UNITED STATES SENATE

SENATOR RON JOHNSON

Senate Homeland Security and Governmental Affairs Committee

328 Hart Senate Office Building
Washington, DC 20510

Affidavit of Justin Leslie in support of Senator Ron Johnson Investigation into the Safety and Efficacy of COVID-19 Vaccines

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I, Justin Leslie being duly sworn, depose and state as follows:

1. I am a living man.

2. I make this affidavit as a whistleblower under the whistleblower provisions of the False Claims Act, Title 31 U.S.C. §3729 et. seq., in support of an investigation or claim as testimony in support thereof. This is a NOTICE OF LIABILITY FOR CRIMES AGAINST HUMANITY for the events that occurred across the United States of America and internationally across the world during the so-called COVID-19 "Pandemic." These crimes include murder and severe injury through mass vaccination and psychological torture to mankind.

3. The COVID-19 vaccines are a biological weapon and should be removed from the market immediately.

4. I communicate in everyday English not legalese; I do not consent to contract or joinder or grant power of attorney or act as surety or trustee.

5. Virology has been proven pseudoscientific and so there are no need for any vaccines ever. See exhibits F, G, H, I and J attached for evidence.

6. The facts are my own and arrived at from my professional and personal experiences.

Experience & Credentials

7. I am fully competent to make this declaration and do so voluntarily and do testify to the facts and matters set forth herein.

8. I began working for Pfizer Inc. as a contractor for Mindlance Inc. in March 2021 in Andover, Massachusetts. I was hired and trained to be a Formulation Analytical Scientist on the mRNA COVID-19 Vaccine platform, as well as mRNA Flu and self-amplifyingRNA Flu formulations ("Vaccine Work"). I resigned my position on April 12, 2022, for reasons described later in the affidavit.

9. I graduated from Rhode Island University with a Bachelor of Science in Biomedical & Pharmaceutical Sciences with a Minor in Business, *cum laud*. My complete *curriculum vitae* is attached hereto as Exhibit A and incorporated herein by reference for all purposes.

10. My initial job description for Mindlance Inc. was as follows:

Job Description:

- The individual in this role will be a highly skilled, meticulous and motivated scientist who will focus on analytical characterization of mRNA and Lipid Nanoparticle (LNP) formulations. He/she willwork cross-functionally to design and coordinate analytical characterization characterize and/or implement fit-for-purpose, robust, characterization methods
- Identify, develop and implement novel biochemical and biophysical methods to further enhance knowledge or to address potential analytical characterization gaps for mRNA and Lipid Nanoparticles
- He/she will participate in performing biochemical and biophysical characterization (HPLC, cIEF, light scattering, particle characterization, spectroscopy, calorimetry) of lipid nanoparticles, lipids and mRNA
- Preparation of samples for characterization studies
- Perform analytical testing supporting process development and product release, including spectroscopic, chromatographic, biophysical, and molecular biology techniques
- Maintain thorough electronic lab notebooks and documentation records
- Deliver reproducible and impactful results under ambitious timelines
- Execute and troubleshoot standard protocols, develop and adapt new protocols into practice, and query literature to incorporate additional assays as needed.

11. During my time working as a contractor for Pfizer through Mindlance, Inc. and based upon my personal knowledge and expertise, my professional opinion was that the entirety of the vaccine roll out was rushed and not being done in either a safe or ethical manner. Given that virology itself is a pseudoscience also creates no need for a vaccine, and should be considered a bioweapon.

12. Based upon the direct information I possessed from the Vaccine Work, I could immediately see that information was being suppressed and withheld from the general public. The suppressed information was important safety and effectiveness data which from both an ethical and regulatory perspective the public had a right to know.

13. During the time I was performing the Vaccine Work, I contacted the investigative journalism group, Project Veritas, on September 27, 2021. Project Veritas immediately reached out to me and asked me to work with them as a "whistleblower" source. Project Veritas represented itself as a media source that wanted to help bring truth and make sure members of public who were interested in receiving these COVID-19 shots were actually receiving genuine informed consent regarding taking them. This occurred simultaneously with more and more entities beginning to mandate taking the COVID-19 vaccines and boosters as well as Pfizer seeking the approval of childhood vaccines.

14. Project Veritas provided me with covert equipment and guidance on the types of information I should elicit from others at Pfizer. As an unwitting whistleblower, I reasonably believed that because of representations made to me by James O'Keefe and various Project Veritas employees, that they would do the right thing and release a story or stories based upon what I ultimately witnessed at Pfizer because the public has a right to know the information being kept from them about the COVID-19 vaccines. I was trustworthy of Project Veritas and gave them all the information I ultimately obtained as a result of my whistleblower work for Project Veritas and kept nothing for myself.

15. After two months of my undercover whistleblower work, James O'Keefe the CEO of Project Veritas, offered me an opportunity to work directly for Project Veritas and to not tell my story, which would have increased vaccine hesitancy on a worldwide scale. At the time I was not fully aware, but now presently believe I was offered this opportunity to work directly for Project Veritas in a full-time capacity, to thwart further my ability to blow the whistle on the crimes against humanity that have been committed by Pfizer, Inc. and all associated businesses who recklessly pushed forward with rolling out the COVID-19 vaccines knowing they were neither safe nor effective. Pfizer had knowledge of the adverse reactions were reasonably caused by its COVID-19 vaccines and/or booster; Pfizer had knowledge that deaths were occurring shortly after people received its COVID-19 vaccines and/or boosters; Pfizer knew the statements released to the public about its COVID-19 products staying at or near the injections sites were completely false and in fact the COVID-19 products were being delivered throughout the body to include the heart, liver, reproductive and other organs. See excerpts from a Pharmacokinetics Organ Distribution study in Wistar Hahn Rats attached hereto as Exhibit B ("Organ Study").

16. The charts on the Organ Study show the distribution of the Lipid Nano Particles ALC-0315 and ALC-0159 (LNPs) through all organs of the body. The chart shows distribution of the LNPs to the heart, testes, ovaries, and other vital organs, in which these LNPs are not supposed to be going. This can be linked to the widely reported myocarditis, pericarditis, and menstrual cycle and fertility issues that have happened since the COVID-19 vaccines have been rolled out.

17. I took the offer from James O'Keefe to work for Project Veritas on May 11, 2022, as a contractor. At no time while I worked as a contractor for Project Veritas did James O'Keefe do a story about the below-referenced materials I obtained at the direction of Project Veritas and in fact, James O'Keefe informed me Project Veritas would not be doing any stories on most of the information I obtained. So, I took the offer to do work for Project Veritas, at least in part, with the intention of obtaining my Pfizer whistleblower information back.

18. The following paragraphs are references and summaries of conversations I took part in all at the direction of Project Veritas. I was physically present during and a member of the discussion during each conversation.

19. kanwal pfizer (rev.com) Kanwal Gill is the Leader of the Analytical Team for Pharmaceutical Research and Development for Pfizer, Inc with her official position being Principal Formulation Scientist. In the meeting we discussed mutations of the COVID-19 virus and potential causes of the COVID-19 virus mutating into variants such as Delta and Mu. It is suggested during this discussion that the virus mutates due to errors in its replication process, which are more likely to occur the more people it infects. It is also discussed the role of vaccines in this process, suggesting that vaccines could potentially influence the virus's mutations. However, it is clarified that vaccines are crucial in preventing the virus from spreading and mutating further. The need for booster shots is also discussed and the ethical considerations of distributing them, as well as the potential side effects of vaccines.

Below are some direct quotes from the conversation:

ME: Why are we worried about booster shots over here if we're going to, shouldn't we be giving them?

KANWAL GILL: Of course, but there's so much of politics. [END QUOTE]

Next, I aske Kanwal about the ethics of giving another experimental injection in the form of the booster shots that were going into people's arms at the time of this:

ME: I mean just thinking ethically, is it okay to give people another experimental injection? Are we sure it's going to be safe and effective? Are we just rolling these out because we want to roll them out for money?

KANWAL GILL: Yeah, I think the drive is to be able to vaccinate everybody and stop it. But also as this is a real time data that we are generating, we don't even know when we started vaccinating people we no idea how it's going to look like mRNA vaccines have been there for 50 years. No, but nothing made to clinical. Why? Because mRNA vaccines have been known to have side effects. Moderna has been working on it for 10 years. This is not the new thing. But why did not reach trial was because of all these side effects. We, Pfizer and Moderna used the emergency and the pandemic to get through it now, but even C D C, not we have.

ME: So, it couldn't ever get F D A approved until we had this emergency. We had to try something new. But why didn't we try a vaccine formulation that has been effective in the past with egg yolk or something that they use with flu, right? I dunno.

KANWAL GILL: Yeah. So, we here did not do it because we already have collaboration with BioNTech on flu before actually Covid happened. So, mRNA, so what the story behind we starting on this because BioNTech tech guy who would the beyond tech, C E O, he called Kathrin Jensen. He called Katherin Jensen and he's like, 'Hey, I have this technology but I don't have resources to make it big.' So, Katherin was like, well, we had a collaboration in place before for flu, so we wanted to give it a shot. It is a new thing. Nobody knew anything about it. We never had this, everything is just fresh from last year.

Learning on the Fly. So it was turning, it was on a fly pandemic. Just roll it out, roll it out, roll it out, and Moderna was there. So we had that, okay, someone is also doing the same thing and they're in business for 15 years working on the same thing. So yeah, let's just take a call. It was a huge risk without anything. There was no time to think it was a time to act. That's how we are operating. [END QUOTE]

These direct quotes are pretty damning. It shows the malfeasance of disregarding safety during the vaccine roll out, and doing this as fast as possible, not regarding the risks of vaccine injury and death on a worldwide scale if something went wrong.

Here is the last direct quote from Kanawl Gill from this conversation:

ME: So, if the vaccine is effective against the variants and everything, then who's being hospitalized? If the unvaccinated people are being hospitalized and why do we care? They have a vaccine. They have a choice. They either can risk it or not. So who's getting sick if they're so worried if the vaccine's so effective?

KANWAL GILL: No, but there are cases right with the vaccine or it is not so effective. Actually there is some level of protection that it is providing a case. That variance there is somewhere here. Pfizer still is saying that it provides some level of, but is that level 90? A hundred percent? No, that could be 50%. So Pfizer is not also confident at this point because this virus is playing

with us. But we are saying at her, we can protect you, but okay, this will help you. You might not die with it if you take Pfizer vaccine and get infected, but you might also die because, so there is a 50 50 chance to it at this point, but I think not 50 50, but majorly if you have a vaccine, so I have a vaccine, I'm not as afraid to get sick because I think based on what I know, okay, maybe I'll get my immunity and my sister, I will. Hopefully it'll not hospitalize me because I have a vaccine and I think if I take a booster, I'll be in a better shape because now I'll have more antibodies to fight against. But again, I don't know how much I even cannot answer that booster, what impact it'll have on my body. Because we already had side effects from two shots, right? There was fever and all, but we don't know the long-term side effects. Of course, we don't know what is going to happen to us. I know.

ME: So we need to make sure that we're doing these long-term studies, right? Yeah.

KANWAL GILL: Which are happening, but there's no long-term data on the side effects of people. I think we will know in a year or two actually what actually is happening. Only Johnson and Johnson side effect came out quickly. There was a clot that there was happening with Johnson and Johnson's, but other than that, mRNA and LNP are very sneaky. I think we are not still seeing a lot of huge complications, but I think we have to really extend this. That is why toxic safety studies, the phase one, there are three types of trials. Phase one, phase two, phase three. We just went straight up, we just did this. But this is why it takes 10 years for a vaccine to come out. It takes years of observation, blah, blah, blah one, then two and three here we are doing everything at the same time. So everybody is as puzzled. Nobody has a hundred even from CEO, Pfizer, CEO, and even Ughur Sahin won't have once because they are looking at it closely watching it. At this point, that's all we can do. And I feel like, yeah, it's also becoming a money game, bringing all these mRNA projects, blah blah, blah. But we are just part of this company and we need to keep our focus on doing that. [END QUOTE]

The conclusion is that that more long-term studies are needed to fully understand the effects and efficacy of vaccines which had not been done, and in reference to the conversation with Bakul Bhatngar asking him about vaccinating the control group, he says that Pfizer vaccinated the control group in the clinical trial. This clearly destroys the scientific clinical protocols of allowing for a control group to naturally play out, which is UNETHICAL. This is regardless of whether there is a virus or not, which there isn't. Kanwal Gill also states that Pfizer CEO Albert Bourla and BioNTech CEO Ughur Sahin do not have answers and the mRNA vaccine rollout was becoming a "money game" with all their other mRNA projects happening. See Exhibit C attached hereto.

20. Ramin Pfizer (rev.com) Ramin Davari is a Pharmaceutical Formulation Scientist for Pfizer. I had a meeting with him to discuss potential adverse reactions of the COVID-19 shots. We discuss a story about a woman who died on the same day she received a Pfizer booster shot. Ramin Davari emphasizes the importance of distinguishing correlation from causation, and notes that it is not their job to investigate such incidents, but that there are people within Pfizer who do. Ramin Davari goes on to say that Pfizer receives "zillions" of reports every single day on the COVID-19 vaccines of adverse events. The conversation ends with a comparison to the food industry in which companies engineer products specifically for the purpose of consumption, regardless of the health risks with Ramin Davari stating that companies are engineering their products "specifically for me to take the next one" to increase my consumption and that the product will ultimately "affect my heart and I'm going to die." We also discussed allergies to certain substances in drugs, and the responsibility of

pharmaceutical companies to disclose these. See Exhibit D attached hereto.

21. Below is a transcript of a conversation I had with Nick Warne the Vice President of Research and Development for Pfizer, Inc. while I was employed doing the Vaccine Work. Exhibit E attached hereto.

NICK WARNE, VP OF RESEARCH AND DEVELOPMENT, PFIZER:
"If we decide that we want to come up with our own proprietary lipids, to get away from let's say the PEGylated lipid which may or may not be related to anaphylaxis in some of the anaphylaxis reactions we're seeing in like 10 per million cases or something like that, then, so we're going to go and do our own work with the research guys, we don't necessarily feel, because we've done the work, that BioNTech should get full access to that, that should be our intellectual property. That should be owned fully by Pfizer, so that's what Patrick's doing with the pure team."

ME:
"I was in a meeting in a hall room the other day just doing stability manager stuff, with my coworkers on the analytical team, and I mean, we were talking about reporting adverse events outside the office, and"

NICK WARNE, VP OF RESEARCH AND DEVELOPMENT, PFIZER:
"Yeah, we're all supposed to do that."

ME:
"They advised me not to."

NICK WARNE, VP OF RESEARCH AND DEVELOPMENT, PFIZER:
"Okay, well you should. Yeah, if you know somebody who's taken a Pfizer drug and has had an adverse event, it's company policy that you should report it."

ME:
"So, what should I say to my coworkers?"

NICK WARNE, VP OF RESEARCH AND DEVELOPMENT, PFIZER:
"Say 'you know what, it's company policy, I got trained on this, you got trained on this, you really should do it.' Absolutely."

ME:
"Well, so, if they're not, essentially. What should we do about that?"

NICK WARNE, VP OF RESEARCH AND DEVELOPMENT, PFIZER:
"Yeah so, so that's an interesting one, because on the one hand. Okay so, do-do you nark on them? Right? Is the question, I get it. I think to some extent, it has to be their responsibility, to report the adverse event or not. If we, if we try to, encouraging is fine. So, I think encouraging somebody to follow the policies is what we should be doing. But it's not necessarily, the compliance, the lack of reporting it, is on them, it's not on you, it's not on me. So if they are not compliant, it's sort of their responsibility, it can't be ours to make them compliant. Now we can encourage them, that's what all the training is for, but we

can't necessarily enforce it, it's really hard. And I don't think there's any way to do it anyway, you know, legally or whatever. So, all I can say is, it is company policy, if you could- if they were sitting here I would encourage them to do that, but I can't tell them to do that. Actually, I think I can, but peer to peer you can't tell them to do anything right? All you can do is encourage them."

ME:
"Another thing that I saw in the vaccine world, I was on Twitter, and I saw that video of Albert actually, I forget what news site it was, and he was, uhm, you know Mr. Bourla, was doing an interview on talking about misinformation, I don't know if you saw it?"

NICK WARNE, VP OF RESEARCH AND DEVELOPMENT, PFIZER:
I heard about it, I didn't watch it, but I heard he had some strong words.

ME:
"I thought so too."

NICK WARNE, VP OF RESEARCH AND DEVELOPMENT, PFIZER:
[laughs] "And I'm like, uh, did you really need to say that. But he's the boss."

ME:
"He talked about dark organizations."

NICK WARNE, VP OF RESEARCH AND DEVELOPMENT, PFIZER:
"Yeah he knows more about that than I do, But I know what we- the only thing I know, is that there have been, because I don't follow a lot of this stuff, I read just a few things. But I know that the organizations working with Russia and with China, specifically, have been spreading a lot of disinformation, and misinformation on whatever media you can find you know? To diss the Pfizer vaccines on a global basis. And, and I'm not shocked because they've been doing this in other areas for a long time."

ME:
"Do we know who they are?"

NICK WARNE, VP OF RESEARCH AND DEVELOPMENT, PFIZER:
"Somebody probably does, you know, but if they're sitting in Russia there's nothing you can do about it."

ME:
"Dark organizations, I-I"

NICK WARNE, VP OF RESEARCH AND DEVELOPMENT, PFIZER:
"I mean Mark Zuckerberg can shut them down off Facebook once he figures out who they are, but then you run into the question of censorship, so it's tricky, it's very tricky."

ME:

"So, is it an international dark organization, or domestic?"

NICK WARNE, VP OF RESEARCH AND DEVELOPMENT, PFIZER:
"I have no idea; I have no idea."

ME:
"You don't know? See I want to know, I want to know who these people are, because I mean, I'm in this fight, you know? I can't stand people are like, I mean, I have friends who won't take the vaccine and I tell them 'It's safe bro' like come on, you know? But, you know, they have their freedoms."

NICK WARNE, VP OF RESEARCH AND DEVELOPMENT, PFIZER:
"This is the issue we run into with Facebook, which I don't deal with Facebook but, you know, as soon as they start taking stuff down, they become a censor."

NICK WARNE, VP OF RESEARCH AND DEVELOPMENT, PFIZER:
"And who's going to become the censor? You know google gets dinged because they're taking down misinformation, are they taking down the wrong kind of misinformation and now it gets to political speech and free speech, some Republican congressman gives a talk and they yank it off your YouTube because they don't like what he has to say. He's a congressman, you know twitter takes Trumps twitter account because they don't like what he has to say, this is the fricken President of the United States and we shut him down, but we're going to have the President of Iran give him a twitter account, you know? So, they've got issues and censorship-so it's tough, it's tough. Because we know the organizati-now if they make lies and we can get our hands on them legally and shut them down we will."

ME:
"Do we have product in China?"

NICK WARNE, VP OF RESEARCH AND DEVELOPMENT, PFIZER:
"Lots of product in China."

ME:
"So why would they be, you know, why would they have disinformation people coming at us then?"

NICK WARNE, VP OF RESEARCH AND DEVELOPMENT, PFIZER:
"They want to play it both ways, the Chinese are, and I got to scoot at five, and I gotta get going. The Chinese want it both ways, they want Western medicines that they will rip off, because the Chinese do not honor US patents, or anybody's patents. So, they will rip us off, but it's a big market, we want to sell some drugs to them because we're going to make some money and so we've-What we'll end up doing, is we will take product to China, and we will actually export manufacturing at some points to China but never the top-the first generation, we'll always give them our older processes because we know they'll rip them off."

ME:
"I see."

NICK WARNE, VP OF RESEARCH AND DEVELOPMENT, PFIZER:
"Yeah, but it's a big market, it's a billion person market."

22. Below is a transcript of a conversation I was a part of while employed doing the Vaccine Work with the following individuals: Bakul Bhatngar, Head of Pediatric Formulations, Cassie Nedved, Senior Associate Scientist, Formulation and Christopher Onyekpe, Senior Associate Scientist, Formulation | DPDD. Exhibit E continued

CHRISTOPHER ONYEKPE, SENIOR ASSOCIATE SCIENTIST, FORMULATION, PFIZER:
"So, Bakul, why are we looking at this covid level concentrations [unintelligible]?"

BAKUL BHATNGAR, HEAD OF PEDIATRIC FORMULATIONS, PFIZER:
"Why? Ah great question! So, we are, for the covid pediatric formulations there are two compositions, and we've seen some effect of, we know that when the, when you go from .1mg per mil which is the adult presentation, down to .033 or .01mg per mil we start seeing unusual behavior in terms of size, percent of capsulation. So, we've done some prework around trying to mitigate some of that. Now uhm, what we eventually decided was before we putz around with covid too much, anything that we do with COVID you need to understand that it connects us back to the mothership in Germany with BioNTech, from IP perspective, so, stays within the room. From an IP perspective, we said, we have a lot more freedom to operate, and work with flu. So, the idea was, we want to set up a study that covers a broad range of concentrations. Starting at .5mg per mil RNA down to .1, to .01, and let's do something out of the box going down to .001. And obviously, our assays are designed for a specific formulation, developed, qualified, validated. And so– but the thought was to cover a broad design space of formulations, under– to see what we can see, and hopefully that broad understanding provides us some, [exhales] I'm exhaling but breathing room, from the perspective of intellectual property. So, these experiments are designed from that perspective to cover a broad design space, now we know there is going to be an impact on size, there's going to be an impact on encapsulation, integrity not so much. But, at the lower concentrations we are running into the lack of reliability of the method itself. And so—"

CASSIE NEDVED, SENIOR ASSOCIATE SCIENTIST, FORMULATION, PFIZER:
"And for like, therapeutic development it's necessary to have lower doses because of its toxicity, right? If we're going into that realm of medicine, I don't know if they are."

BAKUL BHATNGAR, HEAD OF PEDIATRIC FORMULATIONS, PFIZER:
"So, great question."

ME:
"And adverse events."

CASSIE NEDVED, SENIOR ASSOCIATE SCIENTIST, FORMULATION, PFIZER:
"I've read articles on that recently."

BAKUL BHATNGAR, HEAD OF PEDIATRIC FORMULATIONS, PFIZER:
"Yeah so, all of our work in general has been on the modRNA right? Now, Pfizer from a flu perspective is also looking at saRNA of which you've analyzed samples, and so that is supposed to work at much lower concentrations. So, the thought in this craziness as we design this [unintelligible?] we said let's go out do concentrations that we think would be covering saRNA as well. It's mind boggling, you know? So, what I want to tell you is there's a method to the madness. We're not just sitting there and saying, 'well let's try this.' But I think that we, we learn some things from COVID but we don't want to learn too much, yet and [unintelligible] we will develop that database or that, some of the lessons learned with flu and go back and apply them to COVID so that we can buy the– so that we can have the relief that we need from an IP perspective. There are too many strings attached to BioNTech. I'm sure Ramin talks about it, BioNTech is a–

ME:
"I've heard Oskar talk about it actually, where we talk to them when we don't want to be, because they have, like we're sharing half the profits with them right now, or something like that."

BAKUL BHATNGAR, HEAD OF PEDIATRIC FORMULATIONS, PFIZER:
"Yeah, I mean, no matter what we do, they will own things, we will be co-inventor. When it comes to COVID we will be co-inventor, but they'll own everything. And that's not, you know, maybe that's not a fair business practice, but maybe this is something that need to have been thought more carefully by lawyers last year. But last year we were already building the plane as we were starting to fly it."

CHRISTOPHER ONYEKPE, SENIOR ASSOCIATE SCIENTIST, FORMULATION, PFIZER:
"So, when it comes to the IP, how much of the initial formulation, or just the IP in general came from BioNTech, when did Pfizer come in? At what point does Pfizer come in?"

BAKUL BHATNGAR, HEAD OF PEDIATRIC FORMULATIONS, PFIZER:
"Good, good question, so we have to understand that BioNTech is not a formulation company, they are an mRNA company. So, they didn't know how to formulate these lipid nanoparticles they say, 'ha ha we have the best mRNA on the planet!' may not be true, others may have slightly better, more stable mRNA as well. But they went to Acuitas. Acuitas is a Vancouver based company founded by a professor from the University of British Columbia.

CHRISTOPHER ONYEKPE, SENIOR ASSOCIATE SCIENTIST, FORMULATION, PFIZER:
"They made the lipids [unintelligible mumbling]."

BAKUL BHATNGAR, HEAD OF PEDIATRIC FORMULATIONS, PFIZER:

So, they, they had a library of lipids, and so they formulated, they developed these lipid nanoparticles for BioNTech. So BioNTech licensed that formulation from Acuitas. So, it's not just a Pfizer-BioNTech relationship, but there is Pfizer-BioNTech-Acuitas. Now, that's, that's just covid, when it comes to flu, it becomes a little bit different, where we have more flexibility, where whatever we invent, I think we either get the license or we will own it. Then you jump to something called shingles, which, Bob is, I think leading. And so shingles is a slightly different flavor, where I think we are going to work on shingles with, guess who? BioNTech. Okay, but we will be working, maybe separately with Acuitas, the challenges are changes by the day. It's a very complicated IP landscape.

ME:
"Another question, that is kind of family related, my cousin, she took the vaccine but her kid is five, and my, little nephew is five, and she doesn't want to vaccinate her kid because she is concerned about fertility issues. So I was just curious on your thoughts about fertility and long term events that could happen from this."

BAKUL BHATNGAR, HEAD OF PEDIATRIC FORMULATIONS, PFIZER:
"Yeah, and I would say, we don't know. You know, when we sell a drug there's a list of benefits, and there's a long list of side effects. Just reading the list of side effects, many times, scares us from taking the drug. My cousin who's not vaccinated was telling me about, she said 'there's so many women in my group, book reading club or other circle, who've been complaining about irregular periods. Okay, so I asked around in my circle of female friends, I'm a gay man, so I have a ton of female friends, and I said, 'tell me hey,' and this is kind of funny, 'uh, how are your periods?', 'well I don't have periods anymore I got everything removed a few years ago' I said whoa I didn't realize that! But, but the last year has been so stressful, working from home and other things that irregular periods are a norm. So now how do we de-convolute that? So this is such a good question, I think we don't have the data yet, to understand."

ME:
"Well so, are we gathering the data on that? Like, long term, where we're going to be able to check?"

BAKUL BHATNGAR, HEAD OF PEDIATRIC FORMULATIONS, PFIZER:
[Stammers] "I think there are two ways of finding out about it. One is to, look at the, the clinical study protocol, for the, the clinical study, to see how long they plan on monitoring the vaccinated folks, vaccinated subjects, and also I think the list of what they're looking at—"

ME:
"Didn't they vaccinate the control group, too? Because they saw the, the benefit outweighed the risk?"

BAKUL BHATNGAR, HEAD OF PEDIATRIC FORMULATIONS, PFIZER:
"I think eventually they had to. I think from an ethical perspective, I think eventually the control group which wasn't vaccinated, I think they gave, they let, I think, I think, they

don't know what they got, or didn't. But I think there was a question about an ethical obligation of, shouldn't the control group get the vaccine? So, there was talk about this in the first FDA hearings in December, so I know this question was asked, there was a response to it. But I think it's an ethical question on, about, but if you vaccinate them, they drop out of the study. Right, and so then you don't have the dataset that you're trying to generate. And don't they deserve to be vaccinated? Oh my gosh yes! With-with high-risk variants coming up or that, have appeared on the scene, delta, you know, you want to make sure they feel that they're also healthy and safe. That experiment in the real world is occurring, but we are not track-we are not tracking it as we would be in a controlled study. And then some people may say is 'but we should be tracking it' and we are not tracking it."

ME:
"Why not?"

BAKUL BHATNGAR, HEAD OF PEDIATRIC FORMULATIONS, PFIZER:
"I-I-I-I don't know, you know, I think it starts to maybe, encroach on people's freedoms and that how, vision of freedom or definition of freedom is different in different countries you know? Freedom in China vs. freedom in the US is a completely different thing. Uhm."

ME:
"Right, we have the constitution for that."

BAKUL BHATNGAR, HEAD OF PEDIATRIC FORMULATIONS, PFIZER:
"Yeah, so we are not, and we cannot force people. But I think there are states like California where they are saying, even when you're in a store, you wear a mask. It's insane. Okay?"

23. Additionally, on October 14, 2021, I informed the Project Veritas team that I was working with Pfizer vials and was examining them under a microscope. I provided these vials to Project Veritas. Shortly after doing so, Dr. Carrie Madej appeared on the Stew Peters Show discussing having received, through an established chain of custody, vials of the Pfizer COVID-19 vaccines. In the video, which was released on October 20, 2021, Dr. Madej states that Project Veritas is working with a Pfizer Whistleblower which is me. Despite her statements that Project Veritas would soon be releasing a story, nothing was ever released by Project Veritas. Dr. Madej describes in detail how the Pfizer COVID-19 vaccines have ingredients and characteristics that were never disclosed by Pfizer to the general public, which include, but are not limited to, parasitic agents and nanotechnology with transhumanistic purposes. See Exhibit F attached hereto. SHOCKING: Dr. Carrie Madej Releases FIRST LOOK at Pfizer Vial Contents - Forbidden Knowledge TV

24. On or about November 4, 2021, I had another conversation with Nick Warne, the Vice President of Research and Development of Pfizer during which he stated that he had "gamed the FDA" when I asked him about the childhood vaccine approval. Nick Warne explained to me that there was a different buffer system that was used for the approval verses what Pfizer obtained the clinical data from. The clinical data obtained was done using PBS, where the approved vaccine uses Tris. Pfizer and the FDA agreed upon allowing this approval as a "bioequivalence" loophole. This is fraudulent, as the storage temperature of the vaccine was also different and would certainly have impacted the formulation. If there was informed consent, parents of 5- to 11-year-olds after the "approval" would have increased vaccine hesitancy and not injected their child with a formulation that had no clinical data specifically done. See Exhibit G attached hereto.

25. It is currently being reported that roughly 17 million people worldwide have died from being vaccinated with the COVID-19 injections, likely with many more in the years to come. This is equivalent to the number of deaths that happen in global wars. This number is far beyond an amount that demands the removal of these injections from all global markets. See e.g. Exhibit H attached hereto. Study shows COVID-19 injections have so far caused a staggering 17 MILLION deaths – NaturalNews.com

26. Given that it is being reported 17 million people have died from being vaccinated with the COVID-19 vaccines, this is mass-genocide under the guise of a vaccination campaign.

27. "Christine Massey" is an independent researcher out of Canada who has requested from over 220 government and health organizations worldwide, including the CDC, for any evidence of a purified and isolated form of COVID-19 or "SARS-COV-2" from anyone, anywhere on earth, ever with no response. The fact that Christine Massey has gone to so many lengths to ask for the purification and isolation of the virus called "SARS-COV-2" and no organization is able to prove this to be true shows that governments around the world are hiding an experimental injection for a pathogen that is not proven to exist except in computer models. Ultimately, virology is a fraud as well as "germ theory" as a whole. Christine Massey has asked these same government health institutions around the world to provide evidence of the purification and isolation of viruses such as "H1N1, Ebola, HIV and others." The governments around the world are guilty of mass genocide and influencing fear-mongering through the media on the general public worldwide. See Exhibit F attached hereto.

28. Christine Massey has also done FOIA requests to the CDC which took over 6 months to give her an explanation, which they never did for proof of contagion of any "coronavirus" or "influenza" virus that causes respiratory illness. Here is a direct quote from the emails from Christine to the CDC: I've been waiting for 4 months for a response to a request for proof of contagion of respiratory illnesses that are claimed by the CDC to be caused by "coronaviruses" or "influenza viruses" (CDC FOIA #23-00999). Please provide an update or a response. All of the past communications are attached."

29. All of Christine Massey's FOI information can be found at her website with this link : https://www.fluoridefreepeel.ca/fois-reveal-that-health-science-institutions-around-the-world-have-no-record-of-sars-cov-2-isolation-purification/. See Exhibit I attached hereto.

30. Dr. Mark Bailey is a microbiology, medical industry and health researcher who worked in medical practice, including clinical trials for two decades. In 2016, Mark left clinical practice due to the dissatisfaction with the allopathic medical model. Mark is the author of "A Farewell to Virology (Expert Edition)" and co-author with Dr. John Bevan- Smith of "The COVID-19 Fraud and War on Humanity". See exhibit J and K attached hereto.

31. The End of COVID is an online education series with to end the pandemic paradigm with 15 modules outlining the massive fraud of virology, contagion, and more. This resource can be found at this link : https:// theendofcovid.com/learning-center/ . See Exhibit L attached hereto.

32. Project Whistleblower is a documentary of a tell all of my testimony, outlining a large portion of all of these exhibits as well as my experience working for James O'Keefe and other colleagues from Project Veritas and O'Keefe Media Group, who are partly responsible for this information not coming out sooner. See Exhibit M attached hereto.

Here is a quote from Dr. David Martin made in 2021"This was an intentional bio-weaponisation of spike proteins to inject into people to get them addicted to a 'pancoronavirus' vaccine. This has nothing to do with a pathogen that was released...This is about getting people injected with a known to be harmful S1 spike protein...The tragedy is we are in a world where we have hundreds of millions of people being injected with a pathogen-stimulating computer sequence...which is being sold under what the patent office, what the medical profession, and what the FDA and its own Clinical Standards would not suggest is a vaccine, but by using the term we actually are now subjecting hundreds of millions of people to what was known to be, by 2005, a biological weapon." Dr David Martin, 12 July 2021

33. September 21, 2023,an article was published by the Connecticut Centinal by a Staff Writer who has been provided communications about "J O" being James O'Keefe and a Project Veritas Board member communicating about big pharma lying to the public about the safety of the COVID-19 vaccines. This information is on a hard drive with Project Veritas, and potentially Matt Tyrmand according to Exhibit M Project Veritas.
A USB drive will be provided with the evidence from these exhibits. M stand for my code name as a whistleblower source, "Microscope." See Exhibit N attached hereto. Link:https://connecticutcentinal.com/breaking-project-veritas-had-information-in-2021-big-pharma-was-lying-on-covid-vaccine-safety-data-okeefe-told-board-member-never-published/

34. Accordingly, the United States government, Pfizer, Moderna, Johnson & Johnson, Big Pharma, the World Economic Forum, the mainstream media, are complicit in the worldwide COVID-19 fraud and thereby stands accused of reckless criminality, including human rights abuses, crimes against humanity, democide, acts of terror and mass murder.

35. It is my professional opinion that the COVID-19 Vaccines should be removed IMMEDIATELY from the market and public consumption. It is also my professional opinion that all other vaccines should be investigated for safety and efficacy.

[AUTOGRAPH PAGE FOLLOWS]

Justin Leslie

Connecticut State of the Republic

County: Fairfield ss. Brookfield

The undersigned, being duly sworn, deposes and says:

I, Justin Leslie, declare under the penalty of perjury of the laws of the United States of America, and state upon personal knowledge that:

I am an adult of sound mind, over twenty-one years of age, and declare that the information herein is true, correct, and complete and that I have voluntarily affirmed this Affidavit based upon my own personal knowledge, training, experience and expertise, and under the penalty of perjury of the laws of the United States of America.

SUBSCRIBED AND SWORN TO BEFORE ME on the ____ day of March 2024, to certify which witness my hand and official seal.

Notary Public
State of Connecticut

COLLEEN Q. GALBRAITH
NOTARY PUBLIC
My Commission Expires 3/31/26