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17	DEWAYNE JOHNSON,	Case No. CGC-16-550128
18	Plaintiff,	PLAINTIFF'S OPPOSITION TO MONSANTO COMPANY'S MOTION FOR
19	v.	NONSUIT
20	MONSANTO COMPANY,	Hon. Suzanne R. Bolanos
21	Defendants.	
22		Department: 504 Hearing Date: July 30, 2018
23		Hearing Time: 9:00 a.m. Trial Date: June 18, 2018
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#### I. INTRODUCTION

Mr. Johnson has produced at trial the same expert testimony that Judge Karnow considered in rejecting Monsanto's Sargon motions and substantially more evidence with respect to liability and punitive damages than that considered by Judge Karnow in ruling on the summary judgment motion. Monsanto's current motion is simply a rehash of its failed motions infused with gross misrepresentations of the record and should likewise be denied. With respect to causation, Monsanto's motion is inappropriate as it attacks the admissibility of Plaintiff's experts and not whether their actual opinions are sufficient to support a jury verdict. There is no dispute that Plaintiff's experts are eminently qualified to offer causation opinions and did in fact opine that glyphosate based herbicides (GBH) causes non-Hodgkin Lymphoma (NHL) and caused NHL in Mr. Johnson. 5/17/2018 Order re: Sargon and Summary Judgment, (SJ Order) at 7. ("credentials of the expert[s] are unassailable").

Plaintiff's experts considered the whole of the scientific data in coming to their opinions that GBHs cause NHL as required by California law. It is essential that the "body of studies be considered as a whole." *Cooper v. Takeda Pharm. Am., Inc.*, (2015) 239 Cal. App. 4th 555, 589–90. Here, in addition to the epidemiology, there is strong biological plausibility that GBHs cause NHL. Under such circumstances, the Federal Judicial Center's Reference Manual on Scientific Evidence (3rd. Ed.) (Reference Manual) instructs:

... In applying the scientific method, scientists do not review each scientific study individually for whether by itself it reliably supports the causal claim being advocated or opposed. Rather, as the Institute of Medicine and National Research Council noted, "summing, or synthesizing, data addressing different linkages [between kinds of data] forms a more complete causal evidence model and can provide the biological plausibility needed to establish the association" being advocated or opposed. P. 20.

In addition to offering more than sufficient evidence that GBHs cause NHL, Dr. Nabhan and Dr. Sawyer properly conducted a "a differential diagnosis to "rule in" and "rule out" other possible causes of a disease in arriving at an opinion that Mr. Johnson's use of GBHs was a substantial cause of Johnson's NHL. *Cooper*, 239 Cal. App. 4th at 578.

The evidence presented at trial is more than sufficient to support an inference that Mr. Johnson's NHL was caused by Monsanto's products, and is the only logical inference. Mr. Johnson

was diagnosed with non-Hodgkin Lymphoma (NHL) after over two years of spraying dangerously high levels of RangerPro with a power washer that covered his face and clothing with the carcinogens in the RangerPro formulation. This exposure is not Mr. Johnson's fault. July 26 Tr. at 3695:2 (Mr. Lombardi stating "I'm not blaming Mr. Johnson"). Defendant agrees that Mr. Johnson "went well beyond what the label requires" in trying to minimize his exposure. July 9 Trns. Tr. at 1502:15-16. As a safety conscious employee, Mr. Johnson relied on the product label and safety data sheets for RangerPro. July 23 Trns. At 3230:14-16. Unfortunately, Monsanto not only failed to warn about the risk of NHL, but was grossly inadequate in instructing users about how to minimize exposure to its product. 26 Trns. at 3661:17-22. Mr. Johnson's use of a hydraulic nozzle, however, created "a dangerous aerosol." *Id.* at 3694:20-22. Unfortunately, Mr. Johnson "didn't know any better" because Monsanto never bothered to tell him. *Id.* at 3695:1.

Had Monsanto informed Mr. Johnson that GBHs cause cancer, he would not have sprayed the product. Mr. Johnson stated "It's just unethical. It's not what you would do. It's wrong. I have children that go to school, and I've been in schools, and people just don't deserve that. They deserve better." July 23 Tr. at 3235:2-5. Mr. Johnson twice called Monsanto asking if the RangerPro was causing his cancer and Monsanto never called him back. P-Exhs. 332, 334. Fearing that RangerPro might be a cause of Mr. Johnson's NHL, Mr. Johnson's treater, Dr. Ofodile, wrote a letter in March 2015, to the school board asking that Mr. Johnson not be required to spray RangerPro at work. *Id.* at 3154:6-16. That letter did not have the desired effect and in 2016, Mr. Johnson ultimately had to simply refuse to continue spraying. *Id.* at 3236:4-13.

Mr. Johnson's story is tragic and could have been prevented if Monsanto actually showed a modicum of care about human safety. Unfortunately as Judge Karnow ruled, the evidence supports the conclusion that "Monsanto has long been aware of the risk that its glyphosate-based herbicides are carcinogenic, and more dangerous than glyphosate in isolation, but has continuously sought to influence the scientific literature to prevent its internal concerns from reaching the public sphere and to bolster its defenses in products liability actions." SJ Order p. 45. As Dr. Kirk Azevedo testified, Monsanto's Vice President told him "we're about making money. So get it straight." Azevedo Depo. at 50:7-51:25.

#### II. LEGAL STANDARD

Nonsuits are disfavored in jury cases and courts have taken a very restrictive view of the circumstances under which it is proper to take a case away from the jury. "[A] nonsuit in a jury case may be granted only when disregarding conflicting evidence, giving to the plaintiffs' evidence all the value to which it is legally entitled, and indulging every legitimate inference that may be drawn from the evidence in the plaintiff's favor, it can be said there is no evidence to support a jury verdict in their favor". Elmore v. American, (1969) 70 Cal. 2d 578, 583 emphasis added); Alpert v. Villa Romano Homeowners Ass'n, (2000) 81 Cal. App. 4th 1320, 1328; County of Kern v. Sparks, (2007) 149 Cal. App. 4th 11, 16; Carson v. Facilities Develop Co., 2 (1984) 6 Cal. 3d 830, 839 (the inferences need not be the only possible inferences, nor even the strongest inferences; it is enough that there is "some substance to plaintiffs evidence upon which reasonable minds could differ.") Simply stated, nonsuit is an extraordinary remedy with a very high standard and should be denied here where Plaintiffs have presented more than sufficient evidence to support a verdict on all causes of action and punitive damages.

#### III. ARGUMENT

# A. Plaintiff Has Offered Sufficient Evidence For a Jury to Find that RangerPro was a Substantial Cause of Mr. Johnson's NHL.

Each argument offered by Monsanto with respect to causation has already been rejected by Judge Karnow. The issues raised by Monsanto are all foundational issues with respect to the admissibility of the experts' opinions and should not be raised in the context of the sufficiency of the evidence. Once the expert opinion is deemed admissible and presented to the jury, the jury determines the credibility of the expert opinion and not the basis for its admissibility. Here, four highly qualified experts offered admissible opinions that GBHs cause NHL, and two of those experts specifically looked at Mr. Johnson's case and determined that Ranger Pro caused his NHL.

Judge Karnow did indeed make a determination that Plaintiff's experts' opinions are sufficient to support a finding of general and specific causation, holding "Monsanto's motion for summary judgment based on causation turns on the admissibility of Johnson's experts. As discussed above, most of the opinions of Johnson's causation experts are admissible. These suffice as evidence of both general and specific causation." SJ Order at 38. Plaintiff's experts gave the same testimony at trial that

 was considered by Judge Karnow with respect to summary judgment. Defendant goes to great lengths to avoid citing California law or Judge Karnow's directly controlling order. Defendant also misleadingly cite Judge Chabbria's opinion<sup>1</sup>, applying federal law, by omitting the conclusion that "the plaintiffs have presented evidence from which a reasonable jury could conclude that glyphosate can cause NHL at human-relevant doses. Monsanto's motion for summary judgment is denied." *In re Roundup Prod. Liab. Litig.*, No. 16-MD-02741-VC, 2018 WL 3368534, at \*36 (N.D. Cal. July 10, 2018).

Under California law, to create a jury question on cancer causation "[t]he plaintiff must offer an expert opinion that contains a reasoned explanation illuminating why the facts have convinced the expert...that it is more probable than not the negligent act was a cause-in-fact of the plaintiff's injury." *Cooper*, 239 Cal. App. 4th at 578. Furthermore "Under the applicable substantial factor test, it is not necessary for a plaintiff to establish the negligence of the defendant as the proximate cause of injury with absolute certainty so as to exclude every other possible cause of a plaintiff's illness, even if the expert's opinion was reached by performance of a differential diagnosis." *Id.* at 578.

Under California law, causation is not segregated into concepts of "general" and "specific" causation, as Monsanto suggests. Rather, the only causation element that a plaintiff must show is that the defendant's conduct or product was a "substantial factor" in causing the plaintiff's harm. See CACI 430 ("A substantial factor in causing harm is a factor that a reasonable person would consider to have contributed to the harm. It most be more than a remote or trivial factor. It does not have to be the only cause of the harm."). The substantial factor standard "is a relatively broad one, requiring only that the contribution to the individual cause be more than negligible or theoretical." *Hernandez v. Amcord, Inc.*, (2013) 215 Cal. App. 4th 659, 673. Causation involves "factual questions for the jury to decide, except in cases in which the facts as to causation are undisputed." *Ortega v. Kmart Corp.*, 26 al. 4th 1200, 1205 (2001).

Monsanto's arguments related to causation are premised entirely on misrepresentations, an utter disregard of the opinions of Plaintiff's experts, and the misapplication of the nonsuit standard.

<sup>&</sup>lt;sup>1</sup> Judge Chhabria's ruling under federal law is of limited relevance to this case which requires the application of California law. This is particularly true where Judge Karnow considered the same arguments of Monsanto and the same testimony of the experts. Plaintiff does not agree with several aspects of Judge Chhabria's ruling particularly where it deviates from Judge Karnow's opinion.

Plaintiff's evidence is more than sufficient to establish that his NHL was caused by his exposure to GBHs.

## 1. The IARC Monograph Supports Causation

Dr. Neugut, an esteemed oncologist and epidemiologist with forty years of experienced, testified that "I would say that within the scientific and academic cancer community, IARC is recognized as the main arbiter of -- the prime arbiter of what constitutes a carcinogen or a cancercausing agent. I would have trouble even naming a second -- I would have trouble naming a second choice." Tr. at 2550:12-17 (7/18/2018). Dr. Portier also described the IARC process in great detail and explained how it supports a general causation opinion. Tr. at 1718:4-1760:12 (7/12/2018). Dr. Portier rejected Monsanto's arguments that IARC does not consider real world exposure, testifying "Well, of course they do. That's what this chapter is on, and in -- all of the human epidemiology studies are based upon human exposures, which means they're in the real world." *Id.* at 1741:21-24. Dr. Neugut concurs stating "[o]f course it's a real-world carcinogen -- obviously, the epidemiologic studies are relying on how people are really exposed in day-to-day life." 2600:8-2601:21 (7/18/2018).

In fact, the very safety data sheets relied upon by Mr. Johnson require the inclusion of IARC's assessment. (Unfortunately, Monsanto did not include that information while Mr. Johnson was spraying RangerPro.) As explained by Dr. Sawyer "All MSDSes -- that stands for Material Safety Data Sheet -- are required to provide the IARC classification of carcinogenicity, as IARC has been for many years the key agency internationally that determines whether or not a chemical is carcinogenic..." Tr. 3637:2-6. *See also* 29 C.F.R. § 1910.1200, Appendix A. IARC is one "of the most well-respected and prestigious scientific bodies," whose assessments of carcinogenicity of chemicals "are generally recognized as authoritative..." Reference Manual on Scientific Evidence at 20, 565.

Defendant is simply wrong that IARC did not consider the carcinogenicity of glyphosate formulations. IARC explicitly states that it considered human studies looking at "commercial formulations that include glyphosate and other ingredients" and "experimental studies of 'pure' glyphosate and of glyphosate-based formulations." P-Exh. 302. Furthermore, neither Dr. Neugut nor Dr. Nabhan mimicked IARC. Tr. at 2654:2-19 (7/18/2018) (Dr. Neugut reviewed all of the studies in depth after reviewing the IARC monograph); Tr. at 2789:7-2790:18 (7/21/2018) (Dr. Nabhan conducted comprehensive review of the literature.)

### 2. The Epidemiology Supports Causation

Judge Karnow considered Defendant's arguments with respect to epidemiology and rejected them holding that "Johnson's epidemiology experts should not be excluded." SJ Order at 12. In coming to this conclusion, Judge Karnow considered and rejected the exact arguments made by Monsanto in the present motion. Judge Karnow correctly applied California law which holds that it "is generally correct that in many (or even most) instances epidemiological studies provide the best evidence of causation." *Davis v. Honeywell Internat. Inc.*, (2016) 245 Cal. App. 4th 477, 491.

However, it is also proper for experts to rely on "other tools to determine causation" in cases of rare cancer. *Id.; Roberti v. Andy's Termite & Pest Control, Inc.*, (2003) 113 Cal. App. 4th 893, 901 (opinion admissible where expert relied upon animal studies with pesticide and examination of plaintiff). Where the "validity of these studies, and both their strengths and their weaknesses, are subject to considerable scientific interpretation and debate" it is not the court's role to resolve these debates. *Cooper*, 239 Cal. App. 4th at 589–90.

Defendant claims that epidemiology alone must be sufficient to establish causation and that experts can't rely on toxicological data. Judge Karnow and Plaintiff acknowledged that "Johnson's experts do not view epidemiological evidence as dispositive on causation...They conceded that confounding and bias may explain the association found in the epidemiological evidence if the epidemiological evidence were viewed in isolation." SJ at 11. However, it would be entirely improper to view the epidemiological evidence in isolation, and Plaintiff's expert did not do so. *Id.* (Dr. Neugut too does not base his conclusion solely on that epidemiological evidence.)

When asked whether it would be scientifically appropriate to just look at the epidemiology and ignore the animal studies and the mechanistic data, Dr. Portier explained that "Under no condition would it be... it's never good to look at just one set of data" and that its "common practice... it's good practice" to look at all the data. Tr. at 1965:11-1966:7 (7/13/2018). Dr. Neugut concurs. Tr. at 2736:25-2737:17 (7/18/2018). Based solely on the epidemiology data Dr. Portier concluded that that "there's a demonstrated association" and that "causality is reasonable here." Tr. at 1964:1-17 (7/13/2018). Based on all of the data Dr. Portier concluded that "glyphosate is carcinogenic, causing NHL in humans." Tr. at 1994:19-21 (7/13/2018). Dr. Neugut concurred, stating that after applying the Bradford-Hill criteria and factoring in all of the evidence that "there is indeed a causal association"

22.

between glyphosate and NHL." Tr. at 2646:16-23 (7/18/2018)

Defendant claims that there are no statistically significant studies that adjust for other pesticides. This is not true. Judge Karnow already determined Defendant's claim to be false, holding "[w]hile Johnson's experts concede the limitations of the epidemiological evidence, there is at least one study that controlled for other pesticides and still found a statistically significant association between glyphosate and NHL." SJ Order at 7; Tr. at 2700:24-2701:4 (7/18/2018) (Dr. Neugut explaining that Defendant's claim is "not true."). That study is De Roos (2003) and it showed a 2.1 statistically significant increased risk after adjustment for dozens of pesticides. *Id.* at 2736:12-24; Tr. at 1886:9-1894:12 (7/12/2018) (Portier explaining methodology and results of De Roos (2003) study). Judge Karnow further noted that:

Johnson's experts appreciated the risk the confounders could create an unreliable association between glyphosate exposure and NHL but believed, in light of the studies they reviewed and the other information that they considered, that potential confounders were not the cause of the association. Motion, 6 n.16 (citing evidence), 8:9-9:3. And Monsanto has not identified any pesticides that may, in fact, have confounded the data.

SJ Order at 12-13. Dr. Neugut explained at trial that most errors in epidemiology studies push the relative risk down closer to one meaning that the relative risks reported in the studies are actually underestimates of the true relative risk. Tr. at 2584:21-2589:14 (7/18/2018). Dr. Neugut also explained that these errors pushing down the true relative risk are much more of a concern than any potential confounding. *Id.; see also* Tr. at 1965:3-5 (Portier) ("And whereas most of them did a pretty good job with cofounders, some maybe didn't, but I don't think confounders are a big problem in this set of data.");

Although statistical significance is not required to show causation, there are other statistically significant findings for GBHs and NHL, including the meta-analysis. *Id.* at 2682:23, 2683:9; 2685:18-20. As Dr. Neugut explains:

And if you look, all of them are above 1. All of them. That's a phenomenon referred to in causal epidemiology as consistency. They're consistently elevated above 1. Whatever flaws, problems, issues we're all going to raise about these studies, one or the other, no studies are perfect, whatever things each study does, no study is identical. One does

something. One – each study does something differently. Each study — but all the circumstances under which these six studies – some of them control for different things, some of them are done in different populations. Some of them in Scandinavia. Some of them are in America. Some of them in Canada. Some of them are with farmers. Some of them are not. But all of them are consistently above 1, and that's none random.

Id. at 2612:3-18.

Defendants claim "none of the studies show a statistically significant risk ratio that is above 2.0 as required by California law." This is not true. As Judge Karnow explained, California law clearly does not require a risk ratio above 2.0, holding "Johnson's experts discuss epidemiological studies just as one factor in their opinion that glyphosate-based herbicides cause NHL...Cooper does not mandate exclusion of these opinions for this purpose even if none of the studies shows a relative risk of greater than 2.0." SJ Order at 10. Judge Karnow appropriately applied controlling law which holds that "[t]here is no such requirement [for a relative risk of 2.0] in California." *Davis.*, 245 Cal. App. 4th at 493.

In any event there are several studies showing an odds ratio over 2.0, including De Roos (2003). While Dr. Neugut testified that the odds ratio for ever using GBHs from all of the studies combined was about 1.5, he further explained that "if you start to look at dose response of people who are really significantly exposed to glyphosate, got exposed in a more dramatic way, for longer periods of time, for higher doses, they're going to have a significantly higher risk." Tr. at 2617:21-2618-4, 2644:21-2645:1. Dr. Nabhan also reached the same conclusion after his review of the epi studies: "There's a study published by McDuffie and colleagues in 2001. There's another one in 2003 by De Roos and colleagues. There's another one by Eriksson and colleagues that also published in 2008. All of these showed doubling the risk." Tr. at 2825:9-18 (7/20/2018); *Id.* at 2827:15-2830:5 (explaining that Eriksson and McDuffie showed a dose-response); Tr. at 1880:3-1884:24, 1894:13-1898:3 (Portier explaining the dose-response relationship in McDuffie and Eriksson showed over a doubling of the risk for use greater than 2 days per year (McDuffie) and 10 days per year (Eriksson)).

Finally, Judge Karnow explained that "Johnson's experts considered the strengths and weaknesses of the AHS and the strengths and weaknesses of the case control study in reaching their conclusions about epidemiology and causation... This is what was required of them." SJ Order at 13. Further stating that "the absence of dose response in AHS does not foreclose the existence of other data

supporting· a positive dose response finding." *Id.* at 22. Dr. Neugut persuasively explained why exposure misclassification, loss to follow-up and the exponential increase in GBHs during the enrollment period made the results of the AHS study (with respect to GBHs) unreliable. Tr. at 2618:18-2626:13, 2635:8-2640:18 (7/18/2018); Tr. at 1954:3-1959:17 (7/18/2018) (Portier) ("very serious flaws associated" with the AHS study).

#### 3. The Animal Carcinogenicity and Genotoxicity Studies Support Causation.

Again Judge Karnow has decided this issue holding that "[a]s a general matter, Monsanto suggests that it objects to any consideration of genotoxicity or mechanism testimony. This approach is unsupported by Monsanto's citations and conflicts with the Bradford Hill criteria, which require consideration of biological plausibility." SJ Order at 18. Judge Karnow also admitted Dr. Portier's opinion on animal studies because "he concludes the fact that, according to his analysis, glyphosate causes cancer in mammals (i.e., rodents) renders it biologically plausible, under the Bradford Hill rubric, that glyphosate could cause a specific form of cancer, NHL, in humans." *Id.* at 15; *Roberti*, 113 Cal. App. 4th at 901 (opinion admissible where expert relied upon animal studies with pesticide and examination of plaintiff); *Ruff v. Ensign-Bickford Indus., Inc.*, 168 F. Supp. 2d 1271, 1281 (D. Utah 2001) (expert testimony admissible where it is "...based on the genotoxicity and carcinogenicity of these chemicals in animals, the EPA and the International Agency for Research on Cancers classified hydrazines as probable carcinogens posing a significant cancer risk to exposed humans."); Tr. at 2002:10-2003:10 (7/13/2018) (Portier explaining that mechanistic and animal studies support positive epidemiological findings); 2445:6-15 (7/18/2018) (Neugut explaining that toxicology studies in animals are applicable to biological plausibility in Bradford-Hill analysis)

It is entirely misleading to suggest that animal studies are not relevant to humans because they use higher dose. Dr. Portier explained the reason why animal studies use high doses stating:

[t]he reason you do that is because -- what you're interested in, of course, in human populations is much lower exposures. But you're also interested in human risk in the range of 1 in a million to 1 in 100,000 to 1 in 10,000. And we can't use that many animals to get at that type of risk. And so you don't do studies at human exposure levels. You extrapolate them. You take what you see in high doses, and you draw a line or some other technique

to get you into the low region.<sup>2</sup> That's how you estimate human risk. 1806:22-1807:7.

Here the animal and mechanistic studies strongly support causation with respect to the biological plausibility and coherence among the different lines of evidence. There is solid evidence that in addition to other tumors "malignant lymphoma is being caused in these mice by glyphosate." Tr. at 1835:7-9 (7/12/2018). There is also evidence that GBHs are genotoxic in blood cells and lymphocytes. Tr. at 1973:16-25 (7/13/2018). Glyphosate and GBHs have been shown to cause oxidative stress which can operate to promote tumors. *Id.* at 1990:10-1992. GBHs have been shown to promote skin tumors in mice. Tr. at 1857:22-1860:13 (7/12/2018).

Finally, there is clearly a jury issue with respect to who has the correct interpretation of the animal data. Dr. Portier thoroughly explained how the EPA and EFSA failed to follow established guidelines in evaluating this data and was joined by 94 other scientists in a published commentary supporting IARC and Dr. Portier's conclusions. Tr. at 2010:16-2021:2 (7/13/2018).

#### B. Plaintiff Has Established Causation for MF.

Again, Judge Karnow ruled that "I reject Monsanto's argument that there is no scientific basis for Dr. Nabhan to rely on studies that apply to NHL generally in the context of mycosis fungoides.

There is a scientific basis for Dr. Nabhan's opinion - mycosis fungoides is a subtype of NHL." *See also Ruff*,168 F. Supp. 2d at 1285 ("that plaintiffs' expert opinion need not include data showing studies of the exact subtype of plaintiffs' NHL to satisfy their general causation burden."). SJ Order at 23.

Dr. Neugut explained that epidemiologists look at all NHL subtypes as a group:

[E]very disease splits into an absolute panoply of multiple, multiple subtypes. There are more than 60 types of breast cancer. There are more than 60 types of colon cancer. And if you would split and lump into 60

<sup>2</sup> Dr. Portier is referencing the Cancer Slope Factor, which Dr. Sawyer did calculate and came to the conclusion that Johnson's exposure levels were with the range of the exposure levels causing cancer in

animals. In exchange for not referencing the Cancer Sloper Factor at trial, the parties agreed not to "reference, argue or offer testimony that Mr. Johnson's dose/exposure is below or above any threshold

reference dose derived from animal studies." July 24, 2018 Email from Sandra Edwards to Department 504. Defendant's argument that "the level of exposure to glyphosate in the real world is 'very low' is

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actually about even more than 10 million times lower than the quantities that in one day we had to use in animals in order . . . to assess possible carcinogenicity" violates this agreement. Dr. Sawyer would have refuted the defendant's claims by opining that the Cancer Slope Factor demonstrates that the mouse studies, particularly the lymphoma findings, are relevant to Mr. Johnson's exposure levels. (Sawyer Report) Plt.Ex 750 at 145-152.

types every disease, you would know absolutely nothing about any disease if you're going to argue that each one has its own spectrum of causes or outcomes. To some degree, it is true that each one has a unique risk factor or a unique prognosis or a unique treatment, and to some degree one can make universal statements or integrated statements across the -- across the integrated group. We talked about non-Hodgkin's lymphoma as a group. We treat them as a group.

Tr. at 2656:6-21 (7/18/2018). Dr. Nabhan concurs, explaining that it would be impossible to do epidemiology studies for each of the 70 subtypes of NHL and that a scientist should rely on epidemiology studies that look at NHL as a group. Tr. at 2807:22-2809:3 (7/20/2018). In any event, two studies do look specifically at T-cell lymphoma (which is comprised mainly of Mycosis Fungoides). Dr. Nabhan explained that the Eriksson study showed a non-statistically significant odds ratio of 2.29 for T-cell lymphoma. *Id.* at 2828:4-20. Dr. Portier explained that the AHS study, which Defendants rely on exclusively, demonstrated a quadrupling of the risk for T-cell lymphoma and statistically significant tripling of the risk in a lag analysis. Tr. at 2447:10-2449:19 (7/17/2018).

- C. Plaintiff Has Presented Sufficient Evidence Showing That Mr. Johnson's Exposure to Roundup and Ranger Pro Was a Substantial Factor in Causing His NHL.
  - 1. Plaintiff's Experts Testified That Mr. Johnson Had Sufficient Exposure to Roundup and Ranger Pro To Cause NHL.

Judge Karnow has already rules that both Dr. Sawyer and Dr. Nabhan appropriately considered Mr. Johnson's exposure in rendering causation opinions. SJ Order at 25, 28. Monsanto does not challenge that Mr. Johnson was continuously exposed to GBHs from June 2012 through the date of his NHL diagnosis in August 2014. Therefore, Monsanto's sole argument is that "there is every reason to believe that Plaintiff's exposure prior to the onset of his disease was not significant." Def's Mot. for Nonsuit at 10. First, Plaintiff's medical experts offered reams of evidence and testimony detailing the basis for their opinions that Mr. Johnson's substantial exposure to GBHs was a cause of his NHL. Testimony from a medical expert that the plaintiff's exposure to a carcinogen is "almost certainly sufficient" to have caused cancer is sufficient to submit the causation question to the jury. See Sparks v. Owens-Illinois, Inc., (1995) 32 Cal.App.4th 461, 477.

Dr. William Sawyer, a forensic toxicologist, undertook his review of the case in order to specifically determine whether Mr. Johnson's exposure was substantial enough to have caused his NHL.

Hoke Decl. (Sawyer Test. Tr. at 3601:20-3602:8). In reaching his opinions, Dr. Sawyer spoke with Mr. Johnson by telephone and reviewed pertinent medical records, deposition transcripts, published studies, animal studies, and internal Monsanto documents. *Id.* at 3587-3598. Dr. Sawyer testified that 10% of the total exposure dose would have been absorbed through Mr. Johnson's skin each time that he sprayed the GBHs. *Id.* at 3649:8-20. The majority of the absorption would have occurred within the first hour of his exposure. *Id.* 3673:25-3674:6.

Despite Monsanto's claims to the contrary, Dr. Sawyer did compute Mr. Johnson's dose using the available literature and the dermal absorption rate of 10 percent. Sawyer Test. Tr. at 3746:7-19. Dr. Sawyer calculated Mr. Johnson's dose based on days of exposure and miligrams per kilogram per day using a tested model. *Id.* at 3747:2-16. By using this model, Dr. Sawyer was able to specifically compute Mr. Johnson's exposure based on the protective gear he was wearing. *Id.* at 13-19. Dr. Sawyer testified that Mr. Johnson's total exposure was sufficient to have caused his NHL. *Id.* at 3747:13-19; 3791:12-25.

Monsanto's claim that Dr. Sawyer testified that Plaintiff's dose was "less than the average in the peer-reviewed epidemiology studies" is simply not true. After calculating the total exposure levels, Dr. Sawyer opined that Mr. Johnson was "beyond the worst case that I've found in the literature which I used as my basis of calculations." *Id.* at 3674; see also 3596-3597 (testifying that Mr. Johnson was "heavily exposed" at a rate far higher than the individuals in scientific studies). Dr. Sawyer further explained that Mr. Johnson's "Tyvek" suit would have done "very little" in protecting him from exposure to GBHs. The nozzle that he would use would produce a huge aerosol resulting in substantial spray drift. *Id.* at 3663-3664. Furthermore, Mr. Johnson's sweat would have actually created an "immediate diffusion pathway to the skin." *Id.* at 3673:2-11.

When asked whether Mr. Johnson's exposure combined with the 10 percent dermal absorption rate was enough to have caused a carcinogenic response resulting in his NHL, Dr. Sawyer opined: "Yes, absolutely. He is – I can say that emphatically, and base it on the peer-reviewed literature, in that his exposure – his levels of exposure were far higher than that in the literature . . ." *Id.* at 3673:25-3674:16.

Monsanto incorrectly asserts that Plaintiff's experts relied on only "two particular incidents" of exposure in reaching their opinions in this case. Once again, this is a blatant misrepresentation of the

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record; Monsanto did not even bother to cite to trial testimony.<sup>3</sup> In fact, Dr. Sawyer did not even factor the two "incidents" into his exposure analysis lending even more credibility to his opinion that his estimates are actually below Mr. Johnson's true exposure. Based on his experience, education and his review of all of the materials, Dr. Sawyer reached his opinion to a reasonable degree of scientific certainty that Mr. Johnson's NHL was caused by his massive exposure to GBHs. Id. at 3601:9-13; 3606. In reaching his opinions, Dr. Sawyer relied on Monsanto's own official studies as it would not have been possible to replicate Mr. Johnson's' exact exposure amount. *Id.* at 3689; 3691-3692; 3790.

Dr. Chadi Nabhan, Plaintiff's expert medical oncologist, also concluded that Mr. Johnson's chronic exposure to GBHs was a substantial contributing factor to his development of myosis fungoides. Nabhan Trans. Tr. at 2799:4-11 (7/20/2018). He explained that Mr. Johnson had chronic dermal exposure to GBHs including at least two acute high-level exposures. *Id.* at 2867. The epidemiology support a dose-response whereby the more exposure a person gets, the more likely they will develop NHL. Id. at 3027. Based on this information, he was able to conclude that GBHs were a substantial cause to Mr. Johnson's NHL.

## 2. There is Sufficient Evidence That Mr. Johnson's Exposure to GHBs Caused His Cancer In 2.25 Years

Latency is measured from the time of first exposure until the time of diagnosis. Sawyer Trans. Tr. at 3677:4-12 (7/26/2018). Both Dr. Sawyer and Dr. Nabhan agree that the latency for NHL can be much shorter than two years and can vary based on the individual. Sawyer Trans. Tr. at 3781 (7/26/2018); Nabhan Tr. at 2857-2858 (7/20/2018). Plaintiff's experts both provided numerous examples of short latency periods which confirmed their understanding that NHL can be diagnosed

<sup>&</sup>lt;sup>3</sup> The consistent misrepresentations in Monsanto's nonsuit motion is alarming. Monsanto cites to Mr. Johnson missing a "whole season of spraying" to support its argument that Plaintiff failed to specify the amount of exposure prior to the onset of his NHL. Def's Mot. at 10. Monsanto is well aware that this occurred after Mr. Johnson was diagnosed with cancer. Johnson Test. Tr. at 3306:14-3307:23 (7/23/2018). Monsanto states that Plaintiff sprayed for "approximately only one year prior to manifesting symptoms of his MF." Def's Mot. at 10. Dr. Nabhan testified that Mr. Johnson sprayed for over two years prior to his first symptoms of mycosis fungoides. There is no testimony to the contrary other than Mr. Lombardi's arguments. Monsanto cites to Dr. Nabhan's testimony that "it was not until April 2014 that he experienced his first high-level exposure to the formulation" Def's Mot. at 10. Dr. Nabhan specifically testified that he did not remember the exact dates. Nabhan Test. Tr. at 2969-2970 (7/20/18).

within a year of exposure to a carcinogen or other offending hazard. *Id.* at 2855-2859; Sawyer Trans. Tr. at 3676-3677 (7/26/2018). The fact that Mr. Johnson received a very high dosage of GBHs for a shorter period of time, his latency would generally be much shorter. Sawyer Trans. Tr. at 3678-3679 (7/26/2018). Furthermore, the fact that Mr. Johnson's cancer behaved so aggressively would suggest that you would expect a shorter latency. Nabhan Trans. Tr. at 3050 (7/20/2018). Judge Karnow actually excluded the latency opinion of a Monsanto's Expert, Dr. Kuzel, because his opinion that Mr. Johnson's latency was too short was speculative. SJ Order at 36.

It is undisputed that Mr. Johnson's first exposure to GBHs was in June 2012 and he was diagnosed with mycosis fungoides in August 2014. Therefore, the latency for Mr. Johnson's cancer is 2.25 years. Sawyer Trans. Tr. at 3676:8-3677:16. Dr. Nabhan and Dr. Sawyer both discussed the "bell curve" associated with latency periods and both concluded that Mr. Johnson's NHL was caused by his exposure to GBHs. Monsanto cannot cite to any evidence to the contrary. Instead, Monsanto makes the conclusory statement that "Plaintiff introduced no competent evidence on latency" and then proceeds to ignore Plaintiff's evidence on latency. Even if Monsanto disagrees with the opinions of Plaintiff's experts, at this state, "evidence most favorable to the plaintiff must be accepted as true and conflicting evidence must be disregarded." *Miller v. Los Angles County Flood Control Dist.*, (1973) 8 Cal.3d 689, 700.

3. Dr. Nabhan Properly Performed a Differential Diagnosis and Determined That Mr. Johnson's Exposure to GBHs Was a Substantial Contributing Factor to His Development of NHL

Judge Karnow determined that Dr. Nabhan conducted a proper differential diagnosis. SJ Order at 25. Dr. Chadi Nabhan is a board-certified oncologist specializing in the diagnosis and treatment of lymphoma's and leukemias, including Non-Hodgkin's lymphoma. Nabhan Test., Tr. at 2773:10-21; 2776:22-24; 2779:6-12. He has authored over 300 journal articles, abstracts and book chapters relating to cancer with the substantial majority dealing with Non-Hodgkin's lymphoma. *Id.* at 2785:13-2786:4. Dr. Nabhan testified that mycosis fungoides is simply a form of non-Hodgkin's lymphoma. *Id.* at 2780:7-17. As such, it is appropriate to rely upon scientific literature related to NHL generally in reaching causation opinions. Dr. Nabhan explained that NHL is a "large umbrella" and due to the changing nature of classifications and the difficulty in specifically testing each particular subtype, physicians must apply causation evidence to every subtype including mycosis fungoides. *Id.* at 2900.

In reaching his specific causation opinions in this case, Dr. Nabhan reviewed epidemiology, animal studies, toxicology studies, thousands of pages of Mr. Johnson's medical records, correspondence from Mr. Johnson's employer, and relevant deposition transcripts. *Id.* at 2789-2795. Dr. Nabhan also personally met and examined Mr. Johnson. Dr. Nabhan considered the amount of Mr. Johnson's exposure in reaching his opinions. *Id.* at 2831. He considered the amount and duration of exposure as well as the number of reported times that the GBHs would have contacted his skin. *Id.* at 2834-2836.

Next, Dr. Nabhan employed a differential diagnosis in which he considered every possible or plausible cause of Mr. Johnson's NHL. *Id.* at 2841-2842. He considered the known risk factors and causes of NHL including age, race, immunosuppressant therapies, autoimmune diseases, skin conditions, occupation, occupational exposures and viruses. *Id.* at 2842-2852. Dr. Nabhan explained that sun exposure, tobacco, and alcohol are not known causes of NHL and could therefore be excluded. *Id.* at 2852-2853. After properly conducting a differential diagnosis, Dr. Nabhan concluded that Mr. Johnson's only known risk factors were his race (African American) and Roundup exposure. Dr. Nabhan therefore concluded that Roundup was the most substantial contributing factor to Mr. Johnson's NHL. *Id.* at 2853:24-2854:2.

Despite Monsanto's argument to the contrary, Dr. Nabhan did consider whether Mr. Johnson's NHL was idiopathic. Dr. Nabhan testified that because Mr. Johnson was far younger than the typical mycosis fungoides patient this would constitute a "red flag" suggesting to him that there was something behind the NHL. July 20 Trns. 2843:2-2844:19. In other words, Mr. Johnson's cancer was not idiopathic. *Id.* at 2997. Dr. Nabhan is "very certain" that if Mr. Johnson had not been exposed to Roundup he would not have developed mycosis fungoides. *Id.* at 2849:9-21.

Under the applicable substantial factor test, "it is not necessary for a plaintiff to establish the negligence of the defendant as the proximate cause of injury with absolute certainty *so as to exclude every other possible cause of a plaintiff's illness,* even if the expert's opinion was reached by performance of a differential diagnosis." *Cooper.*, 239 Cal. App. 4th at 578. In reaching a specific causation opinion, clearly a medical expert "need not exclude all other possibilities before he or she can express an opinion that the defendant's conduct or product caused the plaintiff's harm." *Cooper,* 239 Cal. App. 4th at 580. It is defendant's burden to proffer "the existence of an alternative explanation,

supported by substantial evidence and not mere speculation..." to defeat Plaintiff's claims as a matter of law. *Id.* Judge Karnow agreed that Dr. Nabhan was not required to rule out every other possible cause to render a causation opinion. SJ Order at 25.

The recent case of *Wendell v. GlaxoSmithKline*, (cited with approval by Judge Karnow, SJ Order at 24) applying California substantive law, is particularly relevant because it involves a rare subtype of t-cell lymphoma. 858 F.3d 1227, 1230, 1236 (9th Cir. 2017). Under *Wendell*, it is not necessary for an expert to "rely on animal or epidemiological studies" for a differential diagnosis to be "found reliable and admissible" particularly in case of rare cancers where it would be difficult to conduct studies powerful enough to create statistically significant results. *Id.* at 1235. In conducting a differential diagnosis one "[a]ssumes the pertinence of all potential causes, then rules out the ones as to which there is no plausible evidence of causation, and then determines the most likely cause among those that cannot be excluded." *Id.* 

Monsanto claims because the causes of t-cell lymphoma mycosis fungoides are mostly unknown (as one would expect with a rare tumor), it is impossible to conduct a differential diagnosis. This is not true. As explained in *Wendell*:

the district court erred when it excluded Plaintiffs' experts' opinion testimony because of the high rate of idiopathic [unknown] HSTCL and the alleged inability of the experts to rule out an idiopathic origin or IBD itself. We do not require experts to eliminate all other possible causes of a condition for the expert's testimony to be reliable. *Messick*, 747 F.3d at 1199. It is enough that the proposed cause "be a substantial causative factor." Id. This is true in patients with multiple risk factors, and analogously, in cases where there is a high rate of idiopathy. ..Moreover, when an expert establishes causation based on a differential diagnosis, the expert may rely on his or her extensive clinical experience as a basis for ruling out a potential cause of the disease. See id. at 1198

*Id.* at 1237. *Wendell* concluded that "[w]ere, as here, two doctors who stand at or near the top of their field and have extensive clinical experience with the rare disease or class of disease at issue, are prepared to give expert opinions supporting causation, we conclude that Daubert poses no bar based on their principles and methodology." *Id.* 

IV. Monsanto's Motion Fails to Consider the Elements for Product Liability Which do not Require Any Evidence of A Safer Alternative Design

Monsanto's Motion for Nonsuit claims that Roundup cannot be found to be defective in design because Plaintiff did not prove that there was a safer alternative design. Unfortunately for Monsanto, California's strict product liability law contains no such requirement. In fact, Plaintiff is pursuing strict liability under California's failure to warn and consumer expectation tests. However, Monsanto fails to discuss the elements of the consumer expectation test because Plaintiff presented ample evidence to support both failure to warn and consumer expectation. Thus, Monsanto's motion must be denied.

California courts specifically exclude consideration of a product's benefits unless the plaintiff pursues a defect claim based on the risk/benefit test. The Supreme Court has repeatedly described two tests for strict product liability design defect:

Design defects appear in products that, although properly manufactured, are dangerous because they lack a critical feature needed to ensure safe use. (Brown, supra, 44 Cal.3d at p. 1057.) We discussed design defects at length in *Barker*, *supra*, 20 Cal.3d 413, establishing two alternative tests for liability. A product design may be found defective if (1) "the product failed to perform as safely as an ordinary consumer would expect when used in an intended or reasonably foreseeable manner" (consumer expectations test) (id. at p. 432), or (2) the risk of danger inherent in the product's design outweighs the design's benefits (risk-benefit test) (ibid.).

Webb v. Special Electric Co., Inc, (2016) 63 Cal.4th 167, 180. Here, Plaintiff is only pursuing a design defect claim under the first test, the consumer expectation test. Under the consumer expectation test, a Plaintiff is not required to "prove that there was a safer alternative design." Sparks v. Owens-Illinois, Inc., 32 Cal. App. 4th 461 (1995). In fact, Judge Karnow explicitly ruled that "Johnson is entitled to an instruction on the consumer expectations test... Monsanto may not defend a claim under the consumer expectations test by relying on the risk-benefit test. Sparks v. Owens-Illinois, Inc., 32 Cal.App.4th 461, 473 (1995); see also Soule v. General Motors Corp., 8 Cal 4th 548 (1994) ('party is entitled upon request to correct, non-argumentative instructions on every theory of the case advanced by him which is supported by substantial evidence."). May 17 Order on Jury Instructions at 5. (emphasis added).

Under these circumstances, the benefits of Roundup are not only irrelevant, they are to be excluded:

...the consumer expectation test applies in 'cases in which the everyday experience of the product's users permits a conclusion that the product's design violated minimum safety assumptions, and is thus defective

regardless of expert opinion about the merits of the design.' A plaintiff may show the objective condition of the product, and the fact finder may use its own 'sense of whether the product meets ordinary expectations as to its safety under the circumstances presented by the evidence.' A defendant may not rebut such a claim with evidence of the design's relative risks and benefits...")

Arena v. Owens-Corning Fiberglas Corp. (1998) 63 Cal.App.4th 1178, 1186 (quoting Soule 8 Cal 4th at 567, 607) (emphasis added). Furthermore Comments K and J are inapplicable as a defense, as "Monsanto does not cite—and the Court cannot find—a California case applying comment k outside the medical context, ... On the contrary, California courts appear willing to apply comment k only where a product is "available only through the services of a physician," Hardeman v. Monsanto Co., 216 F. Supp. 3d 1037, 1040 (N.D. Cal. 2016)

Instead of presenting the elements for the causes of action or tests of product defect that Plaintiff is pursuing, Monsanto attempts to set up a straw man regarding argument of some safer alternative design that does not apply to this case. Monsanto has already admitted, in a Request for Admission entered into evidence in this case, that it did not warn Mr. Johnson of any of the cancer dangers of its glyphosate-based products. July 23 Trns. at 3186:2-13. Knowing that it cannot defeat Plaintiff's failure to warn or design defect claims, Monsanto now attempts to create a test for liability that does not exist here. Plainly, the benefits of Roundup or a safer alternative design are irrelevant.

As discussed below, Plaintiff has presented more than sufficient evidence of the fact that he did not expect Roundup to cause cancer and Monsanto admitted it did not warn of such a danger. July 23 Trns. at 3234:21-25, 3279:1-12 ("Q. Had you seen something, that Ranger Pro could cause non-Hodgkin's lymphoma or cancer, would you have sprayed this product? A. I would never have sprayed that product on school grounds or around any people if I knew it would cause them harm..."). Mr. Johnson also testified that he read the label every time he used the formulation. *Id.* at 3231:3-24. The jury certainly has enough evidence to "use its own sense of whether the product meets ordinary expectations as to its safety under the circumstances presented by the evidence" and whether Monsanto failed to warn. *Arena* 63 Cal.App.4<sup>th</sup> at 1186 (quoting *Soule* 8 Cal 4th at 567, 607). Thus, Monsanto's Motion for Nonsuit must be denied.

### V. Plaintiff Has Presented Sufficient Evidence for Strict Liability Failure to Warn

A plaintiff alleging a failure to warn strict liability cause of action need prove that the defendant either (1) knew of the risks involved at the time of manufacture and/or distribution, or (2) based on the state of scientific knowledge at the time of manufacture and/or distribution, should have known of the risks. Webb, 63 Cal.4th at 180. "Manufacturers have a duty to warn consumers about the hazards inherent in their products. Johnson v. American Standard, Inc. (2008) 43 Cal.4th 56, 64. The California Supreme Court has traditionally imposed strict liability for failure to warn of either known or reasonably scientifically knowable risks of a product. Anderson v. Owens-Corning Fiberglas Corp. (1991) 53 Cal.3d 987, 1000, 1002 ("[t]he rules of strict liability require a plaintiff to prove only that the defendant did not adequately warn of a particular risk that was known or knowable in light of the generally recognized and prevailing best scientific and medical knowledge available at the time of manufacture and distribution."). When adjudicating failure to warn under strict liability principles, "the reasonableness of the defendant's failure to warn is immaterial." Carlin v. Superior Court (1996) 13 Cal.4th 1104, 1112.

# A. The NHL Risks of GBHs Were Known and Knowable by Monsanto Before Mr. Johnson Started Spraying RangerPro

Monsanto contends that a warning requirement was not triggered by the prevailing scientific evidence for Roundup carcinogenicity because 1) the 2015 IARC classification of glyphosate as a probable human carcinogen did not render knowable the risk of cancer; 2) the EPA's review and registration of glyphosate militates against a warning; and 3) Mr. Johnson's injury preceded the IARC classification. Monsanto Brief at 18. Such assertions not only sound in abandoned preemption arguments, but are factually inaccurate. It is well-settled that "reasonably scientifically knowable...refers to knowledge obtainable 'by the application of reasonable, developed human skill and foresight....[t]he actual knowledge of the individual manufacturer, even if reasonably prudent, is not the issue....the manufacturer is held to the knowledge and skill of an expert in the field; it is *obliged to keep abreast of any scientific discoveries and is presumed to know the results of all such advances. Carlin* 13 Cal.4th at 1113, fn. 3 (emphasis added) (quoting *Anderson, supra*, 53 Cal.3d at 1002, fn. 13). The IARC monograph *reviewed* prevailing epidemiological, toxicological, and mechanistic literature associating Roundup exposure with NHL. Evidence presented at trial demonstrated that the data

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reviewed by IARC was known years before the agency classified glyphosate, as testified to by Plaintiff's experts.<sup>4</sup> Neugut, July 18 Trns. at 2614: 17-21, 2617:21-25 ("The point that we should walk away with is that overall, there's a statistically significant increased risk in the 1.3, 1.4, possibly 1.5, range. And that's basically what the case control studies are showing us..." the case control studies range in time from 1999-2008); Portier July 12 Trns. at 1897 (discussing doubling of the risk in epidemiological studies from 2001 and 2008); July 12 Trns. at 1965:9-11 (Dr. Portier's review of the epidemiological literature preceding the IARC decision led him to conclude that a casual association is credible); July 20 Trns. at 2826:18-25 (discussing De Roos (2003) "A. So you will see that it says 2.1. So it doubles the risk."); Portier July 13 Trns. at 1981:14-22 (concluding that Roundup is genotoxic after reviewing over 100 mechanistic studies going back to the 1990s).

Indeed, the toxicologist hired by Monsanto, Dr. James Parry, concluded in the late 1990s and early 2000s that Roundup is genotoxic and can cause oxidative stress, a precursor to cancer based on Dr. Parry's review of toxicological data published throughout the 1990s. P-Exh. 217 at 10, 5 (First Parry Report: "These data indicate that Glyphosate produces oxidative damage in vivo... in vitro evidence of genotoxic effect for Roundup mixture.); P-Exh. 220 at 11 (Second Parry Report: "These studies provide some evidence that Roundup mixture produces DNA lesion in vivo, probably due to the production of oxidative damage."). Dr. Mark Martens, the Monsanto employee who worked closely with Dr. Parry, confirmed Dr. Parry's conclusions. Martens Depo at 65:3-6, 20-25; 72:3-12. Dr. Parry's toxicological findings were also confirmed by Dr. Portier after his review of the animal data spanning from the 1980s to 2010. July 12 Trns. at 1834:16-1837:14, 1863:19-20, 1758:10-11 ("A. glyphosate has the potential to be a promoter of carcinogenesis...[glyphosate] caused the cancer seen in the animals.").<sup>5</sup>

<sup>&</sup>lt;sup>4</sup> Monsanto's awareness of such evidence is discussed in the next section when addressing the negligent failure to warn claim.

<sup>&</sup>lt;sup>5</sup> Dr. Martens confirmed that Dr. Parry was a recognized expert in the field and that the studies reviewed by Dr. Parry employed generally acceptable methods. Martens Depo. at 29:7-9, 238:4-17 ("Q. Was [Dr. Parry] an expert in his field? A. Yes... A. the dosage that was used in these studies that you analyzed, they were following standards and practices that scientists use all over the world, correct? A Yes. O They weren't doing anything abnormal, correct? A No. Q They were following the same practices that scientists follow all over that give us results that we -- that are accepted all over the world, correct? A Yes, insofar they follow the international accepted test guidelines.").

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Given that the Court is obliged to take every favorable presumption fairly arising from the evidence as facts proved in favor of the Plaintiff, nonsuit must be denied when Plaintiff has presented a plethora of evidence from which the jury could conclude that Monsanto should have warned of a cancer risk. Elmore v. American, 70 Cal. 2d 578, 583 (1969). Indeed, Court is not permitted to weigh the evidence and must disregard conflicting evidence such as the EPA's review of glyphosate. Stonegate Homeowners Assn. v. Staben (2006) 144 Cal. App. 4th 740, 745 ("evidence most favorable to plaintiff must be accepted as true and *conflicting evidence must be disregarded.*") (emphasis added). Notwithstanding this, even if the Court were to weigh contrary evidence such as the EPA's classification of glyphosate, the evidence viewed in Plaintiff's favor precludes nonsuit because, to date, the EPA has not reviewed long-term carcinogenicity studies on the formulated product, such as the surfactant, as confirmed by Dr. Sawyer. July 26 Trns. at 3614, 3615:7-10, 3615:11-25 ("Q. [EPA] haven't actually looked at any testing as to carcinogenicity? A. No, it's never been tested."). On this record, nonsuit is wholly inappropriate.

#### Plaintiff Has Presented Sufficient Evidence for a Jury to Conclude that В. Monsanto was Negligent in Failing to Warn

During trial, Plaintiff marshalled a trove of internal Monsanto documents dating back decades which demonstrate that Monsanto personnel were aware, or at the very least, should have been aware of the association between Roundup exposure and cancer. Indeed, as Monsanto's Medical Sciences Lead, Dr. Daniel Goldstein, conceded at deposition, even prior to IARC, a large body of mechanistic studies had revealed the association between Roundup exposure and the biological precursors to cancer such as oxidative stress, of which Monsanto was aware. P-Exh. 315 at 1-2; Goldstein Depo. at 99:1-24 ("You had been dealing with this issue for a while, fair? A. It had certainly come up before, yes."). The jury heard evidence that the epidemiological studies relied upon by Plaintiff's experts were all discussed inside Monsanto at the time of their publication. P-Exh. 282 at 2 (memorandum by Donna Farmer, John Acquavella and Daniel Goldstein stating that "[t]here are now six published studies that arguably associate glyphosate and other pesticides with lymphopoietic cancers...") (emphasis added); Goldstein Depo at 92:22-96:3 (discussion of email from Dr. John Acquavella regarding association between

Roundup and cancer in De Roos (2003) and Hardell (2002)); P-Exh. 309 at 1 (Dr. Acquavella discussing
the results of the McDuffie (2001) paper which found elevated risks of NHL associated with Roundup
exposure; P-Exh. 312 at 1-3 (discussion of results of McDuffie (2001)); P-Exh. 313 at 1-2 (same, Donna
Farmer and John Acquavella celebrating the fact that the reference to glyphosate carcinogenicity was
removed from the paper's abstract); P-Exh. 316 at 1-2 (email regarding "awareness files" sent to Dr.
Goldstein and Dr. Farmer, attaching the Eriksson (2008) paper at the time of its publication; the study
showed an association between Roundup exposure and NHL; P-Exh. 513 at 1 (Dr. Farmer stating that
Monsanto had been aware of the Eriksson (2008) publication for some time). Significantly, Dr.
Heydens admitted that the body of evidence prior to IARC demonstrated that Roundup had
"vulnerabilities" in the areas of epidemiology, genotoxicity and mode of action. P-Exh 294. Dr.
Heydens also admitted in 2015 that the surfactant in the Roundup formulation "played a role" in the
tumor promotion study of George, et al. (2010), discussed at length by Dr. Portier as a basis for his
conclusion that "glyphosate has the potential to be a promoter of carcinogenesis." P-Exh. 366 at 3; July
12 Trns. at 1863:3-20. And, the report submitted to Monsanto by James Parry (discussed above) is also
further evidence that Monsanto knew or should have known that Roundup causes cancer, particularly
since Dr. Parry's conclusions were in regard to the formulated product. P-Exh. 217 (First Parry Report)
P-Exh. 220 (Second Parry Report); Martens Depo at 65:3-6, 20-25; 72:3-12.

Furthermore, Monsanto recognized the impact of the prevailing scientific evidence of Roundup carcinogenicity on regulatory decisions at least five years prior to the IARC classification. P-Exh. 373 at 5 (internal 2010 Monsanto document acknowledging that regulators were growing "increasingly more concerned" as a result of literature indicating an association between Roundup exposure and health risks.) In the same document, Monsanto identifies toxicological studies which have concluded that Roundup poses risks of biological precursors to cancer. *Id.* at 9-10. Monsanto should have known that Roundup can be carcinogenic when its own scientists admit that the formulation – containing the surfactant POEA – used by Mr. Johnson is "hazardous." P-Exh. 383 at 2 ("there are non-hazardous formulations, so why sell a hazardous one?"). Indeed, lead Monsanto toxicologist, William Heydens, admitted in 2015 that the Roundup formulation contains other known carcinogens such as NNG and

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formaldehyde. P-Exh. 357 at 1. Plaintiff's experts testified regarding this aspect of the formulation.
July 18 Trns at 2603:18-25 (Dr. Neugut confirming that formaldehyde is an IARC Class 1 carcinogen);
July 26 Trns. at 3609:21-3610:5 (Dr. Sawyer testifying that the presence of 1,4-Dioxane, ethylene
dioxide and NNG contribute to the carcinogenicity of the formulation: "In fact, ethylene oxide is one of
the most potent carcinogens known to man."). Monsanto's knowledge regarding the carcinogenic
formulation is thus substantiated by Plaintiff's competent expert testimony. Despite such knowledge,
Monsanto refused to conduct a proper long term carcinogenicity study on the formulation, which is why
Dr. Farmer "cannot say that Roundup does not cause cancerwe have not done carcinogenicity studies
with 'Roundup.'" P-Exh. 305 at 1. Such statements evince that Monsanto, at the very least, should have
known that its product could cause cancer.

Monsanto also argues that Plaintiff has not presented evidence regarding the applicable standard of care. However, this is a misstatement of what is required under the law. General negligence, unlike the medical standard of care, does not require the applicable standard of care to be establish by competent expert testimony. Scott v. C.R. Bard, Inc. (2014) 231 Cal.App.4th 763, 787. The Anderson Court made it clear that the applicable standard of care is simply what reasonably prudent manufacturer would have known and warned about." Anderson, 53 Cal.3d at 1002. It is worth noting that Monsanto's citation to Stephen v. Ford Motor Co., 134 Cal. App. 4th 1363, 1367 (2005) is misguided. Stephen is not authority for the proposition that a negligent failure to warn case requires expert testimony regarding the applicable standard of care. Instead, the Stephen court merely reiterated the trial court's finding that "[plaintiff] did not have any expert testimony to show the tire failed as the result of a design defect..." Here, CACI 400 governs, namely the Court will instruct the jury on what the standard of care is and that nothing in this case is so outside common experience that it would require expert testimony; in fact, expert testimony would be prohibited because it would invade the province of the jury to decide the ultimate issue in the case. Ramirez v. Plough, Inc (1993) 6 Cal.4th 539, 546 ("The formulation of the standard of care is a question of law for the court. Once the court has formulated the standard, its application to the facts of the case is a task for the trier of fact if reasonable minds might differ as to whether a party's conduct has conformed to the standard."). If Plaintiff proves

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that Roundup causes cancer, the jury certainly does not need expert testimony to come to the conclusion that defendant's failure to warn of cancer, while representing it as safe, was unreasonable under the circumstances.

Because Plaintiff has presented sufficient evidence that Monsanto was aware of a cancer risk in light of the prevailing scientific literature and its own admissions (discussed above), the trier of fact could properly find that Monsanto was negligent in failing to warn and nonsuit must be denied accordingly.

# C. There is Sufficient Evidence that Monsanto's Failure to Warn Caused Plaintiff's Injury

On the stand, Mr. Johnson testified clearly and unequivocally that he would never have used Roundup had he known it would cause his NHL. July 23 Trns. at 3234:21-25, 3279:1-12 ("Q. Had you seen something, that Ranger Pro could cause non-Hodgkin's lymphoma or cancer, would you have sprayed this product? A. I would never have sprayed that product on school grounds or around any people if I knew it would cause them harm... Q. And if Monsanto had told you at that time that Ranger Pro could have caused cancer, would you have kept spraying their product? A. Absolutely not... Q. Even if Monsanto had said, 'Mr. Johnson, we're not entirely sure, but your cancer might be related to your Ranger Pro exposure," would you have continued to spray the product? A. If they said it might be? Q. Even if they said "We're not entirely, but it might be," would you keep spraying? A. No."). Mr. Johnson also testified that he read the label *every time* he used the formulation. *Id.* at 3231:3-24.

Mr. Johnson's conduct and state of mind *after* his diagnosis are simply irrelevant and cannot be used by Monsanto to claim that a warning would not have been heeded. In fact, when Mr. Johnson was asked by his employer *after diagnosis* whether Mr. Johnson knew that Roundup may be carcinogenic, Mr. Johnson responded: "No, I didn't know that." July 23 Trns. at 3235:21. Similarly, Dr. Ofodile's (Mr. Johnson's treating physician) letter to the school board requesting that Mr. Johnson not be exposed occurred after Mr. Johnson's diagnosis, when the opportunity to heed a warning was long moot. July 23 Trns. at 3154:8-13. Importantly, the letter from Dr. Ofodile was in part prompted by the cautionary principle of preventing repeated exposure to substances that may have contributed to an existing cancer:

"My understanding is that, you know, it is okay and advisable to avoid something that could potentially be -- can exacerbate a condition, even if it's not been conclusive. So, you know, if I had a patient that was exposed to something and I thought may be causing it, my recommendation is to avoid it rather than waiting to see whether or not it truly does cause it later on." *Id.* at 3155:17-25.

Monsanto also suggests that Mr. Johnson's concern for his health after his back-pack leaked, soaking him in Roundup, somehow indicates that a warning would not have been heeded. Monsanto Brief at 20. This makes little sense given that Mr. Johnson would not have even used Roundup, much less been exposed to substantial amounts, if Monsanto had properly warned. Moreover, Mr. Johnson's general health concerns following exposure to a chemical are materially different to knowledge that the product causes a specific type of cancer. For the same reasons, Mr. Johnson's decision to use protective clothing is irrelevant to whether he knew that Roundup could cause mycosis fungoides.

### VI. The Evidence for Punitive Damages is Overwhelming

### A. Monsanto's Conduct Meets the Threshold for Punitive Damages

There is more than sufficient evidence in this case for a jury to find for Plaintiff on his punitive damages claim. "If a company intentionally proceeds with conduct which will expose a person to a serious potential danger known to the company in order to advance the company's own pecuniary interest, punitive damages may be assessed based on a finding that the company has shown a conscious disregard for the person's safety." Ford Motor Co. v. Home Ins. Co., 116 Cal. App. 3d 374, 381–82, (Ct. App. 1981); Boeken v. Philip Morris Inc., 127 Cal. App. 4th 1640, 1690 (2005) (intentionally marketing a defective product knowing that it might cause injury and death is highly reprehensible). Here, Monsanto continues to hide and suppress evidence that there is a potential deadly danger of NHL associated with GBHs and it does so because of its own pecuniary interests.

As an initial matter, Monsanto underestimates the "crux of Plaintiff's case for punitive damages..." *Id.* Certainly, evidence that Monsanto did not conduct the studies suggested by Dr. Parry evinces reckless disregard for human health. P-Exh. 268 ("we are simply not going to do the studies Parry suggests."). Dr. Portier testified that Monsanto only carried out one of Dr. Parry's recommendations: "Q. So of all of Dr. Parry's recommendations asking for affirmative action, only one

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	of them was done in this study? A. Yes." July 13 Trns. at 1997:19-22. Moreover, Monsanto's
	manipulation of the scientific data on Roundup also go to punitive intent, as recognized by Judge
	Karnow when ruling that evidence such as ghostwriting is evidence "that a reasonable jury could find
	amounts to clear and convincing evidence of malice, fraud, or oppression." Johnson Sargon Order at 45-
	46 (The internal correspondence noted by Johnson could support a jury finding that Monsanto has
	long been aware of the risk that its glyphosate-based herbicides are carcinogenic, and more
	dangerous than glyphosate in isolation, but has continuously sought to influence the scientific literature
	to prevent its internal concerns from reaching the public sphere and to bolster its defenses in products
	liability actions.") Id. at 45 (emphasis added). In fact, 6 out of 8 documents that Judge Karnow opined
	the jury could use in deciding punitive intent were admitted at trial, namely: P-Exh. 305 at 1 (Donna
	Farmer stating that "you cannot say that Roundup does not cause cancerwe have not done
	carcinogenicity studies with 'Roundup.'"); P-Exh. 362 at 2 (Dr. Heydens admitting that Monsanto
	ghostwrote Williams (2000) that Monsanto then leveraged with regulatory agencies to support the
	registration of glyphosate <sup>6</sup> "keeping costs down and doing the writing ourselvesrecall that's how we
	handled Williams, Kroes & Munro (2000)") (this occurred during Mr. Johnson's period of exposure); P-
	Exh. 217 (First Parry Report concluding that Roundup is genotoxic and has the potential to induce
	oxidative stress; P-Exh. 220 (Second Parry Report concluding same); P-Exh. 443 (Monsanto also
	ghostwrote the Kier & Kirkland (2012) article, as partly demonstrated by the "manuscript clearance
	form" signed by David Saltmiras); P-Exh. 445 (The plans to ghostwrite Kier & Kirkland (2012) were
	discussed internally by Monsanto). <sup>7</sup>
	Judge Karnow viewed only a limited amount of the evidence supporting punitive damages. The

Judge Karnow viewed only a limited amount of the evidence supporting punitive damages. The Jury has seen much more. In addition to the evidence which Judge Karnow held could support a punitive damages award, Plaintiff introduced testimony by former Monsanto employee, Dr. Kirk Azevedo, who testified that he observed instances when Monsanto's corporate philosophy led him to

<sup>&</sup>lt;sup>6</sup> Such use of Williams is demonstrated by P-Exh. 373, an internal Monsanto PowerPoint presentation which identifies Williams (2000) as an "invaluable asset" that was used for "Monsanto responses to agencies; Scientific Affairs Rebuttals; [and] Regulator Reviews." at 12.

<sup>&</sup>lt;sup>7</sup> The Parties stipulated to the admission of Exhibits 443 and 445 in lieu of playing the video deposition of Dr. Saltmiras during Plaintiff's case-in-chief.

question Monsanto's commitment to safety, such as when a Monsanto CEO told Dr. Azevedo, in
response to the latter's enthusiasm for a sustainable, environmentally-friendly business model, that
"[t]his sort of thing isn't really what we're about. We're about making money, so get it straight."
Azevedo Depo. at 50:7-51:25. This preoccupation with profits over safety is reiterated by documents
introduced through the testimony of Monsanto sales-representative, Mr. Steve Gould, who was
responsible for calculating the costs on Monsanto's Roundup business in California following the 2015
IARC classification, at a time when Mr. Johnson was still using Ranger Pro. See P-Exh. 291.
Monsanto's reckless disregard for human safety is illustrated by the fact that Mr. Gould approved the
statement of a Roundup distributor regarding the impact of the IARC assessment on Roundup's
domestic PR and business: "We're being overrun by liberals, zombies, and morons," who need to be
"taken out one at a time, starting with the elections next year," at a time when Roundup had repeatedly
been associated with NHL, and following the IARC decision. P-Exh. 290 at 1.

Monsanto's efforts to curtail domestic regulatory reviews of Roundup could constitute malice, particularly in light of IARC's decision and Monsanto's reliance on the review of glyphosate by various regulatory agencies for the argument that Roundup is safe. P-Exh. 402 at 2 ("ATSDR Director and Branch Chief have promised Jack Housenger (Director of the US Office of Pesticide Programs) to put their report 'on hold' until after EPA releases its preliminary risk assessment (PRA) for glyphosate."). Indeed, Monsanto's contact within the EPA's Office of Pesticide Programs ("OPP") – Mr. Jess Rowland – bragged that he should be awarded a "medal" if able to "kill" the ATSDR review of glyphosate. P-Exh. 404 at 1-2. Mr. Rowland's role in aiding Monsanto to curtail regulatory safeguards was noted by Monsanto employees. P-Exh. 401 ("Jess doing a nice job at EPA...is there anyone we can get to inside EPA?").

The same evidence discussed in the previous sections in support of Plaintiff's strict liability and negligent failure to warn claims also supports the award for punitive damages, such as Monsanto's awareness of the presence of other carcinogens in the formulated product, and the dangers of POEA. P-Exh. 357 (William Heydens admitting internally in 2015 – only a couple of weeks after Monsanto was notified of Mr. Johnson's enquiries – that the Roundup formulation contains other known carcinogens

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such as NNG and formaldehyde); P-Exh. 383 at 3, 1 ("there are non-hazardous formulations, so why sel
a hazardous one?" Mr. Heydens explicitly states that the reason why it was important for Monsanto to
protect POEA despite its "demise" in Europe was to preserve Monsanto's world business); P-Exh. 282
(Monsanto's awareness of the epidemiology literature supporting causation in a memorandum by Donna
Farmer, John Acquavella and Daniel Goldstein stating that "[t]here are now six published studies that
arguably associate glyphosate and other pesticides with lymphopoietic cancers"). Monsanto's
nefarious intent is clearly illustrated by the fact that Donna Farmer and John Acquavella celebrated the
removal of glyphosate from the McDuffie (2001) paper's abstract, which made it more difficult for
researchers to pick up on the study's association between Roundup exposure and NHL. P-Exh. 313 at 1
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Despite Dr. Heydens admitting that the surfactant in the formulation may have been responsible for the tumors observed in the George (2010) study, Monsanto purposefully manipulated the scientific discourse to disguise Roundup's carcinogenic potential. P-Exh. 366 at 3; P-Exh 368 at 1 (Dr. Heydens drafting expert panel manuscript). *Roeken v. Philip Morris Inc.*, 127 Cal.App.4th 1640, 1690 (2005) (intentionally marketing a defective product knowing that it might cause injury and death is highly reprehensible). Dr. Sawyer testified that Monsanto has never submitted some of its POEA studies to the EPA, nor conducted carcinogenicity testing on them. July 26 Trns. at 3651:15-22. This is while Monsanto is aware that safer alternatives are available. *Id.* at 3627:7-17 and Monsanto's concerted efforts to "orchestrate outcry over IARC decision." P-Exh. 292 at 5.8 In response to IARC, as oppose to taking actions by warning about the dangers of its product, Monsanto ghost-wrote papers for an ostensibly independent expert panel. P-Exh. 336 at 1 ("Manuscript to be initiated by MON as ghostwriters.").9 Monsanto responded to IARC through scientific outreach strategies that can be used for "litigation support...Majority of writing can be done by Monsanto, keeping \$\$ down." P-Exh. 391 at

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<sup>&</sup>lt;sup>8</sup> Despite anticipating the IARC decision and knowing that Roundup had vulnerabilities in epidemiology, genotoxicity and mode of action, Monsanto purposefully attempted to rebut IARC and any suggestion that Roundup may be hazardous in reckless disregard for human health. P-Exh 294 at 1.

<sup>9</sup> Indeed, Ashley Roberts (Intertek) forwarded the expert panel manuscript to Dr. Heydens for review notwithstanding the acknowledgments section of the final published article explicitly stating that Monsanto personnel did not review the paper. P-Exh. 369 at 1.

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Importantly, the jury have been presented with evidence that Monsanto never responded to Mr. Johnson's two enquiries following Mr. Johnson's initial health complications from using Roundup, despite Monsanto being aware of an extensive data set associating Roundup with cancer at the time of Mr. Johnson's initial contact in 2014. P-Exh. 332 ("doctors are saying it is skin cancer...[Mr. Johnson] is just trying to find out if it could all be related to such a large exposure to Ranger Pro since he stated his skin was always perfect until this happened. He is looking for answers."); Goldstein Depo. at 37:13-17 ("As Mr. Johnson has stated under oath that you never called him, can you sit here and say, 'Oh, no, I did call him'? A. I do not recall."); P-Exh. 334 (Mr. Johnson's second enquiry to Monsanto); Goldstein Depo. at 49:20-50:6, 53:11-53:21 (Dr. Goldstein confirming that Mr. Johnson's second call was at the time of the IARC deliberation. "Q. But you don't remember ever getting back to Mr. Johnson and telling him, 'Hey, this group of scientists have just concluded there is a probable association between glyphosate and non-Hodgkin's lymphoma.' Nothing like that ever happened, right? A. I don't recall speaking with him."). Instead, Dr. Goldstein testified that he would tell Mr. Johnson to continue using Roundup notwithstanding Mr. Johnson's prognosis, exposure history, and the mounting evidence for Roundup carcinogenicity. Goldstein Depo. at 56:18-57:12. As a medical practitioner, Dr. Goldstein would be expected to adhere to the cautionary principle, as testified to by Dr. Ofodile, Mr. Johnson's treating physician. July 23 Trns. at 3155:13-3156:4.

Lastly, Dr. Goldstein testified that Monsanto has supported the American Council on Science and Health ("ASCH") over the years, a research organization promoting industry products. Goldstein Depo. at 120:9-16. In fact, Monsanto used the ACSH to respond to IARC. P-Exh. 321 at 1 (Monsanto discussing funding the ACSH). Despite this support for the ACSH, Dr. Goldstein testified that he has

<sup>&</sup>lt;sup>10</sup> Monsanto's initiatives to establish a "global network of third party experts" while Monsanto "works behind the scenes" to manipulate the scientific information on Roundup safety go straight to punitive intent. P-Exh. 378 at 1.

not always agreed with the ASCH's scientific positions on substances: "some of their positions on tobacco, some of their positions on lead, are not positions that I would agree with." Goldstein Depo. at 124:11-13. Although Dr. Goldstein disagrees with the ACSH's industry-friendly position on substances that have since been deemed known carcinogens, Dr. Goldstein has no reservations about his support for the ACSH in their defense of glyphosate. The credibility of Dr. Goldstein and evidence of Monsanto's conduct are ripe for consideration by the trier of fact in determining an appropriate punitive damages award.

#### **B. Punitive Damages Would Not Violate Due Process**

Monsanto argues that "evidence of conduct that occurred after the onset of Plaintiff's MF...cannot support an award of punitive damages without violating Monsanto's federal due process rights because there is no competent testimony that exposure to the formulation after onset of Plaintiff's MF contributed to the progression of the disease." Monsanto Brief at 21. This is false. Evidence was presented during Plaintiff's case-in-chief which supports the notion that Monsanto's failure in responding to Mr. Johnson's repeated enquiries or warning him exacerbated Mr. Johnson's cancer. As Dr. Nabhan testified, "[i]f [a patient] [is] being exposed to an agent that may be causing the cancer, you would tell them not to be exposed to this particular agent because it could make the cancer worse or it could cause another cancer... If anything, it just makes more sense not to spray, if you really have concerns that this is really causing the problem. July 20 Trns. at 2812:21-24, 2865:14-16. Indeed, Dr. Ofodile, Mr. Johnson's treating physician, reached the same conclusion, which is what prompted her to write a letter to the school-board, recommending that Mr. Johnson discontinue spraying:

I did write a letter to the School Board, and I requested that he not be exposed to any airborne environmental allergens as that could exacerbate his condition. Q. And specifically that was referring to the Ranger Pro and chemicals that he's talking about? A. Correct... A. My understanding is that, you know, it is okay and advisable to avoid something that could potentially be -- can exacerbate a condition, even if it's not been conclusive. So, you know, if I had a patient that was exposed to something and I thought may be causing it, my recommendation is to avoid it rather than waiting to see whether or not it truly does cause it later on.

July 23 Trns. at 3154:10-16, 3155:17-24.

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Additionally, Dr. Portier testified that glyphosate can be a tumor promotor, and Dr. Heydens conceded that Monsanto believed the "surfactant in the formulation will come up in the tumor promotion skin study [George et al 2010] because we think it played a role there." P-Exh. 366 at 2; July 12 Trns. at 1834:16-1837:14 ("A. glyphosate has the potential to be a promoter of carcinogenesis."). The Court must view such evidence in a light most favorable to Plaintiff and draw all reasonable inferences in Plaintiff's favor whilst disregarding Monsanto's reference to contrary evidence of the regulatory classification of IARC. Plaintiff's presentation of competent evidence from which a jury could conclude that repeated Roundup exposure exacerbated Mr. Johnson's cancer warrants evidence supporting punitive damages following Mr. Johnson's onset of cancer. Scientific Dispute

# C. Punitive Damages Would Not Violate the First Amendment or Inhibit Legitimate

The common law method of assessing punitive damages—i.e., an initial determination by a jury instructed to consider the gravity of the wrong and the need to deter similar wrongful conduct, followed by trial—is not "so inherently unfair as to deny due process and be per se unconstitutional." *Pacific* Mut. Life Ins. Co. v. Haslip (1991) 499 US 1, 17. Indeed, the Supreme Court has made it clear that the imposition of common law tort liability does not require a pesticide manufacturer to alter its label. Bates v. Dow Agrosciences LLC (2005) 544 U.S. 431, 432 (emphasis added) ("A requirement is a rule of law that must be obeyed; an event, such as a jury verdict, that merely motives an optional decision is not a requirement."") The same logic applies with equal force in nullifying Monsanto's first amendment claims. Finding punitive liability against Monsanto does not preclude Monsanto from continuing to assert that its product is not carcinogenic or continuing to not warn; this is because a jury verdict, as noted by the Supreme Court, is not a requirement, but an event that may incentivize an optional decision by a liable defendant. Monsanto can continue to participate in scientific debate and contend that its product is safe even after a punitive damages award. 11

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<sup>&</sup>lt;sup>11</sup> The two federal cases cited by Monsanto are completely inapposite. ONY, Inc. v. Cornerstone Therapeutics, Inc. (2d Cir. 2013) 720 F.3d 490 concerned claims under the Lanham Act which the court dismissed because "to the extent a speaker or author draws conclusions from non-fraudulent data, based on accurate descriptions of the data and methodology underlying those conclusions, on subjects about which there is legitimate ongoing scientific disagreement, those statements are not grounds for a claim

1	For the above reasons, and the fact that the decision is not binding on this Court, Monsanto als
2	cannot rely on the recent preliminary injunction issued by the Eastern District of California concerning
3	Monsanto's First Amendment challenges to Proposition 65 labeling. <i>National Association of Wheat</i>
4	Growers v. Zeise (E.D. Cal. 2018) 309 F.Supp.3d 842. Monsanto has repeatedly argued throughout
5	this trial that Prop 65 is irrelevant to the claims at bar as well as prejudicial, and this Court has agreed,
6	excluding any evidence of California's decision to list glyphosate as a substance known to the State to
7	cause cancer. Monsanto cannot now assert that Prop 65 is relevant for defending against the claim of
8	punitive damages if the Court has unequivocally held that the decision is not relevant to <i>any</i> issues at
9	trial: "The fact that glyphosate is listed under Proposition 65 is really not relevant in this case and is $n$
10	probative of any of the issues that the jury is going to have to decide." June 28 Trns. at 1187:19-22
11	(emphasis added). Monsanto cannot have its cake and eat it too.
12	Lastly, Monsanto contends that a punitive damages award cannot be supported in light of
13	dispute and debate in the scientific community regarding Roundup's carcinogenicity. The slew of nor
14	binding and unpersuasive federal cases cited by Monsanto have no purchase. For example, in Satcher
15	Honda Motor Co., 52 F.3d 1311, 1317 (5th Cir. 1995), the court noted that the punitive damages awar
16	did not withstand scrutiny because [t]he law of Mississippi allows punitive damages only in "extreme
17	cases;" they "are not favored in the law and are to be allowed only with caution and within narrow
18	limitsunder these standards, we conclude that [plaintiff] was not entitled to punitive damages" Id
19	1316. The record here, unlike <i>Satcher</i> (where only "certain studies" were relied upon by plaintiff), is
20	of false advertising under the Lanham Act." at 498. The case is not authority for the proposition that punitive damages in a toxic tort context violate the First Amendment merely because the liable

defendant disagrees with the scientific evidence which formed part of the verdict. The second

authority for a proposition not therein considered. Stand Up for California! v. State (2016) 6

distinguishable case dealt with defamation claims to which a defense under Wisconsin law was lack of actual malice. Underwager v. Salter (7th Cir. 1994) 22 F.3d 730, 736. Indeed, the 7th Circuit made it

clear "that we speak of Wisconsin law, not of the complex of rules under the first amendment...," much less the constitutionality of punitive liability under the principles of personal injury law. A case is not

<sup>12</sup> Judge Shubb did not have the record presented at this trial or discussed in the instant motion when

ruling on the partial preliminary injunction, did not consider competent causation evidence, and ruled based on otherwise "inadmissible evidence in considering preliminary relief." Nat'l Ass'n of Wheat

Cal.App.5th 686, 703.

Growers 309 F.Supp.3d at 842, fn 6.

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1	replete with studies over the course of decades ass	ociating Roundup with cancer, and troves of internal
2	Monsanto documents that, according to Judge Karnow, may rise to the less stringent California	
3	standards of malice, oppression or fraud.	
4	Similarly, Burke v. Deere & Co., 6 F.3d 49	7, 511 (8th Cir. 1993) was decided pursuant to
5	Iowa's restrictive punitive damages statute, where	, as Monsanto notes, "an award of punitive damages
6	is not appropriate when room exists for reasonable	disagreement over the relative risks and utilities of
7	the conduct at issue." Id. (emphasis added). Calif	fornia law does not require the trier of fact to consider
8	the risks and utilities of the conduct – in fact, this	case, as stated above, is proceeding under a consumer
9	expectation standard and not a risk/utility test. Additionally, the record in Burke, where "there was $no$	
10	evidence that [defendant] knew of any similar [faults with the product] when it sold the [product],"	
11	stands in stark contrast to the matter at bar, where, as discussed above (and affirmed by Judge Karnow),	
12	there is a plethora of evidence from which a jury could conclude that Monsanto was aware or should	
13	have been aware of Roundup's cancer risk and acted with reckless disregard in failing to warn. <i>Id.</i> at	
14	510 (emphasis added). Monsanto's reliance on inapposite, unpersuasive, non-binding case law provides	
15	no grounds for granting nonsuit on this extensive record.	
16	VII. CONCLUSION	
17	For the foregoing reasons, Monsanto's Mo	tion for Nonsuit should be denied and trial permitted
18	to continue.	
19	Resp	ectfully submitted,
20	July 30 2018	/s/ Pedram Esfandiary
21		im Esfandiary, Esq. (SBN: 312569)
22	Time	ent Wisner, Esq. (SBN: 276023) thy Litzenburg ( <i>pro hac vice</i> )
23	David	d Dickens (pro hac vice)
24		rneys for Plaintiff VAYNE LEE JOHNSON
25		WATNE LEE JOHNSON
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