice** 70:1 advising 20:16 advisory 56:20,21 57:1,16,16,17 57:18,18 58:1,6,14 220:14 aerate 171:22 **aerosol** 180:20 aerosols 180:16,22 184:13,16,18 **aeruginosa** 41:11,16 **affect** 145:22 146:9 147:2 172:12 affectionally 99:23 affectionately 130:17 affiliated 20:5 affluent 66:20 **afternoon** 6:1 7:7,10 agency 56:1 **agent** 41:3 106:18 111:14 191:24 191:25 agents 40:17.18.21 41:24 42:1.3 42:4 43:17 192:3 193:8 194:13 **ago** 19:21 44:9 50:6 93:11 94:3 96:11,12 101:3 107:9 109:24 209:9 **agree** 6:7 60:14 104:21 133:23 166:5 182:22,23 198:23 199:21 agreed 5:3 98:5 181:1 **ahead** 34:11 35:3 37:16,16,17 53:12 87:19 89:9 92:22 99:13 102:1 103:6 108:21 115:9 116:18 132:14.15 149:5 153:16 153:19 170:11 178:20 184:5 197:5 204:5 208:13,13,15

aimed 148:13 air 126:25 132:1 139:23 141:2,3 156:25 161:9,14 164:24 166:10 166:15.23 167:24 168:24 178:1 199:4 209:6 **airborne** 164:25 173:6 **airing** 189:10 alcohol-fueled 148:15 alive 140:5.6 178:18 **alkane** 175:6 **Allen** 3:4 6:13 92:14 133:19 148:2,25 152:8 159:9 188:17 193:17 197:8 217:1 allergies 68:14 **Allison** 3:14 147:20,21,22 allow 8:9 192:13 209:5 210:1 allowed 73:2,3 218:1 alternative 15:24 65:24 73:9 alters 173:18 174:21 207:22 **Alvendia** 2:4,4 6:16,16 91:11 92:14,23 96:19 99:11 105:17 107:7,22,24 108:14,22 110:4 110:13,21 118:21,24 127:22 128:1,6 129:19 133:18 138:1,3 138:6 148:2,5,25 149:4,9 152:8 152:16 153:3,9,21 159:7,15,17 159:22,25 160:6 162:24 164:9 164:11 184:3 188:15,23 191:4 191:17 193:15,24 197:7,12 200:8 201:17.20 202:10 215:9 216:21 217:5,8 218:3,6,11,14 218:18,22,24 219:15 **amalgam** 174:10 **American** 32:22 33:13 34:4,9,13 34:21 35:11,11 216:2,6 amicable 57:13 **Amino** 39:6 amount 213:13,14 214:2,4 **analysis** 11:12 38:16 46:18 47:24 48:16,20 49:13 61:6 63:1,10 81:8 105:21,22 123:14,17 134:25 165:18 168:16 172:6 173:15 178:25 179:1,2 181:7 197:1 212:13 217:13,20,24

**ailment** 38:25

219:8 analyst 46:24 **analyze** 16:20 23:13 24:2 39:23 39:25 48:3.7 **AND-** 2:8,14,18 and/or 110:1 **anemia** 73:6 anesthetists 85:5 announcement 12:12 answer 8:4.13 15:21 17:12 27:7 31:18 64:1,11 67:5 77:9,10 82:8 85:17 99:14 105:18 109:4 117:15,24 119:1 129:22 130:7 158:7 160:9.14.17.22.23 163:11 173:9 177:13 184:5 191:20 194:4 200:17,19,20 201:2,3,7,8 202:6,11 208:13 213:22 215:13 answered 128:7 163:1,8 184:4 217:13 answering 9:18 answers 7:19,20 159:20 anthropomorphize 140:24 antibiotics 40:17,23 41:1,20 43:18 anticancer 40:16,18,20 41:2,23 42:2,3 43:17 anticoagulation 125:7 antidiabetic 123:25 **antimicrobial** 42:1 46:12 anybody 126:5 142:16 170:3 211:11,13,17,20 anymore 31:11 215:21 anyway 219:4 **apart** 89:5 apologies 151:1 apologize 11:23 37:24 125:11 196:16 apparatus 46:14 **appear** 23:25,25 68:13 194:25 **appeared** 61:12 195:4 appearing 134:20 appears 168:13,13 applicable 181:7,9 applicants 75:20

**application** 12:14 15:3,19 52:22 53:8 57:12 82:13 **applications** 57:11 73:25 74:1,15 **applied** 18:15,18 53:5 127:12 128:18,20 139:11 158:2 **apply** 178:22 196:4 **appointment** 17:8,8 20:3,4,25 21:13,22 63:18 **appreciate** 11:19,19 50:13 53:14 122:15 **approach** 52:3 132:3,8,17 **appropriate** 21:19 139:2,5 179:17 **approved** 57:13 80:25 85:13 **approximately** 10:22 54:1 84:7 approximation 121:9 **April** 151:6,6 154:3,5,7,8 155:21 155:21 157:14,15,16,17 158:3 158:20,20,21 160:24 arabinoside 41:22 area 15:15 36:20 48:12 54:14 58:25 60:16 61:19,23 114:17 154:5,10,19 155:23,24 158:17 160:21 189:23 areas 15:6,18 22:8 37:6 38:6 148:14 151:23 171:14 argue 106:5 107:18 109:18 159:18 argument 97:12 99:4,25 164:20 164:21 arguments 116:12 arising 189:9,9 **Arizona** 1:15 7:2 arrangement 220:11 arrangements 220:11 arrhythmia 68:15 **arrhythmias** 55:11,14,16 61:21 **arrival** 189:16 arrythmia 69:22 art 132:5 176:22 artery 54:24 55:7 67:8 **article** 4:14.16 5:11 30:10 31:2 33:10 53:7 117:4 166:20 167:2 168:19 169:21,22 170:24 173:4 194:20,25 195:22 196:17,20

197:13,18 202:23 203:8 220:14 articles 30:23 32:11,21 36:17 55:5,13 60:11,15,18,20 61:2,23 117:4.5 133:4.10.12.20 134:12 134:14 143:9 194:22 195:8,11 195:13,20 196:19,23 197:9 201:13 articulate 7:25 articulation 11:20 arts 51:20 **Asha** 53:7 aside 38:19,24,24,25 131:6 **asked** 17:5 25:21 31:3 32:8,23 36:5.24 39:22 40:13 41:3.7.18 42:15 43:9 44:19 48:2,6,7 53:21 63:18 78:25 79:13 95:1,6 96:22 97:8,13 98:4 101:21 117:18,21 118:17 123:10,13,24 124:9,12 126:10,13 128:7 129:3 137:8 138:16,21 159:16 164:17 171:5 175:17 **asking** 8:1,5 64:4 109:21 130:13 152:18 159:7,12,14 193:17,19 193:24,25,25 200:10,11,12 204:13.14 213:23 215:10 218:9 218:12 aspects 136:2 aspirin 82:5 assert 39:15 assertion 201:15 assess 132:21 137:9 assessed 132:20 135:25 assessing 135:21 137:5 assessment 30:10,14 58:4 103:23 assimilated 139:5 assist 87:13 associate 22:7 **associated** 13:16 27:13 35:21 **association** 12:24,25 25:15,22 28:17 32:22 33:14 35:12 124:16 125:17 216:6,6 **assume** 8:4 32:15 74:14 107:25 109:6 111:2 115:2 158:1 176:17 183:7 assumed 219:2

assumes 105:22 108:11,16 218:6 **assuming** 58:23 178:11 188:14 194:4 **assumption** 105:15 106:3 108:19 109:2 111:9 138:22 **assumptions** 105:6 107:25 179:19 **assure** 104:14 astrophysics 43:6 ate 98:14 Atherosclerosis 35:15,18 atmosphere 126:25 atom 205:13 atoms 186:8 205:13.17 206:5 attach 140:11 145:17 146:22 attached 174:7 attachment 140:22 attained 10:23 attempt 12:4 attempted 189:7 attempting 44:8 80:3 81:12 178:3 attend 20:10,11 24:24 attending 25:8 attention 35:20 131:2 attorney 6:12 91:21 220:15,16 attorneys 126:11,14 attractive 136:19 attribute 12:4 August 151:7 155:18,22 160:25 **Austin** 13:11 **author** 59:24 164:18 authored 59:23 113:13 164:6,10 192:24 193:11 authority 220:4 authorized 5:6 authors 30:8,12 **autoclaved** 47:17,18,19 availability 207:9,10 available 207:11 209:4 Avendia's 108:8 avoid 83:8 186:3 award 37:13 71:9.9.10 aware 148:22,24 161:1 216:14 **awful** 43:1

В **B** 214:12 **BA** 10:4 back 17:25 19:25 20:2 29:21 30:22 37:19 41:5 44:8 51:17 53:19 55:3 59:13 62:23 72:10 88:10,17,18,20,20 91:14 97:9 105:11 112:2.22 128:15 131:22 132:11 146:18 147:19.24 148:7 157:15 160:8,13 165:20 167:4 167:16 175:13 197:14 207:15 **background** 9:23 47:10,14 74:20 75:1.6 95:2 97:10 118:14 129:9 138:24,25 144:22 backgrounds 84:9 bacteria 43:19 89:5,6,13 **bacterial** 72:23 74:7 bacterium 41:19 **bad** 26:14 33:9 57:22 72:24 **balance** 47:12 **Baltimore** 41:6 bar 178:1,3,3,6 212:22 bargain/negotiate/fight 57:11 bars 145:8 **basal** 176:20 **based** 53:2 95:2 103:23 108:18 108:19,22 109:6 113:1,6 136:18 140:9 146:10 147:7,8 150:16,17,17 151:2 152:25 170:22 176:7 199:22 203:20 207:9,9 **basic** 71:2 169:14,14 basically 189:5 **basis** 97:6 118:4,19 147:14,17 Baton 3:9 6:3 220:19 bats 111:11 **Baylor** 22:12,16,23 23:2,18,19 23:19.20 **bearing** 172:6 175:3 **becoming** 31:19 145:25 beg 116:15 168:12 began 27:10 28:14 88:15 97:21 **beginning** 55:4 65:2 145:5 **begins** 55:20 **behalf** 1:15 6:14,18 34:12 122:25

Behavior 65:21 behavioral 65:18 **belief** 190:23 **believe** 18:19 62:24 64:7 100:19 122:22 123:25 145:14 149:16 156:19 159:1 165:12 180:11 199:17 206:20 **believed** 96:23 165:13 **BELLE** 1:22 **Ben-Shmuel** 194:24 **beneficial** 81:4,7,18 **benefit** 59:19 78:19 Berlex 77:11 78:24 79:3 80:3 81:12.22 82:13 84:4 85:6 86:5 best 8:3 49:22 50:3 68:17 84:16 103:25 150:24 177:2 179:22,24 190:21 194:13 220:8 Bethesda 71:6 better 9:7 14:11 41:2 80:4 83:10 136:22 140:19 175:19 **beyond** 17:1 25:8 163:7 172:3 194:1 bias 38:5 55:7,18,19 big 93:13 102:22 **biggest** 136:8.17 **billion** 167:10 **billions** 130:1 167:15 183:3 binding 6:9 **Bio-funded** 37:19 **biometrics** 14:17 33:14,15,16,19 **biometry** 10:24 14:8,14,15,17,22 63:10 **biostat** 78:8 84:20 **biostatician** 22:12 23:7 biostatistic 66:10 biostatistical 66:7 biostatistician 18:9 63:21 85:25 biostatisticians 16:15 18:24 **biostatistics** 10:25 14:6,9,10,12 14:14,23,25 15:3,20 17:16 23:23 24:18 36:12 37:22 63:11 63:20 64:23 65:9 66:12 67:20 68:8 71:3.14.16.20 78:1 118:13 127:10 **birth** 38:4 67:10

**bit** 11:21 55:3 56:10 87:16 165:21 176:11 185:24 black 205:21,22 **bladder** 124:1.15 **bleach** 190:15 **bleach-based** 189:11 190:13 **blood** 49:25 71:21 73:22 74:9,11 **board** 18:21,24 57:1 58:14,16 74:8,10,12 77:18,22 78:1,2,4,6 78:7,9,10,15,16,22,22 79:10 80:8 82:12 83:23 84:4,5 220:10 220:14 **Bob** 53:7 body 14:18 103:1 118:5 130:7 132:18,24 203:10 **bold** 52:19 **bolts** 115:13 **bond** 210:23,23 bonded 210:19,21 211:10,22 **bonding** 205:10,12,15,16,19,25 206:18 **bonds** 205:23 207:3 208:18,18 208:19,19,21 **bone** 40:21 **bonus** 122:7 book 35:23,24,25 36:6 56:16 60:7 62:17 194:20 **books** 56:16 62:10,14,14,19 169:14.14 **bottom** 62:13 **BOULEVARD** 2:21 **bounds** 111:7,15,18 **Bourbon** 148:13,20 149:5,7 **boy** 67:5 **Bradford** 12:22 13:8,11 66:16 **brain** 209:8 breadth 71:8 break 8:8,11,12,13 45:5,8 91:9 142:17,22 152:9 178:5 186:7,7 188:12,24 breast 39:7 45:23 46:2,6,8 **brief** 185:4 197:25 198:1 briefly 10:3 **bring** 65:8 75:5 84:10 broad 15:8,17 35:18

broader 43:13 broken 89:14 210:23 brought 46:6 78:6,9 building 138:20,23 154:20 155:15 157:21,24 158:5,13 171:23 175:25 216:18 217:6 Buonanno 165:3,4,4,8,24 175:16 175:17 176:4 burden 63:25 163:5 businesses 93:3 butcher 175:15 butchering 125:11 buttress 113:5 buzz 69:5

 $\mathbf{C}$ 

C 2:1 3:1,4

c-a-r-d-i-o 58:22 c-h-l-o-r-i-d-e 206:16 **CAJUN** 1:3,3,4 cake 29:6 calendar 120:22 121:8 caliber 23:20 **call** 82:2 92:9,11 93:2,8,18,23,25 94:4,12,17,18,23 95:12 96:9,18 98:1,7,9 99:5,23,25 100:9,11 102:24 105:9 114:2 130:17 called 10:24 14:9,13,14 30:11 34:2 57:16 85:2 141:8 179:4 **CANAL** 3:5 cancer 40:15,15,19,20 44:10 86:21 124:1.15 **candidate** 74:22 75:16 candidates 74:22 **Cantrell** 145:21 **Cantrell's** 147:15 **Canyon** 210:13 capacity 85:23 122:20 carbenicillin 41:21 carbon 206:5.8.15 cardboard 180:21 cardiac 54:24 55:11,14,16,18 61:20,21,21 68:15 **cardio-renal** 58:14,22 cardio-type 67:22

cardiologist 84:12,18 cardiology 10:16 15:10,23 35:17 35:18 65:11 80:12 84:11 **cardiovascular** 37:6.10 38:1.3 58:25 59:3,17 67:22 71:21 80:5 80:20 81:4,6,7 82:15,18,20 125:8 care 25:25 28:14,15 47:16 55:19 career 11:3 29:6 38:21 careful 134:20 **carrier** 199:18 **CARS** 134:16 **cascade** 109:13 case 1:4 50:8 51:25 54:16 84:14 89:22 97:9,11,15 99:7,9,16 102:4 113:15 117:20 118:9 122:4 123:10,12,13,20,21 124:14,14,15,19 125:6,13 126:2 132:21 146:13 156:5 163:6 196:5,22 199:20 200:13 202:16 212:1 cases 93:15 114:14 119:5,6 122:17,19,24 125:19,23 **casino** 54:16 catch 198:19 **caught** 141:1 causal 13:4 125:16 causality 12:22 13:8 66:16 causation 124:17,18 cause 1:16 69:3,17 88:9 100:22 101:14 104:14 138:23 176:23 209:20 213:1,3,4 214:3,4 caused 41:17,19 68:15 86:25 causes 13:15 69:15 146:22 214:10 causing 89:2 131:3 145:19 146:5 146:7 147:8 CCR 220:24.24 **CDC** 155:24 158:2 159:5 160:19 179:9,16,23 192:15 204:12,13 204:15,18 CDC-approved 190:25 **cell** 37:10 72:3,12,19 73:6,6,10 73:11 75:13,25 76:5,8 77:13 139:25 140:13 141:10,18

176:23 199:13 cells 141:6,15,15 198:19 199:13 **Celsius** 181:17 center 16:5.10.12 17:3 18:1.3 22:18,22 27:12 120:16 121:1 158:16 193:4 204:7 centers 84:25 190:17 192:9 194:11,12 **cerebri** 124:7.16 **certain** 1:7 41:1 103:12 145:20 146:6 162:17,18,19 163:13 certainly 32:3 38:17 39:14 71:17 85:9,10 117:10,25 118:1 131:1 139:4 163:13 174:14 177:13 certainty 159:11 **certificate** 4:6 220:1,17 certification 18:17.18 certifications 13:25 **certified** 1:17 6:3 220:2 **certifies** 18:24,25 certify 220:3,6,9,12,15 **chair** 84:21 85:2,2 **chance** 183:15 198:10 **Chandler** 1:14 7:2 **change** 8:24 131:9 150:18 151:2 151:12,18 152:25 162:6 166:3 167:20 169:18,23 170:7,15 188:8 209:1,2 213:4,15,21 214:6 **changed** 83:1 141:7 157:24 **changes** 157:10,11 161:17 168:3 188:3,4,5 209:5 **chapters** 56:17 62:18,18 **character** 205:19 206:19 characteristics 102:22 112:6 130:1,2,5 **charge** 52:18 122:12 **check** 32:6 57:3 152:2 188:16 **checks** 217:25 219:9 CHEHARDY 2:19 **chemical** 88:19,20 101:19 161:25 **chemist** 42:18,20 117:23 **chemistry** 41:9 42:16,21 95:18 102:8,12,13,20 103:22 105:2 106:1,2,6,11,11,13 113:24

117:6,8,9,10,11 118:1,3,4,4,14 119:8,10,21 129:6 133:25 139:3 140:8,9 141:2 162:2,4 168:16.17 169:9.14 170:23 178:3 206:11 chemistry/physics 105:22 chemotherapeutic 40:17,18,21 41:3,23 42:3,4 43:17 **children** 72:20 73:10 74:6 75:25 88:4.7.16 90:13 **Chin** 184:22 185:1,3,16,25 China 175:22,25 **Chinese** 176:8 **chloride** 206:16 **cholesterol** 38:5 54:25 55:8,20 59:16 choose 29:9 **chose** 29:10 164:18 circumscribed 135:23 circumstance 178:13 **circumstances** 122:10 145:20 146:6 citations 168:7 cite 156:14 169:11,11 173:3 189:22 **cities** 150:1 city 148:14 149:22 216:10,16 civil 1:1 5:7 163:6 220:14 clarification 92:24 110:8 **clarify** 24:5 80:16 92:16 101:22 108:20 clarifying 187:15 **clarity** 37:24 79:2 108:9 206:24 class 57:24 58:5 63:19 64:18 211:15,18,21 212:6,9,15,18,18 212:20,21 classes 20:17,19 40:25 211:15 classically 141:7,8 classroom 65:9 cleaned 210:14,17 211:12,16,19 211:22 212:10 **cleaner** 191:6 cleaners 160:18 189:11 190:13 190:25

191:24,25 192:3,16 193:8 194:6,13 204:1,8,16,20,22,24 206:12 209:25 210:15,19,20,24 211:4 clear 17:15 19:6 38:19 51:9 58:12 67:2 97:6 102:11 112:25 118:18 120:25 172:17 176:3 195:19 213:11 **clearly** 63:9 108:3 116:1 134:23 152:11 170:6 **cleaves** 173:25 clinic 26:22 27:11 **clinical** 12:8,11,13,19 13:6 15:12 16:5.21 17:3 18:2.4 22:12 23:7 23:14 27:2,23 29:21 31:24 32:2 32:24 33:22,25 34:2 37:12 50:18,23 51:3 64:8,24 65:8 66:3,15 67:21 73:8,14 75:8 76:2 78:8,17 79:5,18,18 84:25 84:25 88:11 127:11 128:17 194:17 clinics 25:19,24,25 26:1,4,6,17 27:17 28:15 **close** 89:7 104:11 145:8 148:18 148:20 179:11 188:20 **closed** 148:16 150:19 151:4,24 152:6 153:2,12 154:5,10,15 155:8,11,23 closely 17:20 **closing** 153:7 189:12 closure 145:9 cloudy 186:19 **CMI** 194:22 co-teach 63:19 coat 177:5.6 Code 5:6 220:13 **cohesion** 173:20 175:3 coiled 141:11,17 **coin** 53:25.25 54:2 Colaneri 203:16,16 **cold** 69:1,3,8,18 70:6 101:10 **collaborate** 22:19 64:17 collaboration 21:8 64:15 colleague 196:15 collect 12:3 43:16 120:18

cleaning 152:4,6 179:15 189:11

collected 23:24 **collection** 13:10 25:18,23 30:7 32:10 76:20 120:6 134:6,7 **College** 216:2 combination 13:1 40:19 49:8 **come** 19:25 26:19 39:22.24 74:13 95:15 103:10 105:6 112:21 133:8 144:2 146:18 156:12 199:24 comes 56:12 111:3.10 158:9 186:23 199:1 214:25 **comfortable** 26:18 183:8 coming 127:4 137:4 168:8,10,11 183:19 **COMMENCING** 1:25 committee 57:16,17,18,19 58:1,6 58:13,16 59:4 84:22 85:3 committees 58:13 **common** 41:11 55:2 69:1,3,8,17 70:6 101:10 129:25 130:2 144:4 commonly 28:18 40:16 48:6 61:8 communicated 128:2 **community** 26:8,9 56:7 70:2 71:18.19 73:16 76:9 77:5.21 companies 57:7,10 97:12,13 123:6 company 57:20 79:3 123:4,4,11 company's 57:12 98:4 comparable 183:12 **compare** 175:25 compared 23:8 comparison 130:16 compiling 39:1 complete 104:18 109:25 116:21 155:2,5 206:1 220:11 completed 27:9 completely 11:14,15 150:4 complexity 33:23 **compliance** 220:9,13 **complicated** 84:23 212:19 complications 86:18 **component** 8:21 66:8 69:25 106:2 119:4 132:19 **components** 11:3 119:3

composition 39:6 **compound** 152:17 **compounds** 113:23 175:7 compromised 199:9 computer 8:23 conceding 187:5 concentrated 213:14 **concentration** 182:1,5,9,15,22 182:24 183:4 185:15 198:17 199:11 213:20 **concept** 28:14 concern 46:11,13 211:21 concerned 144:21 168:21 **concerning** 51:2 196:10 concluded 125:16 219:17 conclusion 13:2 95:15 98:25 99:8 100:18 103:2,10 105:7,13 108:17 109:1,10 113:10,14 114:22 127:4,19,21 128:2,9,19 128:22 129:15 133:8 136:24 137:7.23 138:8 146:18 147:13 167:25 172:18 184:13 192:8 199:25 215:11 **conclusions** 108:3 109:5,19 127:7 133:16 135:5 137:5 138:11,13 139:17 146:17 147:3 147:4 165:14 214:11 216:9 **condition** 150:18 151:3,12,19,20 152:1.20 153:1 161:8 166:3.14 166:22 172:11 184:17 187:19 187:24 190:2 201:16 **conditions** 180:18,20 181:12 182:13 183:10,11 184:2,2 185:1,12,23 187:10,13,16 188:2,7 204:1 211:14,25 212:3 conduct 74:3 78:16 85:1 conducted 161:25 conducting 79:5 **conference** 112:3 128:1 **confirm** 107:3 133:15 **confirmed** 200:6,13 201:5 conflating 153:12 conflict 164:2 confuses 188:9 **confusion** 150:1,5

connected 220:16 conscience 122:12 consider 38:21 40:1 44:12 49:6 54:25 117:22.23 219:8 considerable 24:10 **consideration** 30:15 147:14 216:9 217:14,21,24 218:10 219:4 considered 5:12 15:16 19:14 49:5 114:19 133:2 158:25 consist 99:1 consisted 99:2 180:18 consistent 163:23,24 164:1,21 204:14 constantly 217:5 constrict 14:20 **consult** 137:23 consulted 137:6 **consulting** 22:11 23:2,7,16 contact 199:17 contacted 200:23 contain 142:7 contained 26:25 112:12,13 139:8 156:9 176:12 contains 141:9 contaminate 146:22 **contamination** 45:20 46:13 147:7 203:24 204:9 contemporary 33:25 contending 134:12 content 123:14 141:9 **contentious** 57:14,15 contents 98:3 161:10,18,24 166:16,23 168:4,25 169:2,6 171:1 context 69:8 80:11 126:22 Conti 1:3 105:23 106:19 148:18 149:2 continue 29:8 78:20 184:8 188:8 192:21 continued 3:1 36:12 189:16 continuing 107:8 continuous 45:21 contract 176:25 208:7 209:11,21 209:24 211:3

**contracted** 90:14 151:14,15

154:3,14 155:13 200:6,15 201:11 202:3,8 contracting 201:5 contractual 220:13 contradicts 203:12 contributed 48:12 contributing 59:24 **contribution** 46:15 47:23 64:21 66:6 137:10 contributions 136:2 control 158:16 174:17,18 190:17 192:10 193:4 194:11,12 204:7 control.' 148:16 **controlled** 33:21 34:2 181:14,20 181:25 182:13,14 185:9,10 189:24 controversial 61:10 **CONVENTION** 3:9 **conversation** 102:9 158:2 converted 142:4 **Cooperative** 50:19,22 **coordinating** 16:4,5,9,10,12 17:3 18:1,3 120:16 121:1 **copper** 180:21 copy 141:11 143:2,4,11 cordoning 189:21 core 39:17 corner 149:2,3,11 **corners** 133:17 **coronary** 54:24 55:7 67:8 coronavirus 69:2,6,12 93:3,15 93:22 **coronaviruses** 69:16 93:19 94:9 Corporation 123:11 correct 16:7 19:5 46:22 47:1,4 55:11 57:17 59:22 60:20 62:8 62:25 64:19,22 65:10 69:16 72:13 73:7 98:22 99:8 105:19 106:14,16,17,20,25 108:12 111:21,24,25 113:16 114:11 121:25 128:5 134:4,4 135:14 137:14.14 151:22 154:1 156:16 159:11 165:25 175:22 178:10 178:24 181:12,18,20 182:18

185:1,13 186:13 189:25 192:15 193:3 194:9 196:25 197:22 202:18 207:4 209:13,14 210:5 212:14 217:17.17 220:8 **correspondence** 56:16 61:25 62:7.11 185:5 corrode 212:21,24,25 cost-effectiveness 65:19,22 cough 157:9 198:14 **coughs** 198:12 **council** 56:21 counsel 6:5 220:15,16 **couple** 22:1 48:22 78:13 96:20 101:22 125:21 128:23 133:12 141:7 150:1 166:4 course 8:25 10:16 11:24 12:2 20:21 30:1,1 32:18 35:19 40:10 63:22,24 86:9,16 117:7,8,9,11 117:14 131:2 134:2 140:4,23 141:3 173:22 186:17 206:15 208:22 209:17 courses 24:16,18,23 25:1 65:1 66:10 **court** 1:1,17 6:3,4,8 7:19 55:12 116:22 124:15 144:19 160:9 202:24 220:2 CoV 99:17 129:14 CoV-2 100:4,7,8,10,11,14,19 104:6 105:11,23 106:15 107:3 108:11 110:1 111:4,11,12,12 111:13 113:18 115:7 119:15 126:16 128:3 129:14 130:25 131:3 138:18,22 142:13,14 176:12 177:1,2,10 187:10,12 187:23 201:15 213:13,14,20 214:2,4 covalent 205:15,16,18,23,24 206:18 207:2 208:19 covalently 210:19 **COVID** 119:7 171:24 196:11 197:1 **COVID-19** 4:15 69:10,15 96:24

154:14 155:13 157:20 161:2 170:2 176:25 189:13 200:7,15 201:5,11 202:3,8 203:1 209:21 209:24 **COVID-2** 130:15,19 131:14 crannies 179:14 create 81:12 119:21 193:6 202:17 204:9 **created** 21:13 81:23,25 82:1,14 128:3 131:7,8 135:20 212:2 creates 131:9 192:25 201:16 creating 79:7 **creed** 136:22 critical 132:2 135:25 criticism 185:3 **criticize** 179:23 **cross-pollination** 18:10 **CUISINE** 1:4,4 **culture** 125:22 **cure** 80:3 **current** 22:24 36:4,4 37:4 187:8 187:11 currently 31:14 currents 178:1 curriculum 4:11 13:17 20:11 **custom** 36:20 125:22 customer 103:19 156:23 cut 86:4 CV 9:21 12:6 18:1 21:20,22 22:2 29:22 31:15 32:4 37:1 54:13,13 56:8,11 59:14 60:9 64:5 73:24 cytosine 41:22 D

D 4:1
d/b/a 1:4
damage 95:5,7,11,16,20,23 96:7
96:24 98:25 99:8 100:24,25
101:6,15 102:6,10 104:13,14
105:15 106:11 111:13 119:11
119:21,23,24 128:4 129:12
131:10,16,20 134:3 137:25
138:23 145:19 146:6,8,11,23
147:9 162:7,8,12,13,20,22
169:19,20,24,25 170:7,20,20

99:25 100:4,10 105:10 109:8

110:2 144:15,20 145:4,6,6,14

145:25 150:10 151:14,15 154:4

170:25 171:5,9,12 172:7 174:24 212:14 214:14,20,21,23 215:1 **damaged** 215:20 damages 214:24,25 damaging 162:11 **Dan** 6:17 dangerous 41:16 150:19 151:3 151:13,19 152:19 153:1 157:1 157:11,14,25 158:25 166:11 172:11,11 193:1 201:16,16 202:17 **DANIEL** 2:15 **Daniels-Feasel** 124:20 dare 102:21 dark 186:2,12 187:1,2 189:23 darkness 189:24 **Darwin** 19:19 data 11:12 12:3,25 23:24,25 24:2 39:1,23 40:3 43:15,16 46:17,24 47:24 48:15,20 61:4,8,12 73:1 73:1 74:4,6,6 77:18,22 78:17 78:22 79:9 82:12 83:22 120:16 121:1 123:14,17 165:9,15,23 175:18 dataset 39:25 date 36:15 60:15 dated 37:2 207:13 dates 36:14,17 151:5,8 153:4,5 153:14 155:21 **dating** 62:23 daunorubin 41:22 Dave 91:17,18,20,21 92:3,17 93:10 126:4 **Davillier** 2:15,15 6:17 62:24 day 48:7 158:19,20 186:18 187:4 187:6 188:3,3,3,3,4,4,5,5 220:19 days 27:14 88:13 130:22,22 150:1 157:16,17 186:19 189:19 days' 57:4 ddavillier@davillierlawgroup... deal 24:19 35:20 72:12 **deals** 62:11

**dealt** 59:16 62:8 113:17 114:12 114:14 **debate** 184:7,8,15 **debated** 164:18 decades 35:21 **decay** 180:17 **December** 77:2 127:24,25 **decide** 49:23 **decided** 88:14 **decision** 21:21 22:2 90:16 103:18 145:8 decisions 57:13 decorated 29:16 **deduced** 44:24 50:7 deducted 79:24 deem 60:13 **deep** 131:17 211:18,22 **deeper** 210:15 **deeply** 192:2 211:9 **DEFENDANT** 1:15 3:3 **defendants** 6:14 122:25 **define** 81:17,24 82:24 170:7 214:22 **defined** 81:16 82:24 170:6,8 207:8 220:13 **defines** 170:25 definitely 77:11 **definition** 32:9,15,17 44:16,20 44:25 46:20 50:5 51:18,20 52:11 57:25 66:15 83:2,3 95:19 96:3 101:4 170:10,12,12,13 171:6,11 214:16,19 215:11 **definitions** 205:22 207:7 degree 10:20,21 11:6 19:1,2,4,8 24:8 43:20 154:21 156:24 158:23 159:10 degrees 10:2 19:6,7,10 42:8 181:17,17 182:14 **deleterious** 161:7,8 166:22 deliberately 81:21 **deliver** 81:2,5 delivered 82:10 **delivery** 15:14,16 **DEMARTEST 2:4** demonstrate 105:1

demonstrated 212:6 demonstrates 145:1 denature 178:18 186:9 192:10 192:13.17 193:8 194:8 204:19 denatured 178:16 179:5,6,9 191:22,23,25 193:14 210:2,3,4 211:17 density 103:11 114:14 deoxyribonucleic 141:9 **department** 17:14,16,18 18:14 19:13 20:9 21:5 22:4 **departments** 16:14 17:24 21:17 22:19 **depending** 16:17 200:24 depends 83:24 88:25 89:1 101:12 130:23 131:19 154:22 177:3 **deposition** 1:12 5:4,13 7:13 9:1 115:22 116:1 122:11 219:2 220:12 depressed 40:22 der 206:24 207:1 208:20 **describe** 12:19 13:5,13 20:25 30:17 33:6 37:9 87:24 171:16 205:10 **described** 39:8 77:20 80:11 146:2 174:16 describes 167:7 **describing** 86:23 87:11 137:14 168:22 **description** 20:24 65:6 descriptor 125:5 **design** 12:23 16:20 23:13 38:15 39:16,20 41:7 48:8 49:13 50:22 51:11 64:7,24 75:18 130:25 **designed** 48:2 73:5 **despite** 159:2,3 194:10 201:9,12 destabilize 186:8 **detail** 9:22 36:21 133:2 185:8 detailed 185:24 203:11 details 80:6 123:16 determination 30:6 102:6 determinations 57:21 **determine** 51:11 73:2 78:18 137:6 146:10 173:25 202:2

**determined** 84:16 88:23 89:17 **disciplines** 10:13,14 16:14 17:24 91:13 92:18,22 99:11 107:11 90:9.12 84:2.8 108:9 109:3 110:22 119:1 determines 104:22 **discount** 122:13 128:15 129:22,23 137:18 138:3 **determining** 13:14 131:25 172:4 discovery 5:5 143:2.15 153:21.25 163:11 172:9 **discuss** 101:18 184:5 189:2 191:20 194:5 **develop** 87:10 discussed 22:9 76:22 77:14 98:3 201:23 205:9 216:1 217:12 **developed** 84:19 87:21 120:12 126:15 **Doctor's** 160:8 development 37:21 discusses 120:3 **doctors** 82:23 diabetes 86:20 discussion 166:9.12 189:4 **document** 133:17 diabetologist 16:17 **discussions** 68:16.17 documents 9:2 diabetologists 23:12 disease 13:3,15,16 16:19 44:24 **DODD** 3:8 diagnosing 26:15 45:1 50:7,8,9 51:3,21,25 52:2,2 doing 22:13 23:8 29:7 44:1 83:15 **diameter** 143:24 83:17 90:13 111:24 120:23,24 52:3 54:25 55:6,7,8 64:5 67:8,9 diarrhea 88:7 67:22 69:25 72:19 73:10.11 147:6 diarrheal 88:15 77:13 79:24 80:2,6,20,21 82:15 door 119:7 **diction** 11:19 82:18,21 89:2 124:14 144:20 doors 171:23 172:2 183:18 **died** 19:20 145:25 158:16 190:17 192:9 186:11 diem 121:11 193:4 194:11,12 204:7 **dope** 61:11 difference 60:25 131:1 141:4 diseases 51:25 59:8 69:23 86:17 **Doremalen** 180:4,8,9,11,15 86:18,19,21,23 87:10 186:20,20 187:4 205:1,2 181:2 184:25 **different** 16:14 17:24 22:13 **disinfectant** 179:15,17 **dose** 84:16 33:24 43:13,19 53:5 58:13 61:7 **disinfectants** 158:3,15 179:9 doubt 194:2 61:10 74:1 91:24 112:24 192:16 204:23,24 **Dr** 4:12 6:21 7:7 37:11 115:9,13 113:10 130:2,7,25 131:20,21 disinfected 155:24 159:3 147:23 219:8,16 144:11 152:11,22,23 153:5,6 disinfecting 152:4 draft 98:24 102:14 112:5 153:10,14 171:9 175:5 183:9 dislodged 208:22,23,23,24 **drafted** 116:7 133:11 **drafting** 127:17 215:17 188:9 206:25 207:2 disparagement 160:18,19 **dilate** 14:20 disparagingly 78:5 draw 127:7 128:18 136:24 dissemination 164:25 **Dilemma,'** 68:7 **drinking** 216:25 217:18,19 diligence 76:1 dissertation 53:3,8 drip 45:21,22 46:14 **Dillip** 67:14 drive 1:14 7:2 113:9 **distance** 103:12 **Dillip's** 67:17 distinct 182:1 driven 177:25 driving 145:7 direct 126:14 distinction 12:18 14:7 20:4 23:6 direction 21:6 169:4 220:7 62:1 63:14 89:20 droplets 89:14 173:1,2,7 196:7 directly 115:24 200:20 distributed 15:25 199:2.3 director 16:3,4,4,22 17:1,6 18:1 **DISTRICT** 1:1 **dropping** 210:12 disadvantage 56:11 diverge 14:16 **drops** 65:4 **disagree** 19:9 47:8 66:6 105:25 **divide** 119:3 drought 52:23 135:4 136:6 137:16 192:1 divided 46:4 **drug** 49:24,24 56:20 57:10,11,12 197:17 199:21,23 203:8,19 **DIVISION** 1:5 57:12,20,23 59:2 68:18,20,21 disassemble 209:13,18,19,24 **DNA** 140:17 141:4,8,9,11,15,19 68:22 69:20,24 70:2 81:8,10,11 81:17,19,22,24,25 82:2,3,6,7 disassembled 174:22 207:24 141:22,25 142:7,12 82:14 83:3 120:8 123:4.4.5 208:4.6.9.11.11 209:11 **Doc** 92:15 discarded 135:3 doctor 7:14 9:14,22 11:18 13:23 124:1 **discipline** 11:7 118:8 45:15 48:11 59:15 71:17 87:13 drugs 49:22,23 50:1,2,4 58:5,17

58:24 59:10,11 79:1,6,12 82:4 **DSMB** 77:13,17,18 85:4 **due** 78:20 214:20 **duly** 7:3 220:4 **Dunbar** 3:4,7 8:23 **durability** 102:25 112:8,12

113:18 130:15,16,18,19,24

131:5 **duration** 17:4 180:23 **dying** 140:23

 $\mathbf{E}$ **E** 2:1,1,15 3:1,1 4:1 146:18 **e.g** 189:9 earlier 14:8 61:22 66:14 117:21 120:12 126:4 129:17 133:1,10 133:19 148:17 176:11 early 78:19 87:4,6 91:2 116:8 196:25 **easier** 143:11 easily 208:9,11 209:10 easy 140:23 184:9 eat 29:6 192:11 eating 216:24 217:18,19 economics 15:14 edification 130:14 **editor** 30:14 **editorial** 60:24,25 61:7,7,15 **editorials** 56:16 60:24 61:10 **editors** 32:25 educate 11:22 **educational** 9:23 11:2 27:24 **effect** 40:20 72:19 81:4,6,7,18 82:17 94:9 101:19 105:13 125:7 144:15 150:2 effective 68:18 **effects** 69:24 186:3 efficiency 48:11 **effort** 83:18 135:22 189:14,15 190:21 **efforts** 189:5 egress 102:23 **Eight** 48:22 either 80:2 100:9 168:7 182:11 199:10 210:17

ejection 55:18 elected 145:2 **electron** 205:13,24,25 electronic 143:4.19 electrons 173:24,24 186:8 205:17 207:10.11 **Eli** 123:10 **eloquently** 93:1 136:5 else's 61:13 170:3 embarrassments 136:17 **embedded** 190:14 192:2,20 210:18 211:2,9 emergency 10:16 emit 172:24 **employed** 29:11 55:25 56:8 137:19,21 220:16 **employee** 150:15 **employees** 110:7 150:14 189:17 encourage 22:21 **encouraged** 17:23 21:12,15,17 22:18.21 ended 23:5 28:24 29:3 endocrinology 10:16 enforced 204:2 engage 92:8 **engaged** 91:16 engagement 123:18 **England** 34:19 **English** 11:18 96:5 173:23 **ensure** 32:14 **entails** 11:13 **entered** 126:16 entering 167:7 **entire** 17:3 60:21 73:5 131:25 138:7 entirely 57:23 **entirety** 105:21 entitled 1:16 29:23 33:18,21 123:10 entity 215:20 220:11 **entry** 16:2 17:7,25 20:2 31:25 33:12 35:23 49:2 50:15 enveloping 177:5,6 **environment** 12:20 126:21,23 132:23 134:10 156:25 157:11

157:25 161:6 166:10,13 167:19 167:20,21 172:5,11 174:10 180:4 184:10 189:8 193:1,6 202:18 204:9 217:18 **environmental** 45:20 126:19 157:10,13 158:24 166:10 180:20 183:10 203:24 environments 132:1 **enzymes** 141:25 epi 21:23 78:8 84:20 epidemic 87:21,24,25 88:4,24 89:18 90:2,6,9,15 **epidemiologic** 64:7,10 66:7 103:21.22 146:3 epidemiological 101:19 epidemiologist 12:18 13:14 18:11,13,15,16,22 19:13,14 56:9 85:25 epidemiologists 16:16 17:20 18:3,8,25 20:6,19,21 23:11 25:12 63:22 64:16 90:17 **epidemiology** 17:8,9,15,18 18:14 19:2,5,7,18,20 20:3,17,18,18 20:22 21:1,19,22,25 22:20 23:22 24:8.9.10.12.13.17.19 25:7 29:21 31:25 32:2,23,24 34:4,13 37:8,14,22 38:7,14,17 38:22 39:9,17 40:1,3 44:13,16 44:21,22,25 46:20,21,22 47:12 47:22,22 49:5,6 50:5,18 51:15 52:12,25 53:10 55:1,5,10 56:1 60:14 63:3,16,19,22 65:7 66:11 67:1,3 74:20 75:1,5,6,9 77:25 79:22,23 88:11 102:1,4,5,9,11 103:24 118:12,17,19 119:4,5,6 119:13,15 124:12 127:10 136:23 139:16 216:3 epidemiology-based 55:21 epithelial 176:20 equal 131:7,8 especially 74:16 **ESQ** 2:4,5,10,10,15,20 3:4,8,8 essavs 88:20 essence 60:12 79:16 essential 141:6

exposure/disease 80:19

essentially 10:14 13:12 29:3 33:6 39:23 40:13 48:1 58:15 99:17 131:16 175:4 208:21 210:11 **established** 13:11 148:17 156:22 202:14.15 establishments 193:5 estimated 26:3 estimating 180:17 estimation 163:21 European 35:10 evacuated 87:14 evacuees 88:15 evaluate 79:13 evaluated 180:15 evaluating 58:24 81:9 137:22 event 28:25 53:4 54:10 55:20 events 98:6 120:6 199:1 209:20 **everybody** 78:3 169:16 evidence 132:3,8,17,18,24 133:3 133:7,16 135:20 139:17 213:14 213:21 214:6 evidenced 49:14 135:5 evident 133:9 evolved 36:9 **evolves** 14:11 exact 82:9 113:14 **exactly** 11:9 12:1 14:24 16:8 17:11 29:25 73:22 132:7 150:21 185:7 191:1 **exaggerated** 4:14 196:10 202:25 **examination** 4:4 7:5 20:13 **examine** 102:23 190:12 **examined** 7:3 132:19 **example** 15:10 20:9 22:7 26:12 41:12 70:16 82:4 86:22 89:3,5 122:10 134:16 172:23 174:9 200:22,22 206:2,17 **examples** 39:10 70:17,18 71:15 exams 118:3 **exceed** 163:20 exceeded 163:20 164:4 **exception** 12:24 48:23,23 **exceptions** 33:8 48:22 excluded 26:22 134:25

exclusive 211:25

**exclusively** 59:3 72:22 205:23 **excuse** 51:13 75:8 152:4 157:4 174:12 **execute** 16:20 23:13 executed 48:2 **execution** 38:16 39:16 **exercise** 46:20 132:2 **exhaled** 89:14 **Exhibit** 4:10,12,13,16 13:19,20 115:10 203:1,3 205:5,6 **EXHIBITS** 4:9 exist 190:2,5 200:22 existence 177:9 exists 102:25 **expand** 36:13 83:3 **expanded** 36:10 **expect** 53:25 155:20 182:10 expected 85:25 **experience** 13:24 14:5 16:3 23:22,22 24:20,21 25:11 26:12 27:4 47:6 56:17 57:24 62:22.23 64:23 75:17 79:17 96:5 108:24 120:3 127:8,11,11 128:11,17 129:6 133:5 139:16 183:20 **experiment** 176:6 180:23 212:7 212:8 **experimental** 63:3,16 65:7 180:18 184:17 **experiments** 130:23,24 **expert** 97:9,14,14 98:5,5 105:5 117:25 118:1.6.9.11.12.17 122:20 159:10 193:10 219:1 **expertise** 23:15 26:11,20 48:9 65:8,9 66:3 75:17 95:3 128:17 **experts** 23:12 76:21 174:14 **explain** 130:25 208:5 210:6 **expose** 41:10 209:3 **exposed** 41:21 42:2 50:7,8,10 79:24 83:12 209:4 211:19 **exposure** 13:3,15,16 24:13 41:10 47:12 50:10 51:5,8,22 52:6 80:21,22 81:14 83:8,9 84:13,15 84:19 124:13 131:6.8 145:15 156:25 157:13 158:24 166:10 **exposure**/ 44:23

extensive 8:10 62:22 130:13 extent 129:24 131:20 extrapolate 12:16 **extreme** 89:11 eve 213:5,6,15,21 214:7,10 eyes 199:10 F **F** 120:8 face 199:10 203:20 facetious 85:20 facilities 126:16 facility 16:13 87:9 105:23 106:16 108:12 154:4.10.15.16 **fact** 8:1 17:22 22:4 41:25 48:11 79:20 83:16 98:20 100:13 104:4,15 112:1 129:13 138:9 151:13 156:3 159:2.3 164:19 177:17 178:20 180:21 182:10 183:25 186:1 190:13 200:11,12 200:21 201:4,9 202:8 218:13 218:21 factors 133:2 182:17 facts 163:21 218:6 faculty 16:13 17:4,21,22,23 18:9 20:24 22:3 25:8 29:14 Fahrenheit 181:18 **failing** 149:22,22 fails 49:24 178:4 **fair** 7:22 8:6 15:21 31:1 33:2 53:25 54:2.2 64:25 66:22 71:12 74:16 76:3 79:16 96:16 98:10 99:24 100:1 101:24 111:4 119:12,14 132:6 137:18 138:2 155:1 175:24 186:10 187:25.25 201:25 202:4,5,12 212:15 214:1,1,1 217:7 **fairly** 210:24

**fall** 43:6 63:3 64:3 65:2,3,4 101:4

**familiar** 100:12 165:4 190:6

167:10 210:14

194:17 195:9

falling 67:1

**falls** 39:8

**Family** 34:21 **far** 26:25 60:9 61:2 72:12 76:4 77:9,10 84:19 89:5 198:21 210:18 211:18 Farenheit 182:14 **fast** 58:21 **faster** 52:15 **fault** 23:3 151:2 **favor** 152:9 **FDA** 57:6,10,15,20,25 85:13 120:9,12,21,24 121:2,6 federal 56:20 120:8 feeding 45:23 46:4,14 **feedings** 45:21,22 46:3,7,8 fees 122:2,3,6,7,9,13 **feet** 141:16 **fellow** 18:9 30:12 field 14:23 35:18 37:6,8 56:1 60:14 76:21 136:19 fields 15:9,12,23 33:5 53:5 **fifth** 67:14 **fight** 58:2 **figure** 152:12 **file** 149:22 **final** 138:8.11 **financial** 220:11,17 **find** 80:3 85:15 126:12 146:7 179:13 **finding** 203:12,22 204:14 **findings** 12:4 113:5 136:18 143:8 203:21.23 fine 53:14 91:10,11 97:20 124:24 142:6 fingerprints 14:21 finish 92:22 99:13 135:9 160:11 fire 89:11 firm 92:1.2 **first** 7:3 8:17 9:12 14:5 16:9 17:13 22:2 24:13 27:1 28:15 31:25 40:7,11 49:24 51:21 52:19 58:13,18 60:5 72:10 73:15 88:13 93:2 94:4.14 96:9 96:18 98:14 115:17 119:3 123:2,4 150:1 164:24 166:5,8 170:1 177:21 189:6 197:11,21

201:24 **fit** 159:20 **five** 27:14 45:5,10,12 84:7,8 122:19 124:21 141:16 142:22 142:24 180:20 five-and-a-half 188:18 **five-minute** 91:9 188:24 **flat** 178:12 flexible 10:10 **flipping** 53:24 **flips** 54:1 **flows** 21:6 **flu** 87:4,5,6 101:8,13,13 129:17 130:17.20.24 131:3.12 **fluid** 173:2 **fluids** 172:25 **flv** 140:7 focus 60:19 187:14,16 207:17 212:20 **focused** 59:8,21,24 68:21 72:21 214:13 focuses 36:8 **follow** 33:25 50:1,2 109:12 149:22 200:2 204:13 **follow-up** 64:13 followed 149:15,18 155:23 **following** 12:19 26:16 107:14 112:3 135:19 145:24 158:20 **follows** 7:4 204:12 fomites 4:15,17 196:1,2,4,7,11 199:17 203:1 **food** 56:19,20 120:8 footnotes 143:9 **force** 145:7 forces 43:5 174:17,18 206:24,25 207:1,1 208:20 foregoing 220:5 Forest 124:20 **forgive** 171:10 form 61:4 99:12 105:17 108:15 110:4 118:22 124:7 128:7,12 129:21 133:18 138:15 149:1 160:1 164:11 184:3 201:17.20 202:10 215:14 216:22 218:3,25 219:5

**formal** 11:4 24:8,19,20 25:6,12 44:20 86:1 formalities 5:8 formally 117:9.11 format 220:9 formulate 129:15 **formulating** 133:8 216:8 forth 92:20 108:1,24 110:8 133:5 220:5 **Forty-one** 70:18 forward 24:2 Foster's 67:18 **found** 139:10 143:10 foundation 99:4 four 19:21 85:4 96:12,14 122:18 122:21,23 125:19,24 150:11 fourth 24:15 67:14 fraction 55:18 fractures 26:15 fragment 177:3 fragments 177:1,2 frame 152:21 **free** 43:6 **freestanding** 25:18 27:17 French 148:19 frequency 186:6 **front** 143:3,4,11 **full** 8:15 185:5,6 217:2,2,3 **full-length** 198:1,3 fully 203:18 **function** 38:12 55:17 fundamental 170:22 **funded** 36:4 37:4,8,20 78:24 **funding** 74:23 **furniture** 175:10 **further** 22:5 64:12 198:15,22 220:6,9,12,15 G GALLERIA 2:21

garnered 35:20

Gary 27:21 28:7

gatherings 148:15

**GAUTHIER** 2:9

**gather** 139:17

**general** 10:15 35:12 74:5 79:14 79:15 83:24 95:4,22,22 99:19 101:2 131:4 136:8 generally 9:21 27:3 33:3 105:5 113:17 129:16 130:11 133:1 144:15 148:11 189:10.10 generate 135:24 146:14 generic 58:17 59:11 generics 58:17 59:5 **genetic** 141:10 genre 48:4 76:16 gentleman 194:23 198:8 199:15 gentleman's 165:1 **genus** 82:9 geographic 114:13 **getting** 34:16 42:8 44:1 57:8 79:2 102:15 187:6,7 209:7 219:5 ginger.dodd@phelps.com 3:10 give 7:18 8:15 9:23 20:13 30:14 44:20 45:5 61:25 70:2 86:22 89:3 95:19 96:14.25 123:24 156:14 174:15 206:2 215:22 given 7:13 39:24 43:15,21 76:21 76:24 94:22 97:3 121:11 131:2 138:21 146:1 147:6 156:22 162:12 190:8,17 192:9 204:6 215:12 217:2,2,3 gives 121:21 153:5 204:15 **glad** 27:6 137:1 **globe** 200:14 **go** 8:9 13:21 17:25 20:2 29:10,18 34:11 35:3 37:16,16,17 51:17 52:14 53:19 56:12 60:10,22 62:9,12,17 64:12 65:2 70:12,21 72:6 87:19 89:8,10 92:22 98:17 99:13 102:1 103:6,14 108:20 109:9 112:2 115:8 116:18 119:19 132:14,15 136:7,7 143:6,15 145:9 147:19 149:5 150:2 153:9,16,19 160:7 161:5 167:5,16 170:11 173:16 174:2 175:13 177:21 178:14.20 180:1 180:1 184:5 185:6.18 187:2 188:13 189:6 197:5,6,14 198:6 198:15,21 199:2,9 204:4

207:14,14 208:13,13,15 213:11 214:12 **goal** 132:21 **goes** 13:14 37:19 44:7 58:5 153:3 178:20 going 7:16,17,18 8:4 9:3,21 11:21 13:18,23 20:11,13 21:9 21:24 22:5,9 29:15 37:23 45:4 45:16,24 52:14,15 54:2,8 56:6 56:14 59:13 65:25 89:20 91:8 94:7 100:13,14 103:14,24 107:9 108:25 111:16,17 112:21 115:2 119:19 132:10 133:1 137:17 139:14 140:18.18 141:13 142:19 143:7,8 168:9 171:24 173:16 183:19 185:7 188:18 190:20 191:14 193:7 195:6 197:7,8,9 202:24 205:8 207:17 211:4,11,12,12,16 Goldman 4:14 195:6,16 198:9 **Goldman's** 195:9 **good** 6:1 7:7,10,12 11:19 24:18 33:8,8 75:18 83:9,11 88:25 122:12 150:23 180:12 209:9 **goodness** 77:10 **gotten** 148:15 governed 84:15 government 56:1,7,9 71:18 **governmental** 76:9 governor 103:8,16,17,23 114:21 145:2 governor's 104:17 146:21 graduate 74:18 **graduated** 10:3,5 27:8 28:9 62:24 **graduating** 42:10,13 **Grand** 210:13 **grant** 20:20 71:8 121:20,21 granted 109:7 grantees 71:9 **grants** 20:20 71:8 **GRAVIER** 2:16 **gravity** 177:25 great 9:20 31:10 35:20 greater 12:17

greatly 55:8 **Grill** 1:5 96:10 98:18,21 111:21 112:6 116:25 139:11 148:18 150:8.19 151:4.21.23 155:23 156:24 157:6,12 158:2,12 159:3 161:3 172:10 175:9 181:8 182:18 186:10,23 187:21 190:3 192:5,13,23,25 193:5,11 193:13,13 202:17 217:25 Grill's 106:20 189:4 **group** 2:15 16:25 50:22 75:15 76:7 85:1,2 90:21 91:3 126:6 **GROVE** 1:22 guess 25:2 32:7 41:3 43:10 48:17 78:12 125:21 213:22 guest 67:19,23 68:21 70:7 **guidance** 192:9 204:6,15 guidelines 13:10 70:16 155:24 159:5 160:20 220:10 Η **H1N** 131:11 half 94:2 hand 89:10 135:10 198:17 199:8 199:8,11 209:18 211:3 **handful** 83:12 handling 79:17 hang 101:17 **happen** 9:1 95:3 happened 43:23 44:3 99:5

102:21 134:16

198:4

188:6

harm 78:20

**harmful** 166:14

**Haverly** 114:7,8

heads 54:2,9 88:8

**HAYES** 2:20

havfever 69:4

heads-up 45:6

162:19 167:8 218:15

happy 8:2 110:24 169:13,15

hard 47:9 124:21 143:2 185:7

**health** 15:4,5,8 16:15 17:5 24:17

happens 105:3 158:11 162:11,13

25:3 37:12 63:8 71:5 127:10 216:5.6 healthcare 15:14,16,17 16:16 26:7 27:12 38:20 90:25 91:1 hear 107:16 110:23 209:15,15 213:18 **heard** 117:20 hearing 27:2 heart 55:17 58:19 61:20 73:21 74:9.11 heavily 12:7 **height** 183:13 **help** 40:4 48:7 51:18 63:18 64:14 64:17 73:5.13 82:20 88:9.13 142:5 158:6 210:11 214:20 **helpful** 91:25 100:5 104:2 **helping** 20:17 **helps** 142:2 hematologic 40:15 hemorrhagic 41:17 heptillion 213:24 high 33:7,11 49:25 62:25 182:22 186:6 higher 182:5,10,24 183:21 **highly** 172:23 Hill 12:22 13:8,11 66:16 **hired** 82:13 Hispanic 37:12 **hold** 43:7 55:23 92:15 108:14,22 117:25 honestly 83:6 **honesty** 38:14 **honors** 29:16 hope 156:4,6 172:1 179:5 hoped 82:19 hopefully 188:20 **Hopkins** 10:4 41:6 **hormones** 141:25 horses 71:25 **hospital** 10:7 22:17 27:13 28:6 hospitals 46:5 host 74:1.2 **Houghtaling** 2:9,10 92:10,18 93:23 94:4,13,15 96:19 126:6 128:1

hour 122:13 142:20 hours 28:18,19 150:20 151:4,24 151:25 153:2,13 154:6,11,16 154:18.25 155:12 180:24 188:18 189:13,21 199:18 209:9 **Houston** 25:19,20 28:16 87:14 90:13 91:17,21 **HULLEN** 2:11 **human** 99:18 110:13 118:5 177:22 humans 99:18,22 111:11 humidity 131:7 181:20,22,25 182:15 183:6,16,18,20,23 185:10 189:25 **hundred** 83:10 hundreds 142:10 167:14 182:25 hung 98:12 **Hurricane** 86:14 90:14 **HVAC** 171:19 183:22,23 **hvbrid** 174:22 207:23 208:4,6 209:10.19 Hydrogen 206:4 hvdrology 53:6,9 hypertension 16:18 23:12 37:20 38:3 49:3,16,18,20,22 50:8,9 50:17,24 51:2,3,4,12 52:1,7 59:16 65:12 86:19 124:8 hyphen 58:22 hypothesis 40:14 73:7,8,14 165:16 187:11 hypothetical 104:16,24,25 111:7 hypotheticals 109:16

# Ι

i.e 109:23 ibuprofen 125:15 ICOS 123:11 idea 49:12 136:12 171:2,19,22 identification 87:22 identified 13:20 67:1 72:11,17 76:4 112:20 115:10 117:5 135:20 203:3 205:6 identify 6:6 18:22 36:17 75:16 75:20 88:9 134:6,6 200:5,12 identifying 99:2

**II** 2:10 **ill** 90:15 161:16 214:3,5 illness 67:3 89:24 90:2,5,8 145:6 161:11 166:17.25 167:22 214:10 illnesses 87:14 **imagine** 8:9 18:2 149:24 150:4 immediate 63:10 immediately 199:2 **immune** 40:22 **impact** 52:4 89:24 93:3,18 103:2 114:20,21 126:20 **impacted** 172:19 **impacts** 132:1 **import** 12:7 importance 173:7 **important** 20:14 24:20 69:25 103:15 104:23 193:17 **impression** 30:10 96:7 impressive 29:17 **improper** 160:1 194:2 inaccurate 192:12 **inadequate** 189:15 190:9 inanimate 176:18 198:10 199:4,6 199:7 200:7.15 201:5.11 202:4 202:9,17 209:12 213:15,20 214:5 inaudible 121:2 125:8 139:5 inception 48:8 include 15:6 45:1 81:17 83:3 87:4 133:4.24 134:21 163:15 164:6 included 16:13,15 74:6 104:18 includes 15:8,12,13,14 **including** 5:8 15:24,24 20:19 48:24 72:21 73:1 145:15 incomplete 125:4 incorporate 21:8 incorporated 12:23 incorrect 193:2 increasing 88:8 increasingly 146:1 **Indiana** 10:5 27:21 28:10.15 indicate 14:18 120:4 134:5 162:21 177:17 207:18

**indiscernible** 35:1,4 37:15 70:25 83:19 167:24 190:21 individual 89:25 110:1 130:4 154:6.9.11.14 157:18.23 158:13 167:11 169:21 176:15 176:25 182:6.11 198:16 199:1 199:12 200:6,14 202:3 208:7 209:11,23 214:3,5 individual's 158:4 209:18 individuals 12:15 32:11 39:1 49:20 84:2,5,9 85:7,8 87:13 88:22 90:7 96:20 104:19 115:6 150:12 151:13,15 161:2 189:12 212:4 216:11.18 217:15 218:1 **indulge** 182:9 **infant** 67:10 infants 46:4 infect 99:18,22 140:18 176:14,15 177:11 178:3 199:13 211:11,13 211:17 212:6 **infected** 81:21 88:23 89:16,21,22 90:15 104:5,6 156:22 157:5,6,8 157:8,9,18 158:10 160:21 161:16 177:8,14 183:1 198:11 198:20 199:14.18 210:25 **infection** 68:24,25 87:3 88:15 89:1,23,24 90:2,8 100:10 144:16 145:5,6 171:14 176:24 177:22 178:4 194:18 196:3 212:4 infections 68:13 72:21,22,22,25 74:7 75:25 86:25 150:9 infectious 166:24 **infective** 176:16 infectivity 156:23 161:10 166:16 167:22 inference 11:16 12:1,2,14 **influence** 116:19 **influenced** 116:20 146:13 **information** 26:24 33:4 57:19 99:3 113:21 120:18,19 126:8 126:11 134:8,12,18 141:10,21 141:21,25 142:3,3 144:2 156:9 156:18 197:24 **informative** 201:2,8

**informed** 114:2,3 151:14 154:2 202:6 220:10 **ingress** 102:23 inhibitors 82:5 initial 95:12 131:8 138:9,10 216:14 initially 28:9 32:8 149:25 injected 82:17 injecting 85:7 **injuries** 26:16 27:19 **Innovation** 76:17 **inserts** 208:17 inside 97:22 98:17,21 105:23 111:21 172:10 216:18 217:6.15 217:21 **insight** 74:21 insinuates 134:23 insinuation 174:13,20 207:21 208:16 instance 39:2 154:18 200:5 215:4 **instances** 25:5 66:24 67:2 150:11 160:25 198:11 **institute** 71:5,21 73:22 74:9,11 **institutes** 74:2 75:24 institution 78:23 institutions 22:22 **instructed** 163:15,17,18 **instructions** 94:23 97:1,4 insufficient 194:6.7 insurance 97:12,12 98:4 intellectual 21:6 intended 68:22 215:7 intensive 47:16 intent 51:10 140:12 174:15 intentionally 85:7 **intents** 28:24 interact 175:7 205:13 **interaction** 105:3 113:22 interchangeably 108:16 intercranial 124:8 **interest** 220:17 interested 23:24 74:19 193:21 interesting 82:25 **intern** 10:9 **internship** 10:7,8,10,11 27:9

interpret 40:2 **interpretation** 40:3 61:6 164:1 interrogatories 115:5 126:15 **interrogatory** 116:6 156:13,20 interrupt 107:8 interrupted 37:17 intervals 58:8 interviewing 150:15 introduced 92:17 invalidating 189:20 investigator 71:10 investigators 23:21 46:23 **invitation** 31:11 79:19 **invited** 31:6.13 61:9.14.17 70:16 74:25 invoking 105:2 involve 38:17 60:16 67:3 70:8 involved 20:7,8,16 22:4 25:23 38:15 39:19 40:11 47:16 61:19 65:19 70:5 76:10 79:18 87:22 96:10 97:8 122:18 126:1 150:2 involves 15:15 54:23,24,24 125:14 **involving** 54:25 66:16 67:11,21 67:21.24 70:17 72:17 76:5.8 77:8 94:24 123:13 180:19 ion 205:19 ionic 205:10,12,25 206:18 207:2 **ionize** 186:7 **Irina** 53:7 **irises** 14:20 iron 213:2 irregular 179:13 **irrelevant** 183:5 184:2 **isolate** 88:16 isolated 50:24 51:12 52:7 issue 38:20 44:9 123:4,5 188:10 212:16 **issued** 107:11 issues 38:4 67:23 116:22 it'll 51:18 Italy 203:14 204:11,11 item 215:2 IUD 124:6

J **J** 1:17 220:2,24 **jargon** 71:7 **Jennifer** 2:5,10 6:17,18 jennifer@gmhatlaw.com 2:13 jenniferk@akdlalaw.com 2:7 **iob** 28:11,12 179:23 John 2:10 92:18 john@gmhatlaw.com 2:12 **Johns** 10:4 41:6 **Johnson** 125:12,12 joined 25:19,20 journal 29:20,23 30:3,14,18,24 31:4,5,14,16,19,24 32:2,9,20 32:22,24 33:8,12,13,15,16,18 33:21 34:4,5,9,13,19 35:11,13 35:17,21 36:15 50:16 51:2,15 51:16 52:11 62:2 194:20 195:14 journals 20:22 30:4,4,5,8,9,19,24 31:15 32:10,11,16 33:2,17,24 61:1 iudge 107:19 159:19 **judgment** 116:14 **June** 74:18 151:6,6 154:13,15 155:4,5,12,17,21,21 160:24,25 **junior** 71:9,10 justification 168:13

#### K

K71:6,7,9
Katrina 28:25 86:14 88:14 90:14
keep 22:15 56:14 72:21 142:2
143:17
keeping 186:12
KELLY 2:4
kept 189:23
Kevin 3:8 6:14 9:4 13:21 165:21
kevin.welsh@phelps.com 3:11
kick 57:25
kidney 59:2
kids 72:23
kill 42:4 178:17 191:2,9 193:7
204:20 210:24
killed 178:10 179:3,17 191:23

killing 42:1 kills 191:1.7 kind 9:22 22:15 32:8 36:21 44:7 53:4 69:2 104:12 132:25 144:4 172:19 **kinematics** 43:4 139:1 knew 23:23 40:14 48:5 49:1 know 7:13 11:2 21:9 24:1,11 26:17 32:18 34:7,22 38:2 39:11 47:9,13 57:8 67:5,17,18 74:10 76:11 78:12 80:25 82:3 83:5,10 83:20 85:17 86:1 91:23 92:2 96:4 102:16 105:5 107:20 111:22 112:22 116:5 117:18.21 125:1 130:12 133:11 135:7 140:19 147:15,18 149:1,7,14 149:20,24 150:6 151:24 152:7 156:2,17,17,18,19 157:2,3,23 158:7,8 163:22 171:4,21 172:3 172:15 175:9 176:12 177:12 182:7 183:5,6,22,24,25 185:7 185:15 186:15,21,23,25 187:1 187:3,3,7,8,17,19,22 188:6 189:18 193:22 196:12,22 197:21,22 199:23 204:11 206:4 207:6 213:22 214:13,22,22 217:1 218:11,16 **knowledge** 142:8 144:4 159:4 known 48:4 142:15 **Kuechmann** 2:5 6:17

# L

L 2:5 5:1

L-e-m-u-e-19:12 lab 181:12 182:13 183:11 184:1 185:1,12,22 187:16,22,23 188:2 Labarthe 19:19 Laboratories 123:21 laboratory 189:24 labored 202:20 labs 182:3 lack 80:4 175:18 lactation 38:4 39:7 Lancet 4:14,16 34:24,25 35:6,9

35:10 135:13,14,18 195:7,10 195:13,14,17,19 196:10,17 202:23 land 140:8.11 **landing** 140:16 language 14:10 55:3 96:5 163:23 large 23:14 33:10 48:18 121:20 142:12 175:6 **larger** 12:4 43:10 **largest** 142:15 laser 172:23 lastly 125:12 late 209:7 219:5 law 2:15 92:1.2 161:19 162:1 168:5,21 169:17 170:16,21,22 laws 43:5 105:2 141:1,1 lawsuit 96:10 98:3 **lawyer** 126:3 lawyers 92:4 99:7,9,16 100:16 101:20 112:4 115:15 116:3 122:4 126:1 138:17,17 163:16 164:5 lawvers' 193:21 **lav** 140:2 layman's 140:2 layperson 53:22 70:4 lead 68:17 100:23 109:17 174:7 **leading** 203:24 leads 13:1 90:5,5 96:1 109:10 162:8,22 167:24 169:20,25 170:19 learn 24:12 44:16 102:20 103:7,8 103:9,9,15 learned 24:13 46:22 47:22 112:5 114:20 115:5 158:4 **learning** 12:10 50:12 leave 57:22 171:23 183:17 **lecture** 67:23 68:21 69:19,21,23 70:1 76:24 77:3 **lectures** 67:20 70:7,17 71:13 76:20 lecturing 77:7 led 87:25 103:18 **leeway** 43:21 left 45:16 55:17 59:14

**legal** 107:10,18 159:9,20,21 163:3,19,20 164:1 215:11 **Lemuel** 1:13 7:1 9:12 220:4 **lend** 113:13 **let's** 13:17 14:4 29:21 40:6 45:17 50:15 53:24 60:22 62:12 64:3 66:23 68:3 70:12,14,21 72:3,4 72:8 73:4 77:15 84:2 96:14 99:22 100:11 102:24 111:2 112:1 114:2 115:8 120:2 122:1 123:1 126:18 131:22 142:18,23 144:23 147:19 148:8 150:7 156:21 158:1 167:4,16,23 168:1 173:14 174:19 175:16 177:21 180:1,1,14,14 182:25 183:12,13,14 184:20 185:18 188:11,15,16,23 189:6 192:5 192:23 199:16 207:14,14 212:9 213:10 215:4 **lethal** 41:18 68:16,19 **letter** 62:2.3 164:13 **letting** 140:17 leukemia 40:16 level 23:14 103:19 156:24 157:11 157:14 158:25 166:9,10 193:25 194:6 213:19 levels 157:1 166:11 liability 124:4 **liberal** 51:19 licensed 25:20 27:10 28:10 licenses 13:25 **life** 15:15 17:4 27:24 46:10 65:23 140:5 203:25 **lifetime** 173:6 **light** 172:23 186:3,6,11,14,16,17 186:23 187:1,6 188:3 **Lilly** 123:10 linchpins 30:2 **line** 160:12 212:19 **linen** 47:16 lines 135:20 **lip** 14:21 **list** 4:9 30:18 31:15 78:12 109:17 122:17 125:19 173:19 **listed** 30:24 33:3 60:24 122:24

125:19 **listen** 115:21 122:13 **listening** 122:11,11 174:3 **literature** 85:15 102:16 127:12 127:19 128:18,21,22,25 129:2 129:5,14,16 134:20 156:12 187:9 203:10 **litigation** 93:13 94:24 122:20 124:5 **little** 8:9 43:6 55:3 56:10 87:16 112:24 130:19,20,21,22 131:4 165:21 176:11 207:13 live 142:5 176:14 177:11 **LLC** 1:3.4.4 2:4.9.15 92:4 **LLOYD'S** 1:7 **LLP** 2:20 3:4,7 8:23 local 23:20 **locally** 23:12 **located** 1:14 189:13 location 114:13,24 116:25 126:14 161:15 locked 111:2 **lodge** 215:10 **log** 36:22 **logic** 170:18 **London** 1:7 206:25 207:1 208:20 **lone** 207:10 long 32:6 34:7 55:5,21 78:11 93:11 102:25 130:6 141:16 143:14 163:23,24 164:2 184:16 206:13 **longer** 130:19,20,22 131:4 189:20 **look** 14:4 32:4 34:6 39:15 40:6 45:17 50:15 52:18 53:17 60:17 66:23 67:6 68:3 72:4 76:14 78:17 80:15,24 90:10 96:6 114:7 120:2 122:1 130:23 144:14,23 148:8 150:7,23

**looking** 38:8 39:2 67:19 108:3 125:6 188:19 214:11 looks 8:22 150:22 loose 210:17.24 loss 96:1 100:23 101:1,5,14 105:14 106:10 119:22,22,25 128:4 131:10 145:19 146:6,8 146:11,23 147:9 212:13 214:13 214:17,19 215:1,12,18,19 **lot** 18:10 25:11 26:12 43:1.21 51:19 71:20,20 129:5 131:2 183:20 lots 26:24 42:20,24 62:25 loud 172:24 **Louisiana** 1:2,18,23 2:6,11,16,21 3:5,9 5:6 145:3 220:2,13,19 low 4:17 38:4 lower 23:20 lucky 140:14 lung 67:8 71:21 73:21 74:9,11  $\mathbf{M}$ 

**M-13** 1:5 **M-o-y-e** 9:13 **M.D** 1:13 7:1 220:4 **Madam** 160:9 main 134:17 maintained 186:2 maintenance 150:14 major 10:14 11:18 106:8 212:17 **Maker's** 68:7 makeup 175:1,2 making 24:21 97:13 179:19 220:11 **manage** 75:25 **managed** 83:15 **management** 55:7 64:23 manages 58:16 mandate 216:10,17 217:3 **maneuvered** 174:6.16 maneuvers 174:7 189:8 manifest 206:7 manner 82:24 215:7 manufacturer 215:7 manuscript 30:3,15 135:25

152:9 156:21 157:15 165:7

215:22 219:4

114:24 201:13

173:14 182:7 184:20 187:18

188:11 189:2 197:10,12 209:7

**looked** 21:2 61:1,2 83:16 114:19

136:1 137:10 198:1,4 manuscripts 30:4,13 126:13 map 116:24 **marbles** 210:12 March 60:7 103:8 145:11 147:16 mark 13:18 205:4 marrow 40:21 mas@chehardv.com 2:22 mask 216:25 217:6 masks 216:12.19 217:15.19.22 master's 10:20,21 11:6 masters' 19:6 maternal 67:9 math 47:9 49:4 53:9.15.17 mathematical 10:4 11:8,9,11 15:1,3 42:9 47:6 mathematics 36:11 48:5 49:7 54:19 Matt 6:18 matter 35:8 65:11,13 66:2 91:16 220:17 matters 20:14,15 65:16 **MATTHEW** 2:20 Matthews 91:17,18,20,21,22 92:4,7,17 93:9,14,23 94:5,8,13 96:19 126:4 mayor 103:8,15,23 104:17 114:20 145:3,12,21 146:21 147:15 148:12 216:17 mayor's 103:17 147:2 149:15,21 McCormick's 37:11 **MD** 27:2 mean 17:11,14 20:4 21:24 34:10 42:23 66:19 67:13 68:23 69:23 71:15 78:5 85:19,19 87:20 99:22 102:21 106:12 110:6 135:8,16 137:14 138:11 139:12 146:12 152:3,18 153:18 156:1 156:11 158:9 161:20,23 162:15 163:12 168:12,14 169:13,15 174:15,15 176:13,21 177:10,13 184:7 187:20 188:8.20 197:22 200:22 207:6,6 208:10 210:25 211:1 212:23,25 213:1,23 214:25 215:3

meaning 49:19,21,21 53:4 79:23 81:11 131:7 132:1,5 201:4 215:6,20 means 10:12 20:5 29:25 58:23 89:22 95:23 140:3,8 145:15 161:12,13 172:22 173:21 186:7 188:7 195:4 200:23 meant 150:16,21 measurable 199:19 213:13 measured 55:17 measurement 188:7 measurements 14:18 measures 72:20 103:25 mechanism 58:4 207:5.12 mechanisms 207:7,8,8 medical 15:9 22:18,22 24:9,14 24:15,24 28:10,22 30:2 32:22 35:11 42:10,13 44:9 47:1,11,11 47:13 48:4,25 75:7 87:13 108:24 159:10 medication 9:15 medicine 10:6,13,15,15 15:25 27:16,17 29:2 33:9 34:19 35:12 40:25 65:24 76:18,22,23 83:7 118:12 124:14 127:8 136:9,18 medicines 49:19 **MediClinic** 25:14 28:17 meeting 71:6 75:15 121:1 138:10 meetings 20:10,12 77:8,12,13 member 17:2,14,16,17,21 18:9 18:14 19:19 25:9 29:14 56:19 57:1 78:1,2,3,6,7 120:16 216:2 216:5 members 16:13 17:22,23 19:13 19:18 20:24 22:3 83:22 217:21 membrane 178:5 membranes 198:18 199:12 memory 45:25 65:17 124:25 141:13 185:6 mentioned 42:14 126:4 206:21 mere 69:1 merely 13:16 214:12 merits 136:1 **METAIRIE** 2:11,21 method 220:6

**Methodist** 10:7 22:17 27:12,13 methodology 132:21 135:21,23 135:24 136:11,13,14,19,24 137:4.13.15.19.21 139:10.16 165:12 173:12 175:20,21 180:12,13 181:1,11 197:24 199:24 methods 63:3,16 65:7 meticulously 137:22 metric 137:9.15.16 162:16 micro 178:1 microbes 42:4 microbiology 40:12 194:18 microcurrent 209:6 mid-1980s 27:18 milk 39:7 mill 28:4 **Miller** 3:4 4:5 6:13,13 7:6 13:18 13:21,22 42:6 45:7,12,14 91:7 91:13,15 92:21 93:7 99:20 105:20 107:16,23 108:7,20 109:14 110:10,11,14,17,24 111:1 115:8,11 119:9 127:23 128:13,14 130:10 134:1 139:6 142:16,20,23 143:1 147:20 148:1,4,8,10 149:3,6,12,13 152:14,24 153:7,20,23,24 159:13,16,18,23 160:3,7,11,15 163:9,10 164:15 165:20,22 184:19 188:14,20 189:1 191:12 192:4 193:19 194:3 196:9.13 197:10,16 200:10,25 201:18,21 201:22 202:13,22 203:6 205:4 205:7 215:15,25 217:4,7,10,11 218:4,9,13,16,20,23 219:6,7,13 millera@phelps.com 3:6 million 210:12 millions 142:10 167:14 mills 27:20 28:1,8,11 Milt 19:17 mind 51:20 62:10 66:1 70:4 98:24 **Minimal** 163:5 **minimum** 163:4 minute 91:8 109:24

minutes 45:5,10,12 97:7 124:22 142:19,24 188:13 215:22 **minutiae** 184:15 Mirena 124:6.11 **Mischaracterization** 118:24 mischaracterize 120:14 mischaracterizing 108:2 109:1 153:17 191:5,18 216:23 misleading 136:11 179:21 **misled** 136:12 **missed** 53:20 missing 36:1 199:24 **mission** 78:15 mistake 24:22 33:9 mistakes 136:8 mitigates 179:25 190:22 194:14 194:15 modeling 49:3,10,15 50:2 176:8 176:8.9 molecular 173:20 205:18 206:17 **molecule** 82:6 173:25 **molecules** 82:4 142:4 174:8 178:2 186:8 209:3 **Mondelli** 4:16 202:23 money 121:21,23 122:16 monitor 82:13,16 monitoring 77:18,22 78:22 79:10 82:12 83:23 84:3 month 74:17 183:13 months 19:21 117:20 183:16 **motion** 116:13 Motrin 125:15,17 move 21:7,8 24:2 51:18 70:10 160:6 moved 176:16 movement 21:6 140:21 141:3,3 movements 14:21 173:24,24 Move 1:13 4:12 6:21 7:1,7,8,9 115:13 219:8,16 220:4 Move's 115:9 **multiple** 11:2 144:3 184:11 **MURPHY** 2:9 **MURRAY** 2:19 **mute** 147:21,22,22 **mutually** 211:25

Ν N 2:1 3:1 4:1 5:1 naked 213:5,6,14,21 214:7,10 name 6:2 8:15 9:10,12,13 14:23 52:19 68:11 91:4 165:1,2 173:5 175:15 194:23 names 92:5 94:21 nanometers 143:23 144:7.9 **narrow** 32:17 40:2 **nation** 14:13 national 71:5 73:21 74:1,9,11 **nationally** 6:2 23:13 **nature** 44:23 50:7 79:24 197:23 205:20 near 86:12 111:10 200:18 **nearby** 177:18 necessarily 176:13 177:10 necessary 31:11 106:7,9 144:24 144:25 147:3,5 **need** 8:8 19:8 24:5 29:18 31:10 43:16 60:16 64:12 84:12.13 110:23 129:13 133:23 142:4,16 144:5 158:7 167:9 190:7 203:17 204:21 213:18 219:12 **negative** 17:13,13 **neither** 220:15 **network** 37:10 never 17:17,17 55:25 62:10 80:24 95:14 98:21 106:19,23 107:1 117:6.9.11.13 186:24 208:10 211:8 212:7 nevertheless 214:10 **new** 2:6,16 3:5 34:19 57:11,12,23 58:3,3,4,5 87:20 88:7,14,14 103:10,24 106:20 114:16,17 117:1 145:3,4 172:1 183:16 187:2 216:10.16 **newborn** 47:16 Newton's 43:5 nice 7:11 66:19 122:14 Nickerman 19:17 **night** 186:18,19 **NIH** 71:3 120:21 non 77:20

non-lab 187:10,12 **nongovernment** 71:19 77:6 nongovernmental 77:21 78:21 nooks 179:13 normally 190:2 **NORTH** 2:11 nose 198:16 notes 215:23 **notice** 150:17 151:2 **noticed** 46:7 88:6 **notices** 150:17 noticing 6:11 **notified** 154:2,9,13 155:12 **notion** 27:17 184:8 **notwithstanding** 66:2 137:12,12 **novel** 83:5 **November** 1:25 94:2 219:18 220:20 nucleic 141:5 **nucleus** 141:12 **number** 19:9 36:19.21 40:7 46:7 61:22 74:14 88:6 103:20,20 119:5,6 122:17 126:15 130:21 145:1,5,7,24 156:8 163:18 166:2,6 167:6 173:3,19 189:22 196:23.23 number-cruncher 38:13 **NUMBERED** 1:16 **nursing** 15:13 nuts 115:12 0 O 5:1

O 5:1
o'clock 191:13
oath 6:9
object 107:7,9 108:25 129:20
200:7,16 201:6,11 202:4,9
209:12 213:16,20 214:5,6
objected 174:14
objection 6:8,15,20 92:17 99:12
105:17 108:15 110:4,19 118:21
118:21 127:22 128:7,12 129:21
133:18 138:1,4,6,15 149:1
159:22,25 160:1 162:24 163:1
163:8 164:9,11 184:3 191:4,15

non-hospital 199:20

191:17 201:17,20 202:10 215:10,14 216:21,22 217:9 218:3,25 219:5 objections 5:12 objects 139:23 173:19 174:21 202:17 207:22 observation 40:25 observations 172:24 **observed** 180:25 **obtained** 12:5.25 obtaining 11:1 **obvious** 70:5 obviously 92:19 178:17 **occasion** 32:1.3 occasional 35:5 Occasionally 57:14,23 occupational 27:15,16,19 occur 203:25 occurred 48:18 51:14 occurrence 51:13 54:10 occurs 170:21 Oceana 1:4 93:4 94:25 96:10 97:22 98:18,21 102:21,22 103:9,18 104:19 105:10 106:20 111:21 112:6 114:4 115:6.14 116:25 119:5,7 126:16 127:1 129:1 138:20,22 139:11 148:18 149:9,10,14,20 150:8,19 151:4 151:20,23,24 154:3,5,8,13 155:23 156:24 157:6,12 158:2 158:12 159:2 161:2 171:17 172:10 175:9 181:7 182:18,24 183:6,23 184:2,16 186:10,23 187:21 189:4,7 190:3 192:5,13 192:23,25 193:5,11,12,13 202:17 204:21 205:3 216:11 217:16,25 218:15,16 Oceana's 105:12 106:24 114:24 October 37:2 94:5,10,12,17 96:15,17,18 97:25 98:1 99:6,7 100:17,18 105:8 125:2 137:24 137:24.24 offered 118:9,11,11,17 **office** 150:14 **officer** 220:3

Oftentimes 7:24 **oh** 19:15 28:13 36:8 45:19 51:9 57:7 61:16 62:10 64:6 67:5 71:7,15 73:25 74:16 75:3 77:10 77:10 86:11 87:4 89:8 90:19 92:3 94:1 121:7 124:6,11 125:3 136:5 140:4 146:7 150:25 168:8,11,15 189:6 206:9,15 211:6 okay 7:10,16,24 8:8,15 9:5,9,20 10:18 11:6 12:10 13:8,17 14:2 14:4 15:18 16:2,8,12,22,25 17:7,12,25 18:6,20 19:4,16,22 20:23 21:11 22:5,11 23:6 25:1 25:10,13 29:11,15,19 30:17,21 31:5,8,24 33:2,12,18 34:3,7,18 34:23 35:14,23 36:5,14 37:18 39:5 42:12 43:3,8,25 44:15 45:3,9,10,11,19 46:1,6,25 47:25 48:10,21 50:12 51:16 52:5,22 53:1,11,21,23,23,25 54:4,12,18 55:4,6,15,16,22 56:6,22 57:6 58:7 59:10,23 60:3,21 61:17,22,25 62:4,12,21 63:9,12 64:9 65:1,13,15 66:13 66:22 67:19 70:3,21,23 71:2,6 71:17 72:6 73:4,12,21 74:8,24 75:19 76:3,7,14,15,19 77:2 78:10,14 79:2,12,16 80:1,7,14 81:11,16 82:7,12,23 83:4 84:1 84:8 85:15,18 86:8,12 89:19 90:12,24 91:6 92:7 93:17,21,25 94:22 95:25 96:13 97:22 98:10 98:16 99:15,21 100:16,22 101:2,10,17,24 102:7,14 103:3 106:15 107:6,22 110:10 112:1 112:18 113:3,17,20 114:12,15 115:1 116:5,10,23 117:6 119:13,13 120:1,2,15,25 121:4 121:18 122:1,6 123:7,20 124:24 125:20,25 126:7,18 127:2,16 128:13 130:11,18,21 131:5,22 132:13,17 133:7

officials 145:2

138:5 139:7,7,14 140:4 141:20 142:13 143:13 144:8,12,23 147:11,19,23 148:1,5 150:7,25 151:5.10 152:24 153:9 155:1.4 155:7,20 156:4,21 157:7,19,22 157:22 158:5,6,10,14,18 159:2 160:17 161:5,17 164:16,22 165:7,17 166:7,8 168:18 169:10,18 170:5,14,15 171:3,7 171:13 172:4,15,20 173:14 174:25 175:12,24 176:25 177:21 178:11,18 179:16,20 180:13 181:10 182:8 184:20,21 185:18 187:8 188:12 189:3 190:6 193:4 194:10 195:5,16 195:24,25 196:9,16,22 197:15 198:6,7,25 201:1,9 202:14,20 203:7,13 204:18 205:4,15 206:17,21 207:5 208:12,21 210:7,15 211:6,24,25 212:9 213:4,10,19 215:22 217:20 218:18 219:13 **old** 207:7 once 101:17 133:19 135:7,9 158:3 159:4 193:17 200:8 ones 30:6 48:24 70:24 72:11 212:9,12,17 **ongoing** 156:23 onsite 28:7 **open** 26:6 140:12 171:23 172:2 183:18 **opening** 140:16 216:15 openings 145:10 186:12 opine 139:4 202:16 203:18 **opined** 201:14 **opines** 198:8 opinion 95:2 97:6 99:9,16 102:17 123:9,15,23 124:9 138:10 145:22,24 146:9 157:12 163:25 172:13 179:8 192:8 193:14,15 193:20,23 198:9 200:11 opinions 98:2 138:14 193:21 220:14 **Opportunities** 68:8 **opportunity** 34:17 101:18

134:22 135:4,16 136:7,16

102:19 **optimal** 50:3 **optimize** 49:21 oral 172:25 173:2 order 32:7 48:10 139:24 191:22 204:19 209:11 orders 104:18 148:12 149:15,21 149:23 150:2 ordinarily 190:4 organic 113:22 119:8,10 129:6 133:24 139:2 141:2,2 161:24 168:17 169:9 170:23 175:7,7 **organism** 111:10 **organization** 71:3 75:12 121:15 121:20,24,25 organize 24:1 **organized** 90:21 91:3 original 61:3,4 174:23 208:2,2 220:18.18 **Orleans** 1:1 2:6,16 3:5 103:10,24 106:20 114:16.17 117:1 145:3 145:4 172:1 183:16 187:2 216:10,16 **orthopedics** 15:11 26:13 outcome 122:7.8 220:17 outlined 133:1 outside 102:25 111:6,14 112:19 112:21 130:7 187:23 188:1,1 overall 39:12,14,15,20 overlap 206:1 oversee 78:16.25 owner 25:14,22 150:10 owners 111:3 115:15,19,20 P

P 2:1,1 3:1,1 5:1 p.m 1:25 219:17 page 4:2 29:16 36:3 59:13 60:22 62:9,13,13 70:12,21 72:10,10 76:16 146:17 197:6,14 198:6 220:18 pages 220:5 paid 57:1 80:7 92:19 122:4,4 pairs 23:7 207:11 pandemic 131:2 183:14 **panel** 57:17 paper 53:18 135:9 169:22 185:5 185:6 papers 134:7,19,22,24 135:4,13 135:18 paraffin 175:5 paragraph 62:5 122:2,2 126:18 127:6 128:15 131:23 132:11 134:5 136:21 143:17.22 144:14 144:14.23 145:11 147:19 148:9 150:7,8 154:1 156:7,9,21 164:22 165:17,19,23 166:1 167:6 168:2 171:13 172:20 173:14.17 174:19 175:13.14 180:2 184:20 185:18 189:2 207:15,16 paragraphs 62:5 **pardon** 116:15 168:12 PARISH 1:1 part 15:16 20:20 21:4,17 22:17 23:15 29:1 63:17 72:18 74:25 79:9 102:6 103:21,22 115:17 121:14,16 177:6 186:22,25 **partial** 206:18 participants 9:5 participate 16:25 **participated** 10:12 85:8 participating 77:7 **particle** 176:16 205:19 particles 82:6 167:10 179:25 particular 15:18 26:4 43:8 46:15 47:19 49:11 52:10,17 59:7 61:15 68:21 70:15 75:12 82:14 91:3 114:2 123:15 136:9 174:8 207:17 particularly 12:11 93:4,4 148:13 parties 5:4 116:13 220:15 party 107:2 patent 123:5,15 **patents** 123:13 pathogen 41:17 pathogens 41:11 pathophysiological 35:19 pathophysiology 198:24

patients 25:24 26:8,16,19 27:5 27:10,11,14,15,19 28:12,21,23 29:7 40:14,22,24 41:1,9 50:10 73:6 81:21 82:14 83:12.14 86:17 87:8 157:3,5,8,9 patronage 156:24 patrons 150:9 157:5,6,8 167:7 171:20 189:17 192:11 193:6 pay 73:17.18 121:21 paying 121:20,23 pediatric 48:9 **pediatrics** 15:24 45:21 47:17 48:13 peer 30:11 134:7,19,24 135:2,8 135:10,14 195:14,17 peer-review 196:19 **peer-reviewed** 166:20 167:2 168:18 169:10,22 170:25 **peers** 30:13 **Pena** 1:17 6:2 41:14 220:2,24 pending 8:12 9:8 160:14 penetrate 198:18 **people** 32:19 74:19 81:3 89:4,7,8 89:16 92:13 102:23 103:11 104:1.4.5.6 126:16 137:1.16 140:23 161:15 171:10 176:6 183:1,7 184:11 200:24 212:6 percent 63:13,15,21,25 64:18,21 65:4 121:7,10,10 181:22 182:15 183:15,18,24 201:24 percentage 209:25 **Perez** 2:10 6:18 perform 43:9 **performs** 158:15 **period** 145:9 155:22 212:2 periodic 199:16 **periods** 145:18 152:10 persistent 88:7 persists 49:25 **person** 14:19 84:20 89:23 111:3 145:16,16,19 146:4,4,4,16 155:25 164:10 169:4 170:25 198:12 199:5,5,7 201:4,10,12 201:12 202:7 209:20 218:8 220:11

patient 26:18,22 81:18 158:9

**person's** 165:1 199:10 211:3 **physicians** 24:11 25:23 26:10 216:7 personal 198:18 74:14,20 84:14 90:16,19,21 pointed 39:21 66:14 177:14 **perspective** 132:9 146:14,15 127:9 183:21 215:13 physicians' 25:14,22 28:17 policy 26:21 68:7 persuade 136:9 **physicist** 42:23,25 43:7 117:22 political 33:23 **perused** 54:22 **physics** 42:22,24 43:2 95:18 politics 34:1 **pestis** 89:13 102:8,12,13,20 103:21 105:2 polysaccharides 141:24 **Pfizer's** 12:12 106:1,2,6,12,13,16 117:13,16 poor 72:23 **Ph.D** 1:13 3:14 7:1 10:23 11:1 117:19 118:1,2,3,4,15 119:7,10 poorly 86:20 201:18 119:21 129:6 133:24 139:2 **population** 12:5,17 68:7,19 25:2 28:20 74:17 75:9 220:4 89:12 145:15 **Ph.D.s** 19:7 140:8,9 141:1 162:2 168:15,15 **Pharma** 93:13 **portion** 106:7 158:12 173:17 168:16 pharmaceutical 77:12 79:3 physics/organic 113:23 181:10 81:12.22 86:5 pick 183:13.13 portions 106:8 150:19 151:4 Pharmaceuticals 78:24 84:4 picornavirus 69:2 **pose** 14:1 199:18 124:20 picornaviruses 87:1 posed 129:11 PharmacoEconomics 31:3 piece 132:19,19 **position** 16:8 57:2 80:7 98:4 pharmacologic 49:17 **pieces** 94:8 107:17,21 150:16 174:17 pharmacology 49:3 **place** 161:24 178:14 positions 174:6 Pharmacy 85:6 positive 115:7 150:10,12,13 **placed** 213:20 **phase** 91:2 149:18 154:22,22 **places** 169:7 189:16 **plague** 89:11,12,12 155:6 potential 173:7 **phased** 149:19,23 **plaintiff** 6:19 124:2 125:9 pour 140:13 **PLAINTIFFS** 2:3 **phases** 152:10,11 153:6,11 power 42:1 **Phelps** 3:4.7 8:23 141:14 **POYDRAS** 2:5 **plaque** 89:13 **Phoenix** 149:25 **plastic** 180:20 practical 12:14 **phone** 92:9,12,13 96:20 98:7,9 play 81:9 129:8 **practice** 28:22 29:2 101:17 105:9 **played** 76:9 practicing 18:15 **phrase** 69:11 80:4 126:19 146:5 **Pradaxa** 125:2.3.7 plays 118:13 207:17 214:18 215:17,19 please 8:2,9,16 9:24 22:25 34:11 **PRAIRIEVILLE** 1:23 **phrased** 201:19 37:16 60:23 62:9 71:24 86:11 pre-COVID-19 189:8 **physical** 38:25 95:5,7 96:1 99:8 102:3 103:6 109:4 172:8 pre-stage 168:9 100:22,25 101:5,14,19 102:6 191:18 197:13,15 208:14 210:8 pre-staging 198:3 102:10 106:23 128:3 131:6,9 precautions 204:2,8 213:17 131:10,17,18 161:19 162:1,8 pleased 23:15 predicate 166:8 162:22 166:3 168:5,20 169:16 **plus** 60:11 86:18 **predicated** 106:3 161:19 168:4 **pneumonia** 41:17,19 169:20,25 170:16,19,20,21,22 predicted 161:21,22,22 168:20 174:17,24 212:13 213:4,15,21 **pneumonias** 72:23,24 predictions 52:23 214:6 215:12 pneumonic 89:11 **prefer** 63:10 **physically** 107:3 161:9 166:15 point 9:3 21:4,9 24:11 29:5,8,13 preference 26:23 166:22 168:24 169:1,5 31:9 41:4 44:8 57:8 64:10 70:8 preliminaries 7:17 physician 25:19,20 28:4 34:21 84:11 86:6 88:16 98:9.13 105:2 preliminary 9:14 prematurity 16:19 41:8 75:4,4,5 79:21 85:24 137:10 146:3,11 150:23 166:18 86:16 89:19 91:1,1 169:4,7,8,8,21 170:24 178:15 **premier** 35:12 physician's 89:20 184:17 192:11 200:24 211:7 prenatal 67:10

preparation 200:4 **prepared** 220:7,9 **preparing** 114:18 115:25 116:3 126:7 prescribed 49:20 prescription 9:15 **presence** 86:20 114:3 157:24 158:4 161:13 177:13,16 **present** 3:12 23:1 104:20 105:10 105:12 106:4 107:4,12 108:11 110:2,3 111:5 154:10 158:13 161:2 171:20 182:17 186:17,18 186:18,19 201:15 **presented** 105:10 203:20 press 194:22 195:3 pressure 49:25 presumably 98:13 **presume** 36:12 preterm 39:6 pretty 48:25 prevalent 146:1 prevent 9:17 51:13,13 preventing 82:18 **prevention** 50:23 51:4 prevents 204:9 previous 166:9,12 197:14 **previously** 44:19 126:1 **primarily** 27:15,16 36:10 40:15 52:20 59:1,3 114:12 164:23 **primary** 18:8 25:25 71:13 75:11 78:14 **principally** 64:8 162:4 principles 181:4,6 **prior** 8:13 17:25 93:8,22 94:9 125:23 199:8 **prism** 136:14 **private** 78:22 120:20 **probabilities** 53:15 54:20 **probability** 11:13,14,15,25 36:11 53:10,17 54:13 104:9,9,11 **probably** 92:5 142:14 176:15 198:22 209:8 **problem** 26:8,15 43:15 68:14 145:4 152:16,17 164:3,20 214:9,9

procedural 66:3,5 **Procedure** 5:7 220:14 **procedures** 204:2,8,16 proceed 73:3 proceeding 219:17 **process** 12:3 13:13 30:2 35:19 44:23 50:6 79:23 109:10 116:8 132:18 135:20 138:7 148:4 161:19,21 168:4,20 173:19 189:20 205:12.16 processes 161:25 **proclamation** 145:12,21,23 147:2,16 **produced** 170:16 **produces** 183:23,23 producing 82:18 **products** 124:4 175:6,7 profession 13:24 professional 10:2 14:4 16:3 33:4 220:3 **professor** 14:6 22:8 **program** 14:12 28:20 37:11,11 38:16 50:20 72:4,18 73:5 74:3 75:9,18 83:6 **programs** 14:13 19:1 37:20 prohibition 220:13 **project** 48:8,8 projections 178:2 prolonged 145:17 **promise** 60:22 111:18 **prong** 182:8 **prongs** 51:20 **pronounce** 34:24 173:5 180:6 **proof** 163:5 **propensity** 145:16,17 146:1 **property** 100:20 101:15,20 105:14 106:10,12 145:19 146:6 146:8,23 147:9 150:12,18 151:3,12,16,18,20,25 153:1 172:7 189:9,10 214:13,17,20 215:1,17,19 proprietary 81:1 prospective 150:15

**proud** 52:8 **provide** 39:25 74:4,21 144:5 **provided** 5:11 46:4 57:19 113:21 116:5.10.12.24 117:3 126:8 173:12 204:1 **provides** 51:10.10 136:2 165:15 **providing** 144:18,19,21 **prowess** 74:4 76:1,2 **pseudo** 124:7 **pseudomona** 41:11,16 pseudotumor 124:7,16 **public** 16:15 17:4 24:17 25:3 26:7 63:7 103:25 110:6 127:10 216:5.6 **publication** 24:1 30:7,16 39:3,7 39:21 40:7,11 42:15 43:8,14,15 44:10,13 46:16 47:3,15,20,23 48:12,18 49:11 52:10 59:24 60:1 194:18 195:17 196:18 **publications** 16:1 38:9,12 39:3 39:20 45:17 46:8 48:3,19,20 52:19 54:14 59:15,18,19,21 60:2 **publicly** 195:1,4 **publish** 20:21 30:5,8 32:1 33:5,9 34:4 36:7 publishable 24:3 **published** 13:12 30:9 32:5,16,21 41:4 43:23 44:2 45:18 51:14 56:16 60:6,7,8,15 61:5,8,23,25 62:14.16 134:7 156:12 publisher 36:8 publishes 195:19 **pull** 13:17 115:9 196:9 202:22 pulmonology 15:11 punctuated 54:14 **Purdue** 10:19,20 purpose 71:13 75:11,14,15 144:16 purposeful 140:22 **purposes** 5:5,6 28:24 99:25 212:13 219:1 put 21:20,21 22:2,3 23:4 31:16 36:20,24 38:5,23,24 39:12 43:12,12 51:20 75:14 88:8 92:9

**protect** 104:1

**proteins** 141:24

110:18 147:3 150:24 164:19 radiation 186:6 145:13 146:21 186:22,25 188:8 213:24 raised 220:18 190:10,15 192:6 197:17,19 putting 38:19,19,25 73:23 131:5 reasonable 13:13 194:1 **rampant** 86:18 131:5 ran 136:20 reasoned 124:17 randomized 50:23 **reasoning** 72:16 147:12 0 reasons 36:19 119:18 125:21 range 144:11,13 qualifying 20:13 **ranges** 144:6 190:8 quality 15:15 33:7,11 65:23 **rapidly** 89:15 recall 61:14 79:12,15 84:2 92:5 **Ouarter** 148:19 rare 68:15,15 136:1,1 123:23 156:11 185:17 question 8:12,13,17,18 9:8,10,14 rate 14:19.19 38:4 **receive** 121:19 15:22 18:23 27:7 31:17,18 rates 180:17 received 79:19 34:11 35:3 57:15 64:1,11,13 rationale 50:22 98:3 Recess 45:13 91:12 142:25 66:18 70:1 74:16 82:8 88:25 raw 176:21,21,23 177:14,16,17 188:25 215:24 104:24 105:18 109:4 110:17,22 **reach** 179:15 190:13,15 192:3,17 **RECILE** 2:20 112:17 115:17 121:4 124:12 192:18,19 **recognize** 123:2 206:3 129:3,11,22 138:16,17,19,21 reached 138:8,11 192:25 **recollection** 77:6 80:1 98:11 152:18 153:8,22,23 158:7 reacted 212:3 recommend 204:18 159:8,8 160:2,4,8 163:2,8,11 reaction 207:5,7,12 recommendations 190:16 166:6 172:8 173:9 177:12 read 70:4 113:4,9,13 115:4 120:5 194:10 184:4 191:16,21 193:12 194:2 129:4,14,15 131:24 132:10,14 recommended 158:3,15 194:4 197:2,9,19 200:1,19 132:15 134:23,24 135:2 150:8 record 6:7 9:11 91:14 110:19 201:23 211:16 212:5 215:2,14 151:1 156:22 160:13 168:2 160:13 190:11 218:7 217:13 218:5,17,20 172:21 177:24 189:7 195:23 Recurrent 55:20 questioning 160:12 197:3 198:4 199:22 203:5,7,17 reduce 51:12 122:9 questions 7:18,24 9:18 14:1,1 203:17.22 reduction 180:24 24:6 95:2 96:23 159:24 164:17 reading 5:8 50:19 152:14 **ref** 166:5 191:14,15 202:21 205:9 216:2 readings 200:4 refer 195:6 203:15 **quibble** 127:16 reads 145:13 150:8,25 164:24 **reference** 143:9 165:18 166:1,6 quick 7:16 76:14 205:8 ready 45:8 167:9 175:2 quicker 51:19 **reagent** 206:11 **referenced** 113:2,6 114:9 127:3 quickly 15:22 76:15 213:12 real 23:21.22 76:15 188:7 203:25 127:3 quite 11:21 21:4 62:22 83:6 206:8 references 115:4 143:16,21 185:24 207:16 **reality** 178:19 145:11 165:8 quote 120:5 133:3 148:15,16 realize 24:6 reflect 122:3 197:23 **quoted** 145:12 really 10:21,24 14:9 23:4 24:1 reflecting 134:8 quotes 131:24 132:4 27:17,22,24 28:13 37:18 40:1 reflective 61:3 43:21 46:24 48:9 53:17 62:2,6 refresh 45:24 120:13 R 71:8 87:5 88:11,25 89:7 118:13 refreshed 124:25 R 2:1 3:1 129:13 130:13 144:22 146:9,12 **regarding** 59:20 96:9 100:18 **R-o** 88:3 154:22 187:6,20 192:6 197:25 112:11 **r-o-t-a** 88:3 198:1 201:18 206:12 207:20 Regardless 39:18 **r-o-t-o** 88:3 208:8,9 211:22 215:20 219:11 regenerative 76:18,22,23 **R.S** 220:4 Registered 220:2 **realm** 102:8.9 **R0** 71:6 reask 191:19 regulations 120:19 **R01** 71:7,10

**reason** 36:16 78:18 125:18 143:6

rehash 22:6

rejected 135:10,12 rejecting 134:15 **related** 46:16 64:22 67:22 71:22 121:6 122:8 129:16 134:14 220:16 **relation** 39:2 45:22 129:1 **relationship** 13:2 44:24 67:9 80:20 124:1,13 125:8,14 132:22 134:9 136:3 relationships 220:13 relatively 26:13 32:25 89:6 148:19 released 209:6 211:8 relevant 176:9 184:14 213:8 reliable 33:4,7,11 175:18,20,21 reliance 184:1 relied 112:12,15,22 127:4,7 128:9,16 134:13 175:14 184:22 **relies** 133:21 rely 30:7 36:2 111:23 129:5 133:22 139:8 180:4,9 185:19 remain 210:16 remainder 143:7 remained 180:22 184:13 **remediation** 152:3 189:5 remember 41:24 56:13 59:1,9 65:20 73:18 80:6,19 81:1 82:11 84:6,7 85:9,11 90:11,11 94:21 115:23 123:12,16 124:22 125:23 126:17 135:1,2 152:5 156:10 181:24 185:4 207:7 208:24 remembered 82:9 **reminded** 183:21 remote 6:10 remove 9:4 renal 58:22 render 123:10,15 124:10 184:1 rendered 26:5 reopened 154:19,21 reopening 149:19 154:23,23 216:15 repaired 215:1 repeat 15:2 55:12 66:15 110:24 repeatedly 24:22

repeating 37:25 rephrase 8:2 replicated 43:20 report 4:12 8:10 97:20,21 98:6,8 98:13,17,24 101:21 102:14 104:2 106:1,6,7,9 107:12,20 108:2,10 109:5,7,8,16 111:7,15 111:16,18,19,24 112:5,13,16 112:16,21 113:1,6,6,10,13,14 114:10,19 115:9,13,25 116:3,7 116:9,19,21,25 118:18,19 119:3,19 120:3,19 126:7,20,23 127:5,17 128:16 130:12 131:23 133:9.12 134:13 135:6 138:9 138:12,14 139:9 143:3,7,16,20 143:23 144:17,24 145:13 146:17 147:4 152:2,5,15,24 155:10 156:15 161:1 162:20 163:16 164:13 168:14 169:12 169:22 170:7 171:1 178:19 180:10,11 181:10 182:2 184:22 184:25 185:1,3,16,20 186:13 192:24 193:11 195:7,9,16 196:10 198:2 200:4 206:22 207:15 214:12.18 215:17 **reported** 1:16,21 120:7 150:12 189:12 220:6 **reporter** 1:17 6:1,3,8,21 7:19 41:14 55:12 160:9,10,13 202:24 220:2,3 **REPORTER'S** 220:1 Reporters 6:4 **reporting** 121:6 220:6 represent 6:6 42:25 124:2 representative 116:2 representatives 115:14 represented 125:9 reprimanded 149:21 reproduce 140:19 **reproduces** 141:17,18 reproduction 141:6 request 35:5 71:9 require 18:21 required 40:16,18 216:10,18 220:10

requirement 163:3 requirements 18:20 requires 216:11 rereleased 210:21 research 12:20 15:4,5,8,9,10,10 15:11,13,13,15,17,22 29:7,10 30:2 36:4 37:5,8,12,19 39:12 39:13,14,15 41:5 48:1 49:12 50:20,22 61:3 74:2,4 75:17,21 76:1 83:18 85:1 102:15 120:3 127:11 128:17 130:13 133:21 134:17 135:22,22,24 136:8 165:5 180:18 189:17 202:16 researched 201:10 researchers 30:13 reserved 5:12 resources 126:12 respect 12:11 38:4 44:12 49:15 49:17 50:16 75:21 79:6 93:22 94:24 95:14 144:17 146:11 173:15 181:7 187:9 193:11 212:4 **respiratory** 86:24 87:2 responding 61:12 responds 61:8 response 40:22 41:2 95:10,11 152:19 responses 116:6 responsibility 63:13,15,25 83:14 121:2 responsible 52:20 63:21,23 64:18 rest 139:24 146:19 restaurant 93:5,6 97:11,23 103:12 104:7,10,12,19,23 105:11,12,16 106:21,24 107:4 109:9,25 110:12,16 111:2,4,11 111:12 115:15,20 119:16 139:11 155:11 157:1,13 158:23 158:24 159:4 161:3,9,14,15,18 161:24 166:3,15,21 167:7 168:4,24 169:2 172:16 175:25 176:1,9 178:23 183:1 192:7,12 201:14 204:21 216:11 217:16 217:22 218:2 219:10

restaurant's 161:5 166:13.23 169:6 172:5 restaurants 93:4,19,22 94:9 95:4 95:15 145:8 192:10 restore 180:3 189:8 restriction 217:3 restrictions 148:13 152:22 153:14 result 24:2 40:1 96:24 135:23 151:19 198:25 resulting 105:14 212:14 results 16:21 88:20 131:9 136:10 136:11,15 137:17 176:5 197:19 197:21 resumption 154:24 retained 92:19 **retinal** 16:19 retired 23:4 31:16,21 revealed 172:24 **reveals** 189:19 reverse 147:5,6 172:18 reversed 8:20 review 30:2,11,13 32:3,12 34:8 34:17 35:5 36:6,6 73:1 74:3,15 102:20.20 123:14 127:12 128:18,21,21 136:15 reviewed 32:20 34:12,14 36:18 37:5 59:2 73:25 112:8 115:3 127:18,19 128:24 129:1 133:5 133:12 134:7,19,24 135:2,8,10 135:14 195:8,11,12,13 reviewer 29:23 30:18,25 31:4,5 31:14,20 32:9,24 33:13 34:8 35:24 reviewers 31:10 33:1 36:15 135:17 reviewing 34:1 reviews 31:17 57:19 195:14,18 rhinitis 68:12.12 rhinovirus 69:1 rhinoviruses 86:25 **Rick** 19:17.20 **Rico** 2:4 6:16 92:10,11 94:18 110:18 126:6

rico@akdlalaw.com 2:7

**rid** 160:20 **Riddle** 185:19 186:1,13 189:18 189:18 **right** 7:14 8:21.24 9:3 11:17 14:24 22:25 23:3 29:15 32:15 32:17 33:20 40:6,8 42:19 45:7 45:15 50:12,15,19,25 51:1,9 52:9,9,14 53:2,23 55:4 56:18 60:2,7,22 65:14 66:18 68:1,9 68:20 69:15 70:12 72:4,9,14,15 73:7,16 75:13 77:24 78:4,8 80:14 81:7,24,25 82:19 84:16 88:5 90:4 91:13 92:5,21 94:19 96:8.9.22 97:3.18 98:16.23 100:2,11 106:11,17 107:23 108:4 109:13,15 113:25 114:17 115:8 116:16 120:17,22 122:1 123:17 125:1,25 127:20 131:16 131:16,22 132:4,7,12,24 133:14 134:4,5 135:15,19 136:4.21 139:14.20 141:22 142:23 143:20,22 148:5,19 149:4,12,17 151:8 152:5 153:3 153:8,20 154:8 155:3,4 156:7 156:14 160:3.7 161:23.23 162:18 164:5 165:9 168:1,6,20 170:6 172:17 174:4,4,4 178:13 178:18 179:7 180:1,2,14 181:14,23 182:3,6,12 183:8,14 183:25 184:23 185:4,10,23 186:2 188:23 195:25 196:4 199:4 200:23 201:25 203:21 204:23 205:8 207:3,14,25 209:15 211:9,17 212:11,21 214:2,24 215:5 216:1,8 217:7 218:22 219:6,13,15 risk 4:14,17 171:13 196:11 199:19 202:25 **RNA** 140:13,17 141:4 142:2,7,12 142:14,15 176:12,13,21,21,23 177:6,10,14,16,17 **ROAD** 1:22 **RODERICK 2:4 role** 39:13,18 43:13 46:23 47:20 65:6 66:25 72:25 73:15 76:5,23

76:24 77:3,24 81:8 82:17 94:23 102:4 118:14 146:13 **roles** 76:9 room 10:16 137:2 189:21 rooms 189:12 rose 157:1,13 166:11 **rotating** 10:9,11 rotavirus 87:25 88:3,4,17,21 89:17 90:8.14 Rouge 3:9 6:3 220:19 rough 209:1,1,2 row 54:9 **RPR** 220:24,24 rules 220:10.14 run 54:5,9 55:5,21 **running** 209:3 runs 52:22 53:3,3,20 54:7,8 rush 134:17 rust 213:1,3 S **S** 2:1 3:1 5:1 **S1S2** 178:1 safe 150:18 151:3,13,19 153:1 192:11 193:6 **Safeguarding** 68:6 safely 215:21 safety 77:18,22 78:22 79:10 82:12 83:23 103:25 120:18,19 161:10 166:16,24 167:21 sake 158:1 salary 121:14,16,22,22 sample 12:3,5 70:16 **SARS** 100:7,8,9,9,19 **SARS-2** 96:24

**SARS-CoV-1** 180:19

204:10

136:3

**SARS-CoV-2** 4:17 69:14 95:8,14

99:17,23 104:6 128:25 129:4

131:25 132:22 134:9,14,16

177:25 180:19,22 203:24

SARS-CoV-2/environment

**SARS-COVID** 96:23

143:23 144:13 146:21 173:8

**SARS-COVID-1** 180:16,25 seasonal 68:13 separately 101:3 **separating** 174:23 208:1,2 **SARS-COVID-2** 150:9,13 second 24:14 33:12 45:17 50:6 **September** 87:5 202:23 156:23 180:16 189:16 51:21 55:23 59:4 72:8 92:15 satisfactorily 29:9 97:9 101:3 103:5 107:9 111:17 sequence 49:22 50:3.3 53:4 98:6 saw 27:14,18,22 61:22 135:2,7,9 127:6,6 139:9 143:13 167:5 98:11 158:6,8 192:16 198:25 saving 18:13 63:10 105:25 172:21 173:2 182:9 198:6 series 7:18 48:3 202:21 209:20 118:16 139:15 147:6 153:6,25 209:16 serious 145:2 162:16 163:7 172:18 179:11,12 second-to-last 207:20 seriously 21:19 191:6,6,8,10 192:2,15,22 secondary 17:7,8 20:3,4,25 seriousness 145:1 218:14 21:12.22 63:18 serve 77:21 97:14 says 14:5 22:11 23:1 25:13 49:2 Secondly 146:2 served 18:7 66:24 84:3 85:22 50:21 78:21 107:20 127:7 section 29:18,22 37:4 56:13,15 122:20 143:23 152:25 153:15 167:9 62:17,21 113:24 155:11 service 56:7 71:18,19 73:16 76:9 sections 158:12 169:5.23 173:17 177:24 179:9 77:5.21 179:16 189:6 192:24 199:15 see 7:11,12 9:4 25:13 26:19 services 220:12 27:12 32:4 34:6 41:8,9,18 46:9 serving 85:22 86:1 203:23 scale 210:12 51:16 54:9 55:22 56:7 67:23 set 78:7,9 220:5 **scary** 83:7 70:3,14 71:22 72:3 81:3,5 **setting** 27:1 40:23 scattering 172:23 83:14 90:10 100:13 118:13 **settings** 199:20 scenario 105:9 108:8 109:23 **settle** 210:9 120:10 123:1 127:14 133:1 111:6 160:24 178:22 212:2 142:18 146:25 173:4 178:7 **settling** 210:10 199:16 204:3 212:3 213:8,10 seven 25:24 26:3,4 83:25 84:1 scenarios 109:17,22 **scheme** 43:10 214:9 Seventeen 68:3 school 10:6 14:12 17:4,21 21:3 seeing 8:22 25:23 26:19 27:4,6 Seventy-four 77:11 22:20 24:10.14.15.17.24 25:3 27:10,11 28:12,20,23 29:7 sex 38:5 55:7.18 127:8 **shape** 61:4 28:10 42:11,13 47:1,11,11,13 48:5 62:25 63:7 75:7 seen 12:7 27:23 61:5 107:1,5 **share** 26:12 205:17,24 schooling 11:4 171:3,4 182:24 196:20 **sharing** 9:1 26:20 science 11:11 14:22 15:4,5,8 **Seldane** 68:6,12 Shekelle 19:17 37:13 42:9 47:9 49:4 52:8,12 selection 55:19 **SHEP** 50:19,21 55:19 108:23 146:10 163:21 206:23 **self-evident** 166:18,19 169:16 **Sherman** 2:19.20 6:18 sciences 10:4 47:7 self-explanatory 35:8 shifts 28:7 scientific 99:4 147:9,16 163:24 self-propelled 139:22 **short** 22:16 62:6 142:2 186:9 seminars 25:8 77:7 165:8 175:1,2 187:11 short-circuit 32:8 **scientifically** 99:24 146:23 senior 19:18,19 23:11,11,14 shorten 57:9 scientist 84:19 46:23 71:11 **shorter** 61:11 scientists 84:15 sense 136:25 213:18 shorthand 100:5 scope 60:13 sensitive 41:20 83:17 172:23 **shortly** 27:10 **SCOR** 73:22 sentence 127:2,6 128:20 131:23 **show** 9:2 41:25 75:24 191:1 showing 74:4 screen 8:20,20,23 9:2,4 143:17 131:25 148:12 150:24 153:10 screenshot 8:22 153:15 164:24 166:9 168:1 **shut** 110:6,6 scroll 36:1,2,3 55:3 56:11 71:24 169:18 172:21 177:24 189:5.6 **shutdown** 104:18 109:25 110:5 72:5.10 76:15 86:11 165:20 203:22 207:18.20 208:16 149:18.23 155:2.5 216:16 **seal** 220:18 sentences 177:22 shutting 150:4 **season** 87:5 **separate** 33:17 58:7 sic 138:20

sickle 72:3,12,19 73:6,6,10,11 75:13,25 76:5,8 77:13 sickness 99:24 110:2 side 34:3 66:3.11 69:24 sifted 132:19 signature 220:18 signing 5:9 **signs** 144:18,19 **siloed** 21:4.7 similar 100:19.19 104:16 109:21 180:24 184:25 185:22 Similarly 111:19 **simple** 26:14,14 53:24 simplified 65:25 **simply** 18:13,25 19:1 23:23 33:18 46:17,17 87:20 89:22 140:24,25 163:17 178:6 179:21 190:13,17 192:1,15 **simulations** 176:5,7,7 **single** 67:23 164:13 sir 7:12,15 8:19,24 9:16,19 11:5 12:9 13:7,11 14:3 16:11 17:10 18:5,7 19:3 22:10 25:4,16 26:2 28:2 29:24 30:20 31:7,23 32:13 33:23 34:20 35:16 42:17 44:11 45:2 47:2,4 49:9,17 52:13,21 56:3 58:9 59:12 64:20 75:23 79:4,8,11,25 80:13 81:14 82:22 94:6,16 95:24 96:21 97:24 98:19 100:21 101:7,9,16 102:13 112:7,10 113:11 114:23 114:25 116:4 119:17,20 120:11 123:19 125:14 131:15 132:16 144:10 148:21,23 160:10,16 162:4 164:8 165:6 180:5,7 181:19 185:21 204:25 214:8,15 sit 31:1 98:20 126:17 site 28:4 73:22 sites 74:22 sitting 140:15 156:2 211:5 216:24 **situation** 26:7 171:24 situations 27:23 **six** 54:8 84:7,9 122:19 209:9 **size** 70:16 131:19 144:6,9,11,13

174:5 179:24 190:22 194:15 sizes 131:20 **SJS** 125:17 skill 78:6.8 **skills** 79:22 **skip** 22:9 29:15,18 139:14,20 177:23 **slowly** 89:6 small 32:25 173:6,23 198:11 213:9 smaller 85:11 smallest 214:4 **smoking** 59:17 67:10 **smooth** 178:12 208:25.25 **SN1** 207:8 **SN2** 207:8 **sneeze** 157:8 167:11,13,14 182:11 198:14 sneezed 182:6 sneezes 167:8,13 198:12 sneezing 183:2 **solely** 38:12 60:19 solid 127:1 136:19 139:23 173:18 174:21 207:22 **solution** 206:12 **solving** 54:12 **somebody** 32:20,21 40:2 43:14 104:22 122:12 167:8,13 169:23 209:2 213:23 **someplace** 36:25 140:9 soon 198:13 sophisticated 88:12 **sorry** 9:9 12:21 19:23 22:25 34:10 35:2 37:16 42:12,12 47:4 51:6 55:13,19 58:21 71:7 80:16 86:4 88:19 89:8 92:2 102:1 103:5 105:11 115:16 124:23 125:3,4 131:11 137:19 147:25 148:6 151:11 161:22 170:11 174:12 175:13 178:20,21 179:3 181:11 190:7 195:3 197:2 203:14 204:4 sort 27:2 **sound** 139:10,18 165:13 173:12 198:24

**sounded** 38:8 58:18 sounds 66:19 195:22 **source** 173:3,10,11 174:23 sources 33:4 144:3 168:19 **South** 1:14 7:2 spaced 89:5 speak 43:11 115:13,18 116:2 157:9 speaking 46:23 130:11 182:11 special 26:11 specialist 16:18,18 84:13 183:22 specialists 16:16 specialized 15:19 specialty 26:4 35:17 species 79:15 82:10 130:2 142:11 144:10 **specific** 18:20 36:17 43:9 75:15 79:15 118:8 125:18 129:14 134:22 214:16 **specifically** 64:22 71:22 95:4,6 102:18 121:5 126:24 134:25 150:10 **speech** 172:24 173:7 spell 41:13,15 206:14 **spend** 11:21 52:16 120:23 149:6 **spew** 157:9 **spewing** 184:11 **sphere** 79:1 **spoke** 58:21 **spoken** 94:7,8,14 115:19,24 sponsor 120:7,7,20 **sprains** 26:14 **spread** 89:13 104:12 145:14,16 146:2,3 164:23,23 165:16 172:22 173:13 196:2,7 **spreading** 90:5 145:18 **spreads** 89:6,11,15 **spring** 196:25 **Springer** 35:25 36:6,7,8,9,16 stability 180:15 Stadnytskyi 173:5 staff 110:7 216:17 217:21 stage 40:23 154:23 staged-to-phase 145:10

stainless 180:21 STAKELUM 2:20 **stand** 170:12 standard 73:9 107:10.18 132:20 159:9,20,21 163:5,19,21 200:9 204:1,8,16,20,22,24 standardized 188:2 start 6:11 9:25 10:1 22:25 started 14:11 17:20 27:6 98:5.12 98:14,17 112:4,4 127:17 starting 40:7 **state** 1:2,18 9:10 49:25 134:9 139:21 176:22 187:8 197:23 220:2 **stated** 148:12 statement 55:24 103:18 151:18 164:6 166:21 168:19,21 **statements** 137:3 166:2 states 148:12 176:1 statins 37:21 82:4 **statistical** 11:16 12:1,2,14 33:13 63:1.9 **statistics** 11:8,10 15:2,4 36:11 48:6 49:1 53:15,16,18 108:23 **status** 67:9 **statute** 220:10 stay 168:1 178:15 210:13 stays 178:6 steel 27:20 28:1,8,11 180:21 steering 84:22 85:3 stem 37:10 Stenotype 220:6 step 17:5 34:10 198:15 **stepped** 35:2 steps 198:22,23 202:2 Stevens-Johnson 125:15 stick 64:3 108:5 stint 59:4 **STIPULATED** 5:3 **STIPULATION** 4:3 **STOCK** 3:14 147:23 **stood** 67:11 **stop** 19:23 26:10 31:19 72:8 76:13 77:19 78:19,20 83:16 103:5

**stopped** 80:14 stops 210:19 straight 53:9 Street 2:5,11,16 3:5,9 105:24 106:19 148:14,20 strength 12:24,24 stricken 160:4 **stroke** 37:7,10 38:1,3 50:23 51:2 51:4,14 52:1,4 54:23 55:6,16 59:17 stronger 139:12 structure 91:4 161:18 162:6 168:3 169:19,23 170:7,16 **student** 24:16 25:4.6 28:19.20 43:22 48:25 66:23 117:19 students 20:17,18 66:25 117:10 117:16 **studied** 118:2 studies 64:7 93:21 95:14 102:24 112:8,11,14,15,19,20 113:1,4,8 113:9,12,17,21 114:1,19 187:18 200:3 203:15 **study** 12:23 41:7,8,9 42:16 43:18 43:18,19 46:12 51:10 73:2 78:19.20.20.24 107:1 114:1.5.9 120:20 135:21 136:10 165:10 175:13,22 176:4,5 181:12 183:12 184:1 200:21 stuff 50:12 **subject** 35:8 65:11,13,16 66:2 158:9 subjects 26:8 **submitted** 30:3,4,23 31:2 32:11 32:19 56:15 113:15 **subset** 12:15 **substance** 131:6,18 substantial 24:9 27:8 203:10 successfully 74:5 successive 54:9.10 **sufficient** 75:17 78:18 160:20 198:17 199:11 200:18 **suggest** 188:12 193:9 203:23 suggested 88:17 suggests 140:12

summary 116:13,14 summer 183:14,16 supervised 120:6 **supervision** 67:3 220:7 supervisions 66:23 supervisory 66:25 supplemental 46:3 **support** 103:19 121:22 133:16 139:17 143:10 147:10 168:19 181:4 **supported** 135:11 146:24 **supports** 102:16 165:16 166:20 sure 9:12,25 10:3,21 11:11 13:10 15:1 16:12 17:2 19:15 21:23 23:10 27:6 31:2 33:25 37:11,23 38:18,23 44:18,22 55:13 57:7 58:20 63:17 64:6 67:13 68:11 71:15,25 72:2,18 77:16 80:18 86:24 87:17,25 88:6 96:8 99:5 102:19 107:13 108:1,5 112:2 114:7 115:18.23 125:23 126:24 133:24 134:21 141:5 142:9 152:20 153:11,16 156:18,19 158:8 163:18,23,25 172:9 182:23 183:10 194:19.21 195:21 197:4 206:12,24 208:8 209:17 **surety** 200:19 surface 114:20 132:2 139:24 145:18 146:4 156:25 161:18 166:11 168:3 170:2 173:17.18 174:6,8,21,21,24 176:13,17,18 177:10 178:1,2,6,10,12,15 179:14 190:14,25 192:25 198:13 199:6,7 201:12,15 207:22,22,23 208:3,24,25 209:1,2,5,21,25 210:9,10,13 213:25 surfaces 103:1 106:24 111:20 119:8 126:25 127:1 145:17 146:22,22 147:8 161:14 173:16 178:19 179:18 180:17 189:11 192:7,12 193:9,12 198:10,12 199:4 **surgery** 10:15

**SUITE** 2:5,16,21 3:5,9

surprise 142:9 surprised 21:3 surrounding 132:1 survey 200:18 **surveyed** 201:13 **survivability** 187:9,12,23 survives 130:6 sustain 103:19 **swath** 15:8 swear 6:22 **swim** 140:6 sworn 7:3 220:4 Symposium 76:18 **symptoms** 144:19 **Syndrome** 125:16 **system** 171:17,19 172:12,16 **systolic** 50:24 51:12 52:7

## T

**T** 5:1.1 146:2 t-e-t-r-a 206:15 **table** 140:15,16,17 178:23 179:14 199:2,3 210:10,25 211:5,9,10,10,15 212:22 215:4 215:6 216:24 **tabletop** 179:8,12 take 7:21 8:8,11,12 12:15 27:18 32:5 40:6 45:4,7,17 52:18 57:15 63:23 73:4 80:15 88:22 89:16,21 90:7 91:8 105:11 109:7 120:2 122:1 142:16,22 142:23 143:13 144:14 149:8 180:3 188:12,23 189:2 190:25 197:4 198:22 216:9 217:14,20 217:24 taken 1:15 5:5 45:13 91:12 118:3 130:5 140:25 142:25 188:25 202:2 215:24 219:3 220:3 takes 89:4 161:24 talk 21:18 66:10,11,21 70:15 77:15 102:24 111:17 126:18 143:8 146:19 148:11 152:3 166:2 172:20 176:3 177:22 180:14 192:5,23 208:15 212:9 talked 15:1 20:10,12 26:25 60:12

70:23 93:2,18 97:7 109:24 120:5 129:17 133:10 176:11 196:23 199:1 207:16 214:14 talking 28:16 64:6 69:10,13 76:17 107:15 109:15 126:23,24 126:25 131:11,12 132:25 133:13 139:21 143:18 152:21 162:3 174:9 175:12,21 176:17 177:4 183:3 194:19.21 196:14 196:14 199:3 talks 54:7 60:8 144:15 153:10 156:7 164:22 167:6 171:13 task 42:15 43:9 46:16 task-oriented 47:20 taste 57:22 taught 65:2 117:6,8,9,11,13 teach 20:17 teachers 52:8 teaching 56:17 62:22,23 63:6 64:17 team 91:1 92:18 technical 8:18 10:10 11:22 technology 88:13 **telephone** 112:3 127:25 tell 11:9.25 13:9 14:24 16:9 27:3 29:25 34:14,15 40:9 42:18 52:23,24 56:19 57:6 63:2,14 64:2,4 65:15,17 67:7 68:10 72:16 78:12 81:1 83:6 86:13 88:5 91:23 102:18 105:6 110:20 118:10 126:22 132:7 136:4 140:3 142:8 149:10 150:21 158:10 161:12 173:21 177:2 190:7 195:5 196:1 204:16 215:16 telling 136:4 209:10 218:7 tells 176:22 temperature 131:7 183:5 188:4 217:25 219:9 temperatures 181:15 182:14 185:9 189:25 ten 75:5 180:18 tenet 170:22 tenets 12:22 13:1

tenure 14:6 terfenadine 68:5,6,11,14 term 10:10 21:12 39:6 132:5 214:13 terminology 163:3 terms 11:23 87:8 107:14 140:2 184:15 tersely 27:7 test 12:15 72:19 106:23 107:2 150:13 182:2 tested 115:7 150:10 **testified** 7:4 95:13 testify 133:19 220:5 testimony 7:21 75:19 118:25 153:17 183:11 191:5,18 220:3 testing 111:20,22,23 151:20 tetra 206:15 tetrachloride 206:6,8 **Texas** 10:23 14:7 16:6 18:21,24 19:12 22:14,16,17,18 23:9,10 29:12 63:7 64:16 121:17,21 Texas's 20:24 text 169:8,9 textbooks 62:16 thank 7:9,10 17:2 23:19 91:14 92:25 116:16 163:9 187:14 197:15 219:15 theft 205:25 theory 11:12 52:22 53:3,19,20 54:5.7 therapeutic 85:20 therapies 75:13 therapy 72:19 73:9,9,11 thing 14:5 51:6 52:24 53:12 60:5 68:17 80:17 83:9,11 86:6,7 140:10 163:12 182:21 things 20:9 22:1,9 26:14 28:13 34:8 101:23 103:7,13 127:3 128:23 131:6,8 132:25 166:4 196:24 think 7:24 9:6 10:21 11:25 12:10 13:12 19:21 24:11 25:24 29:1 29:17 31:1 32:5,9,21 37:2,3 41:20 43:16 44:5,5 47:21 53:16

tens 182:25

53:18 56:12 59:13 60:4 64:25

70:10,14,17,23 73:19 76:17,19

78:13 79:20,20 80:9 82:4 83:2 83:9.11 85:4.20 92:3.10.10.11

treat 68:22 74:5 75:25

treated 86:20

95:13 96:4 97:5 100:4,12 101:22 107:16 109:9 113:5 114:6,6,8 115:22 116:20 117:3 118:10 119:2 123:2,3,5 126:4 126:10,12 129:17 139:15 141:14 148:17 150:16,25 153:25 156:1 164:3 172:14,18 174:13 175:24 176:8,9 181:22 183:4,12 184:9,14 190:8 196:6 196:6 200:21 202:20 203:2 205:1,1 206:4 212:17 213:17 214:8 215:12 216:7 217:12 219:14 thinking 88:12 132:2 third 24:15 107:2 177:24 190:10 190:15 **Thirteen** 50:18.19 **THOMAN** 3:13 thought 13:13 43:25 99:3 109:9 125:20,22 144:4 147:24 148:4 148:6 173:12 181:6.9 **thousands** 85:10 142:10 172:25 195:20 threat 192:21 three 19:21 34:16 54:8 57:4 62:5 83:25 84:1,21 94:2,2 96:11,12 122:18 180:23 191:10 211:14 211:14,25 212:3 throw 137:1 ticarcillin 41:20 **TIFFANY** 3:13 tight 211:23 tightly 135:23 141:11,17 211:10 time 6:5 8:11 11:21 14:2 27:16 31:9 36:22,23,24 37:1,24 46:25 47:3,5 52:16 93:11 94:14 95:12 98:16 120:23 121:7 124:17 126:14 127:17,18,21 145:18 149:6 152:10,10,21 153:13,19 188:16 197:4 206:13 207:16 times 57:13 83:7 126:20 152:23

163:18 191:10 **tipped** 47:11 tires 88:17 title 30:25 49:14 70:5 173:6 **titled** 71:16 titles 60:17 today 9:15 11:23 75:19 98:20 187:12 **told** 24:7 39:25 88:21 97:5,6,19 155:9 202:1 top 62:13 67:15 209:5 212:22 topic 146:20 total 154:11 170:18 touch 191:7.8 199:7 touched 164:13 191:23,24 touches 149:4 198:13,16 **touching** 209:21 tough 179:23 track 145:6 tract 86:24 87:2 **trade** 68:11 **tragic** 27:25 training 24:9,10,19,21 25:6,12 75:6,8,9 127:8,9 trans 168:23 transcribed 220:7 transcript 1:12 5:9 220:8,9,9,18 transferred 155:10 **transformation** 96:1 100:23 101:1,5,14 105:14 106:10,12 106:18 119:22.24 128:4 131:18 161:17,23 162:7,10,11,12,13 162:19,21 168:2,22 169:19,24 170:17,18,19,20,21 transformations 131:17 transformed 161:6,9 166:14,15 166:21,23 168:24 169:1 172:5 172:10 transforms 169:6 translational 37:13 **transmission** 4:14,17 173:8 196:11 198:10 199:19 202:25 203:25 204:10 transparent 132:20 travel 139:22

treating 26:15 27:4 49:22 86:16 127:9 treatment 13:3 49:3,18 50:10 51:11,22 52:6 68:12 70:6 81:2 81:5 87:22 treatments 50:11 treats 69:24 trial 5:12 12:20 13:6 16:17.21 43:10 50:23 65:8 66:4,15 78:17 85:8,12,16 133:22 160:5 trials 12:8,11,13,25 16:5 17:3 18:2.4 23:14 33:22.25 34:2 51:4 55:20 64:8,24 65:18,18,19 65:21,21,22,23,24 67:21 70:18 70:19 78:25 79:5,18,19 **tried** 175:15 trier 100:13 triggers 90:1 **trillion** 81:3 83:13 trojan 71:25 trouble 68:4 92:6 true 44:23 50:7 60:18 79:24 163:4 178:9 179:19 190:4 197:23 220:7 **Trust** 87:17 truthfully 9:18 **try** 7:16 22:15 65:25 66:9 76:14 140:12,12 141:13 142:5 173:22 205:8 trying 22:6 27:12 44:16 49:21 59:18 60:4 124:21 128:11 142:2 147:24 152:12 153:11 156:10 185:5 187:17 197:3 206:22 209:8 tuberculosis 89:3,6 **Turn** 146:16 turned 14:17 33:24 141:22 turns 87:15 130:8 tutored 117:10,16,19 two 25:5 33:17,24 34:16 36:19 36:21 37:6 41:23.23 44:6 49:8 51:20 58:7,12 62:5 67:14 84:20 89:4 92:12 94:20 103:20 117:4

117:20 119:3 122:18 145:7
150:13,14 167:10,14 177:21
180:19 189:17 190:8 198:22,23
205:12,16 215:22
two-prong 52:3
two-thirds 120:4
type 26:4,22 45:1 49:15 51:22
59:7 68:25 77:7 87:3 93:12
123:9,23 124:9 171:19 172:12
175:9 177:3 213:15
types 40:24 43:19 49:19 88:10
93:21 175:5
typically 48:15 208:19
typo 150:22

**U** 5:1 **U.S** 176:9 **Uh-huh** 121:3 **ultimate** 99:23 110:1 ultimately 98:2 100:13 101:6 138:19 209:19 211:23 212:18 umbrella 38:7 39:8 unambiguous 100:6 underestimate 184:10 undergrad 10:1 **undergraduate** 41:5 42:8 43:22 44:2.3 **underlying** 11:12 38:19 44:7,9 64:5 80:2 **underneath** 30:19 33:12 **understand** 8:1.14 11:25 18:23 21:24 38:23 42:22 43:4,5,5,6 51:24 53:23 54:17 60:13 66:5 95:17,17,18 103:1 104:3 105:3 107:19 109:3.19 110:20.21 118:7 120:1 128:11 129:4 133:14 136:13,23 139:1,1,2,3,7 153:22 187:17 190:12 191:21 194:5 201:23 210:6 211:6 understandable 189:14 **understanding** 11:13,15,16 95:23 102:5 116:22 118:5 135:15 146:12 147:7,8 154:19 155:9 171:9 175:4 207:12

220:8 understands 11:14 167:18 understood 8:5 24:4 49:2 54:11 54:15 163:19 165:14 167:12 undertaken 104:1 **UNDERWRITERS** 1:7 **undone** 189:15 **unfortunately** 11:17 19:20 87:16 179:24 194:14 unhealthy 180:3 **United** 176:1 units 47:16 universe 32:25 49:23 60:1,19,22 115:3.3 122:19 134:8.11 universities 75:16 university 10:5,6,19,20,23 14:7 16:6 19:12 20:23 22:13,16,16 23:9,10 29:12 63:7 64:16 74:2 121:17,21,23 unraveled 141:16 unrelated 130:12 **unsafe** 161:15 unstandardized 187:19 unusable 215:2,3,6 updated 37:1 **Upper** 86:24 **urgent** 28:14,15 use 5:13 14:18 38:2 49:24 55:2 70:2 96:2,5 100:23 101:1,5,13 101:14 105:14 106:10 108:8 118:4 119:22.23.25 128:4 131:10 137:15 140:11 143:9 163:22 165:18 167:19 174:13 193:8,8,18 196:24 204:4,19 212:13 214:17 215:4 useful 134:8,11 uses 175:10 utility 187:22 **utilize** 165:8 171:20 173:10 utilized 137:4 173:11 181:14 **UV** 186:3,6,11,14,16,17,23 187:1

**V vaccine** 65:19,21 70:18,19

valid 220:17 value 122:15 135:22 187:18 van 180:4,7,9,11,15 181:2 206:24 207:1 208:20 variability 188:3,9 variation 122:6 **varied** 65:13 varies 144:9 various 10:13 30:18,19 65:1 145:15 180:17 **varying** 55:6,8 vector 173:13 vehicles 196:2 **vendors** 110:7 ventilated 171:14 ventilation 171:16 172:6,12,16 172:19 ventricular 55:17 veracity 209:23 verbally 7:20 **verge** 140:5 version 35:10 53:22 66:1 143:20 186:9 versus 1:6 123:11 124:20 130:24 131:13 168:16 184:16 viability 188:4 **viable** 177:18,19 180:22 184:13 184:16,18 189:19 vicinity 104:7,8,10 109:8 114:4 138:18 victim 140:25 **victims** 140:7 videoconference 1:12 6:10 view 64:11 137:10 146:3,12 200:24 violation 149:20 viral 68:13,24,25 72:21,22,25 74:7 80:10 82:6 86:17,18,19,20 86:22,24 87:2,3,10 88:10,15,19 88:19,20 105:3 167:10 177:1,2 VIRGINIA 3:8 virologist 85:23 86:2 virology 76:23,25 77:3,12,13

80:10 85:21 86:1 139:20 169:8

169:14 176:22

187:5 188:3

volunteers 91:2

virus 67:4,12 68:22,24 69:20 153:11,17 162:2 176:3,17  $\mathbf{W}$ 70:15 80:21,22,23 81:10,13,15 177:3 178:11 188:17 192:6 **W** 2:10 3:8 81:17,19,20,21 82:2,3,10,17,19 193:7 194:21 199:3 W-a-a-l-s 206:24 82:24 83:2.3 84:17 88:2 89:22 we've 22:8 26:25 60:12 61:1.2 Waals 207:1 208:20 90:5 93:5 95:8,8 100:4,9,14,19 70:10 76:16 83:2 91:7 120:5 wait 99:11,11 108:14,14 128:6,6 101:12,20 102:25 103:12 142:18 164:4,17 171:24 188:17 129:19,19,19 138:3,3 162:24 104:14,15,20,20,23 105:1,16 202:14 214:13 162:25,25 216:21 106:3 107:12 110:2 112:9,12 weak 162:16,16 208:19,19,21 waiting 139:24 148:2,6 203:4 113:22 114:4 119:8 129:16 weakens 164:20 **waived** 5:10 130:6 131:12,19,19 134:18 wear 216:12,19,25 217:19 walk 9:22 13:23 142:7,14 143:23 144:6,8,10 wearing 217:6,15,22 walked 109:22 160:21 161:8,13 164:23,23,25 website 191:2 walks 104:22 109:16 165:16 166:14,22 167:24 Wednesday 67:18 115:23 want 17:15 21:23 24:6 32:14.15 week 27:14 28:19,19 94:2 115:22 168:23 169:1.5 172:22 173:13 38:18 51:17 52:3 53:22 57:22 173:15,18 174:5,9,10,15,20,22 116:1 63:2 71:25 83:8 84:22 96:8 weeks 34:16 94:3 96:11,12,14 174:23 176:12,14 177:11,18,25 101:24 102:2,11 107:13 108:1 178:5,5,10,14,14 179:13,16 weighed 133:8 108:5 110:8 112:2 120:13 182:2,16,22 184:9,10 186:2,9 weight 67:9 132:3,8,17 133:3,15 133:15 137:1 140:11 146:19 well-established 162:1 186:12 189:19,23 190:14,15,22 152:20 153:16 159:23 190:11 191:1,2,7,22 192:2,10,14 193:7 well-phrased 7:25 190:19 191:13 193:20,22 194:8,14 196:8 198:17 199:6,9 Welsh 3:8 6:14 206:11,23 208:10 213:10,11 204:19 207:21 208:1,2,6,17 went 10:19,22 20:23 59:15 67:7 215:16 209:10,19,24 210:1 212:3,21 71:17 104:19 152:6,22 153:13 wanted 58:12 74:2 163:22,25 virus's 194:15 were-- 75:24 warrants 73:23 viruses 59:20,22,25 60:8,16 62:8 weren't 85:22 135:2,7 155:4 wasn't 27:22 41:8 65:10 69:3 62:11,15,16,19 65:20 67:24 whatsoever 147:17 70:5 72:22 73:19 76:1 86:1 69:3,21 70:9,20,24 71:1,16,23 Wheaties 98:14 87:5 135:9 138:19 185:5,6 72:17 73:1 75:22 76:2,10 77:8 white 205:21,22 water 139:23 77:14 81:3,5 83:8,9,13,13 85:7 wild 89:11 wax 174:9,10,22 175:2,4,9 86:15 95:3,17 99:18,18,21 WILLIAMS 2:19 207:23 208:18 101:2,4 129:25,25 130:8,8 willing 97:8,14 100:3 waxes 175:5 131:9 139:3,21 140:4,10,14,15 wind 208:23 way 39:12 43:12 61:4 66:9 75:14 windows 171:23 172:2 183:18 140:16,21 142:9,15 157:10 91:24 117:24 138:14 142:5 167:18 173:23 177:19 180:19 186:11 145:24 147:18 151:1 152:12 183:3 185:16 192:17,18,19,20 wipe 190:25 193:9 170:8 171:1 188:8 190:20 192:21,24 193:13 209:3 210:9 wish 54:6 196:6 212:19 210:10,16,17,20,21,24 211:2,7 witness 5:9 6:9 41:13 45:4,10 **Wayne** 1:14 7:2 211:15,18 212:6,16 213:24 91:10 92:25 99:15 110:18,20 we'll 6:11 9:1 19:25 32:7 38:1 Visit 73:22 110:23 118:23 138:5 142:18,21 60:21 83:10 108:20 139:9,20 vitae 4:11 13:17 203:4 146:18 177:23 vitro 41:7,9 42:2 witness's 193:23 we're 9:2,21 11:21 13:18 30:13 **volumes** 55:18 **wonder** 82:23 38:25 53:24 56:6 100:14 wood 174:9,10,22 207:23 208:18 volunteer 87:12 90:22,23 103:14 107:14,15 111:17 115:2

119:18 131:11,12 132:24 133:1

139:14 143:17 146:20 152:20

wooden 140:15

word 69:5 95:23 108:16 110:5

116:16 125:11 126:19,21 **13** 4:10 29:2 50:16 51:1 Y 140:11 149:8 174:13 179:3 **14** 52:14,17 54:19 **Y** 3:8 204:20 210:3 **1434** 220:14 **veah** 19:25 36:22 40:10 42:22 words 107:25 127:16 131:24 **1455** 5:11 65:3 70:14 84:24 86:11 91:11 164:6 167:20 170:3,4 193:7 **15** 53:12 54:19 55:5 65:4 121:7 92:11 93:10 119:24 121:16 work 10:2 12:8 16:19 17:23 18:8 121:10 125:3 127:24,24 134:2 142:21 20:1,6,20,20 21:18 22:21 23:15 **150** 57:4 73:20 146:7 148:4 149:3 154:7 23:16,19 25:2 27:2 28:6,15 **16** 55:13 147:16 164:25 185:2 187:14 188:15 **16-hour** 28:7 32:1,19 38:5,20 43:23 44:3,12 197:4.7.10 206:10 217:10 **1625** 2:5 45:12 48:25 52:20 54:12 55:21 year 16:24 17:5 24:23 27:21 57:4 60:19 61:13 66:12 71:21 **16th** 145:11 62:24 73:20 120:22 121:8 **17** 54:23,23 55:13 68:2,3 76:7 86:13 87:11,12,12 90:12 **years** 24:15 29:2 36:10 44:6 90:22,23 91:22 92:4 93:12 **1702** 2:16 69:17 75:5 78:13 83:10 118:3 **17th** 96:18 98:1 99:7 100:18 97:21 98:8,23 99:1,2 120:12,16 122:22,23 125:19,24 128:10 120:23,24 121:5,12 196:22 105:8 127:25 137:24 141:8 171:8 212:1,1 **18** 122:2,2 125:2 **Yep** 66:17 **18487** 1:22 worked 15:23 19:23 22:23 23:11 Yersinia 89:13 23:21 25:21 27:20 28:3,18 93:8 **19** 54:23 122:17,25 vesterday 12:12 93:10 125:25 126:3,5 **1965** 13:12 **Yolanda** 1:17 6:2 220:2,24 worker 150:14 **1973** 44:5 **you-all** 93:12 working 17:19 19:11 20:18 **1974** 10:5 24:25 25:1 44:5 25:11 60:6 90:19 93:14 98:5,13 **1978** 10:6,7 24:25 27:8 42:7 98:15,17 116:9 121:1 178:12 **1979** 45:18 zero 104:10,11,11 209:8 **zoom** 8:21 92:11 93:2,18,23 **1980s** 21:16 works 137:5,22 139:7 **1981** 10:21,22 47:15 0 Workshop 68:8 **1984** 68:8 world 200:5 206:8 **1986** 23:1 1 worldwide 201:4 202:7 **1987** 10:25 17:21 37:20 62:23 **1** 1:4 2:21 4:10 13:19.20 43:15 worried 159:21 73:23 74:18 75:1,10 48:23 61:2 70:10 151:7 155:6 **1992** 28:25 29:3,11,13 worse 14:11 155:22 211:15 **1996** 56:23 58:8 worst 27:23 **10** 1:25 39:4,22 43:14 219:18 worthy 30:6,15 74:23 1st 155:18 160:25 **100** 54:1 63:13,21 wouldn't 18:12 74:17 83:20 **102** 76:16 2 117:23 190:20 213:2 **10th** 94:1 96:15,17 97:25 99:6 **2** 4:12 29:16 115:10 211:21 wrap 209:8 215:23 100:17 105:8 127:24 137:24 212:6.9.15.18.20.21 write 101:21 103:2 114:18 151:6 154:8 155:21 160:24 **20** 54:24 60:23 62:13 121:10 writing 112:15 **1100** 2:21 3:9 126:18 127:6 128:15 131:23 **written** 62:18 **112** 76:13,17 **200** 60:11 wrong 64:22 154:1 179:18 **115** 4:12 **2000** 3:5 56:23,24 58:8,8,10,11 190:24 191:3,11 200:24 212:14 **11th** 220:19 58:15,15 59:8 218:4 **12** 48:19,24 **2001** 63:4 64:3 65:2 wrote 53:7,7 111:19 136:5 **12:11** 1:25 **2002** 56:24 58:8,11,11,15 59:8  $\mathbf{X}$ **120** 144:6 **2003** 79:12 80:2 84:3,4 85:6 **125** 143:23 144:6,8,13 **2004** 80:2 **X** 4:1

 16:23 65:3 155:4,5,12,21,21 157:17 158:3 **7:00** 191:13 158:20 160:24 44:4 185:18 16:23 65:4 2:21 77:2 2017002 220:24 2:11 4:16 203:15 205:5,6 23:5 31:22 125:2 2:6.16 28:18,19 181:22 182:15 1:25 37:2 94:10,12,17 95:13 3:5 183:15,18,24 201:24 145:12 147:16 151:7 157:14 1:23 3:9 122:12 196:25 219:18 220:20 3:9 72:7 **2020-02558** 1:4 43:23 188:11 189:2 147:19 148:9 60:7 **73.4** 181:17 182:14 **456-8624** 2:12 4:13 105:23 106:19 150:7,8 154:1 43:24 44:4,4 77:15 80:14 86:3 4:16 **482-5811** 2:6 54:24 62:13 72:5 132:11 77:11 80:14 86:3 156:9 167:6 86:6 220:5 4th 158:20 61:3 72:5 134:5 10:8 25:1 27:9 41:5 42:10,12 220 4:6 43:23 44:2 3:10 10:7,8 27:9 4:3 59:13 203:15 135:19 137:3 181:17 **5:30** 188:17 54:25 72:5 139:14,15 150:20 54:1 156:21 165:17.19 166:1 849:2 151:4,24,25 153:2,7,13 154:6 168:2 833-5600 2:22 154:11,16,18,25 155:12 189:13 2:6,12,17,22 3:6 1:15 7:2 189:21 72:7 164:22 165:23 175:13,14 86:9,10 **24-hour** 155:22 77:10 75:3 63:15,25 64:18,21 70:11 173:3 171:13 70:12 77:9 139:21 172:20 72:10 142:19 173:14 207:15 9/29/20 4:16 **27th** 151:6 155:17,21 160:25 70:23 71:12 **566-1311** 3:6 70:17 189:19 1:14 7:2 220:24 **28-day** 189:22 70:13 177:21,22 909 2:5 29th 202:23 **582-6998** 2:17 70:23 71:12 70:21 2:16 **5th** 157:14,16 158:21 58:10,11,14 4:13 203:1,3 211:18 212:18 70:22 30 28:18,19 72:6 143:22 6 120:2.4 60:23 72:11 **6:00** 188:21,21,22 **346-0285** 3:10 **6:25** 219:17 72:6 144:14,14 144:6,13 2:11 72:7,8 76:13 72:7 180:2 3:5 184:20 144:23 **37:2554** 220:4 145:11 4:5 72:4,4 73:21 **3rd** 151:6,6 154:3,5,7,13,15 7/3/20 4:14