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1	Curtis G. Hoke (SBN 282465)	ALAMEDA COUNTY August 30, 2018
2	THE MILLER FIRM, LLC 108 Railroad Ave.	CLERK OF
3	Orange, VA 22960 Phone: (540) 672-4224	THE SUPERIOR COURT By Dajuana Turner, Deputy
4	Fax: (540) 672-4224 Fax: (540) 672-3055 choke@millerfirmllc.com	CASE NUMBER: RG17862702
5	Attorneys for Plaintiffs	
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8	SUPERIOR COURT OF THE STATE OF CALIFORNIA	
9	FOR THE COUNTY OF ALAMEDA	
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11	COORDINATION PROCEEDING SPECIAL TITLE (RULE 3.550)	JCCP NO. 4953
12	ROUNDUP PRODUCTS CASES	Case No.: RG17862702
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14	THIS DOCUMENT RELATES TO:	DECLARATION OF CURTIS G. HOKE IN
15	Pilliod, et al. v. Monsanto Company, et al.	SUPPORT OF PLAINTIFFS' MOTION FOR LEAVE TO FILE A FIRST
16	Alameda Superior Court Case No.: RG17862702	AMENDED COMPLAINT
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20		Indeed Hon Johns Datroy
21		Judge: Hon. Ioana Petrou
22		Hearing Date: October 9, 2018 Time: 9:00 a.m.
23		Department: 17
24		Reservation No.: R-1993571
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28	DECLARATION OF CURTIS G. HOKE IN SUPPORT OF PLAINTIFFS' MOTION FOR LEAVE TO FILE AMENDED COMPLAINT	

DECLARATION OF CURTIS G. HOKE

I, Curtis Hoke, declare and state:

- 1. I am an attorney at law admitted to practice before all of the courts in the state of California. I am an attorney at The Miller Firm, LLC, attorneys of record for plaintiffs Alva Pilliod and Alberta Pilliod. I am over eighteen years of age and am fully competent to make this Declaration in support of Plaintiff's Motion for Leave to File a First Amended Complaint (the "Motion"). Except as otherwise expressly stated below, I have personal knowledge of the facts stated in this declaration, and if called to testify, I could and would competently testify to the matters stated herein.
- 2. On or about July 26, 2018, via a telephone conversation with Mrs. Alberta Pilliod, Plaintiffs' counsel discovered that Mrs. Pilliod in addition to her husband, Alva Pilliod was also diagnosed with non-Hodgkin's lymphoma. Plaintiff's counsel had previously believed that only Mr. Pilliod was diagnosed with non-Hodgkin's lymphoma, and believed that Mrs. Pilliod was diagnosed with brain cancer, which Plaintiffs' counsel did not know to be non-Hodgkin's lymphoma. In August, 2018, The Miller Firm received medical records showing that Mrs. Pilliod's brain cancer is actually a form of non-Hodgkin's lymphoma called diffuse large B-cell CNS lymphoma. Plaintiffs therefore seek leave to amend their Complaint to include these and related allegations.
- 3. On or about August 15, 2018, Plaintiffs' counsel met and conferred with Monsanto's counsel in order to determine whether Monsanto would stipulate to Plaintiffs' filing of their proposed First Amended Complaint. Monsanto objected to the proposed amendments, forcing Plaintiffs to file the accompanying Motion for Leave to File a First Amended Complaint.
 - 4. This Court has not yet set a trial date for this action.
- 5. Plaintiffs' Motion for Leave to File a First Amended Complaint seeks leave to amend Plaintiff's Complaint initially filed June 2, 2017. *See*, **Exhibit 1** appended hereto. Plaintiffs wish to add language to their Complaint in order to 1) add newly-discovered factual allegations that Mrs. Pilliod was also diagnosed with non-Hodgkin's lymphoma, allegedly due to her exposure to Roundup, 2) remove unnecessary allegations (Complaint ¶15-19) relating to co-plaintiffs Charles Baker, John Novak, Sharon Rowland, Sharon McClurg, and Marjorie Grubka, who have since filed their own separate Complaints, and 3) add Loss of Consortium claims held by each Plaintiff.

- 6. Attached hereto as **Exhibit 2** is a "strikethrough" draft of Plaintiffs' proposed First Amended Complaint. Additional allegations are highlighted; deleted paragraphs are stricken through.
- 7. Attached hereto as **Exhibit 3** is a "clean" version of Plaintiffs' proposed First Amended Complaint.
- 8. The effect of Plaintiffs' proposed First Amended Complaint is shown, in detail, on **Exhibit 3**. Plaintiffs' proposed First Amended Complaint would add the following additional allegations to Paragraph 14:

"Plaintiff ALBERTA PILLIOD is a competent individual over the age of 18, a resident and citizen of the United States, and hereby submits to the jurisdiction of the Court and alleges that jurisdiction and venue in this court is proper. Mrs. Pilliod currently resides in Livermore, County of Alameda, California. Mrs. Pilliod used the same Monsanto manufactured glyphosate-containing Roundup® products as Mr. Pilliod and suffered from severe physical, economic, and emotional injuries as a result of her use of Roundup® between 1975 and 2011, including but not limited to non-Hodgkin lymphoma, diagnosed in April, 2015 in California. Mr. and Mrs. Pilliod are lawfully married spouses and each seek damages for loss of consortium herein."

- 9. Plaintiffs' proposed First Amended Complaint also specifically makes a claim for Loss of Consortium held by both Mr. and Mrs. Pilliod. *See*, **Exhibit 3** at ¶¶ 163-166.
- 10. Finally, Plaintiffs' proposed First Amended Complaint would remove unnecessary allegations (Complaint ¶¶15-19) relating to co-plaintiffs 1) Charles Baker, 2) John Novak, 3) Sharon Rowland, 4) Sharon McClurg, and 5) Marjorie Grubka, who have since filed their own separate Complaints.
- 11. The above-described amendments are necessary and proper because they seek in good faith to add and amend language and causes of action that are consistent with the facts now known to Plaintiffs' counsel, but that were not originally pleaded, and failure to plead them now would deprive Plaintiffs of the right to compensation for meritorious claims and causes of action in this case.

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EXHIBIT 1

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FILED BY FAX Curtis G. Hoke (SBN 282465) ALAMEDA COUNTY THE MILLER FIRM, LLC June 02, 2017 108 Railroad Ave. CLERK OF THE SUPERIOR COURT By Burt Moskaira, Deputy Orange, VA 22960 Telephone: (540) 672-4224 CASE NUMBER: 4 Facsimile: (540) 672-3055 RG17862702 E-Mail: choke@millerfirmlic.com 5 Attorneys for Plaintiffs 6 7 8 SUPERIOR COURT OF THE STATE OF CALIFORNIA 9 IN AND FOR THE COUNTY OF ALAMEDA 10 (UNLIMITED JURISDICTION) 11 12 ALVA AND ALBERTA PILLIOD; CHARLES Case No.: 13 BAKER, JOHN NOVAK, SHARON 14 ROWLAND, SHARON MCCLURG, COMPLAINT FOR DAMAGES AND MARJORIE GRUBKA, 15 DEMAND FOR JURY TRIAL lб Plaintiffs, BY FAX 17 VS. 1. Strict Liability - Design Defect 18 2. Strict Liability - Failure to Warn 3. Negligence MONSANTO COMPANY; 19 4. Breach of Implied Warranty WILBUR-ELLIS COMPANY, LLC; and 20 5. Punitive Damage WILBUR-ELLIS FEED, LLC, 21 JURY TRIAL DEMANDED Defendants. 22 23 24 25 26 27 28 COMPLAINT FOR DAMAGES AND DEMAND FOR JURY TRIAL

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COMPLAINT AND DEMAND FOR JURY TRIAL

The above-captioned Plaintiffs (hereinafter, "Plaintiffs"), by attorneys, THE MILLER FIRM, LLC, as and for the Complaint herein allege upon information and belief the following:

SUMMARY OF THE CASE

- In 1970, defendant Monsanto Company, Inc. discovered the herbicidal properties of glyphosate and began marketing it in products in 1974 under the brand name Roundup®. Roundup® is a non-selective herbicide used to kill weeds that commonly compete with the growing of crops. By 2001, glyphosate had become the most-used active ingredient in American agriculture with 85-90 millions of pounds used annually. That number grew to 185 million pounds by 2007. As of 2013, glyphosate was the world's most widely used herbicide.
- Monsanto is a multinational agricultural biotechnology corporation based in St. Louis, Missouri. It is the world's leading producer of glyphosate. As of 2009, Monsanto was the world's leading producer of seeds, accounting for 27% of the world seed market. The majority of these seeds are of the Roundup Ready® brand. The stated advantage of Roundup Ready® crops is that they substantially improve a farmer's ability to control weeds, since glyphosate can be sprayed in the fields during the growing season without harming their crops. In 2010, an estimated 70% of corn and cotton, and 90% of soybean fields in the United States were Roundup Ready®.
- Monsanto's glyphosate products are registered in 130 countries and approved for use on 3. over 100 different crops. They are ubiquitous in the environment. Numerous studies confirm that glyphosate is found in rivers, streams, and groundwater in agricultural areas where Roundup® is COMPLAINT FOR DAMAGES AND DEMAND FOR JURY TRIAL

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used. It has been found in food, in the urine of agricultural workers, and even in the urine of urban dwellers who are not in direct contact with glyphosate.

- 4. On March 20, 2015, the International Agency for Research on Cancer ("IARC"), an agency of the World Health Organization ("WHO"), issued an evaluation of several herbicides, including glyphosate. That evaluation was based, in part, on studies of exposures to glyphosate in several countries around the world, and it traces the health implications from exposure to glyphosate since 2001.
- 5. On July 29, 2015, IARC issued the formal monograph relating to glyphosate. In that monograph, the IARC Working Group provides a thorough review of the numerous studies and data relating to glyphosate exposure in humans.
- 6. The IARC Working Group classified glyphosate as a Group 2A herbicide, which means that it is probably carcinogenic to humans. The IARC Working Group concluded that the cancers most associated with glyphosate exposure are non-Hodgkin lymphoma and other haematopoietic cancers, including lymphocytic lymphoma/chronic lymphocytic leukemia, B-cell lymphoma, and multiple myeloma.
- 7. The IARC evaluation is significant. It confirms what has been believed for years: that glyphosate is toxic to humans.
- 8. Nevertheless, Monsanto, since it began selling Roundup®, has represented it as safe to COMPLAINT FOR DAMAGES AND DEMAND FOR JURY TRIAL

humans and the environment. Indeed, Monsanto has repeatedly proclaimed and continues to proclaim to the world, and particularly to United States consumers, that glyphosate-based herbicides, including Roundup®, create no unreasonable risks to human health or to the environment.

JURISDICTION AND VENUE

- 9. The California Superior Court has jurisdiction over this action pursuant to California Constitution Article VI, Section 10, which grants the Superior Court "original jurisdiction in all causes except those given by statute to other trial courts." The Statutes under which this action is brought do not specify any other basis for jurisdiction.
- 10. The California Superior Court has jurisdiction over the Defendants because, based on information and belief, each is a California resident, a corporation and/or entity organized under the laws of the State of California, a foreign corporation or association authorized to do business in California and registered with the California Secretary of State or has sufficient minimum contacts in California, or otherwise intentionally avails itself of the California market so as to render the exercise of jurisdiction over it by the California courts consistent with traditional notions of fair play and substantial justice.
- 11. Venue is proper in this Court pursuant to California Code of Civil Procedure Section 395 in that the subject injury occurred in Alameda County.

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12. Furthermore the Defendants have purposefully availed themselves of the benefits and the protections of the laws within the State of California. Monsanto has had sufficient contact such that the exercise of jurisdiction would be consistent with the traditional notions of fair play and substantial justice.

13. Plaintiffs seek relief that is within the jurisdictional limits of this Court.

THE PARTIES

Plaintiffs

- 14. Plaintiff ALVA PILLIOD is a competent individual over the age of 18, a resident and citizen of the United States, and hereby submits to the jurisdiction of the Court and alleges that jurisdiction and venue in this court is proper. Mr. Pilliod currently resides in Livermore, County of Alameda, California. Mr. Pilliod used Monsanto's glyphosate-containing Roundup® product and suffered from severe physical, economy, and emotional injuries as a result of his use of Roundup® between 1975 and 2011, including but not limited to non-Hodgkin lymphoma, diagnosed in 2011 in California.
- 15. Plaintiff CHARLES BAKER is a competent individual over the age of 18, a resident and citizen of the United States, and hereby submits to the jurisdiction of the Court and alleges that jurisdiction and venue in this court is proper. Mr. Baker currently resides in Raymond, California. Mr. Baker used Monsanto's glyphosate-containing Roundup® product at work and at home from 1995 to 2016 and suffered from severe physical, economy, and emotional injuries as a result of his use of Roundup®, including but not limited to non-Hodgkin lymphoma, diagnosed in June, 2015 in California.

- 16. Plaintiff JOHN NOVAK is a competent individual over the age of 18, a resident and citizen of the United States, and hereby submits to the jurisdiction of the Court and alleges that jurisdiction and venue in this court is proper. Mr. Novak currently resides in Vista, California. Mr. Novak used Monsanto's glyphosate-containing Roundup® product at work and at home from 2002 to 2010 and suffered from severe physical, economy, and emotional injuries as a result of his use of Roundup®, including but not limited to non-Hodgkin lymphoma, diagnosed in 2011 in California.
- 17. Plaintiff SHARON ROWLAND is a competent individual over the age of 18, a resident and citizen of the United States, and hereby submits to the jurisdiction of the Court and alleges that jurisdiction and venue in this court is proper. Ms. Rowland currently resides in Fontana, California. Ms. Rowland used Monsanto's glyphosate-containing Roundup® product from 1985 to 2015 on her 10 acres and suffered from severe physical, economy, and emotional injuries as a result of his use of Roundup®, including but not limited to non-Hodgkin lymphoma, diagnosed in June, 2015 in California.
- 18. Plaintiff SHARON MCCLURG is a competent individual over the age of 18, a resident and citizen of the United States, and hereby submits to the jurisdiction of the Court and alleges that jurisdiction and venue in this court is proper. Ms. McClurg currently resides in Roseville, California. Ms. McClurg used Monsanto's glyphosate-containing Roundup® product at home from 1974 to 2016 and suffered from severe physical, economy, and emotional injuries as a result of his use of Roundup®, including but not limited to non-Hodgkin lymphoma, diagnosed in 2003 in California.
- 19. Plaintiff MARJORIE GRUBKA is a competent individual over the age of 18, a resident and citizen of the United States, and hereby submits to the jurisdiction of the Court and alleges COMPLAINT FOR DAMAGES AND DEMAND FOR JURY TRIAL

that jurisdiction and venue in this court is proper. Ms. Grubka currently resides in Martinez, California. Mr. Cole used Monsanto's glyphosate-containing Roundup® product at home from 1986 to 2016 and suffered from severe physical, economy, and emotional injuries as a result of his use of Roundup®, including but not limited to non-Hodgkin lymphoma, diagnosed in December 2016 in California.

Defendants

- 20. Defendant Monsanto Company ("Monsanto") is a Delaware corporation with its headquarters and principal place of business in St. Louis, Missouri. At all times relevant to this complaint, Monsanto was the entity that discovered the herbicidal properties of glyphosate and the manufacturer of Roundup®. Monsanto has regularly transacted and conducted business within the state of California, and has derived substantial revenue from goods and products, including Roundup, used in the State of California. Monsanto expected or should have expected their acts to have consequences within the State of California, and derived substantial revenue from interstate commerce.
- 21. Defendants Wilbur-Ellis Company LLC is a California limited liability corporation with its headquarters and principal place of business in San Francisco, California. At all times relevant to this complaint, Wilbur-Ellis Company, LLC sold and distributed Monsanto products including Roundup, within the State of California.
- 22. Defendants Wilbur-Ellis Feed LLC (with Wilbur-Ellis Company LLC, hereinafter "Wilbur-Ellis") is a California limited liability corporation with its headquarters and principal place of business in San Francisco, California. At all times relevant to this complaint, Wilbur-COMPLAINT FOR DAMAGES AND DEMAND FOR JURY TRIAL

Ellis Feed, LLC sold and distributed Monsanto products including Roundup, within the State of California.

- 23. Defendants Monsanto Company, Wilbur-Ellis Company, LLC, and Wilbur-Ellis Feed, LLC are collectively referred to throughout this Complaint as "Defendants."
- 24. Plaintiffs are informed and believe, and based thereon allege, that in committing the acts alleged herein, each and every managing agent, agent, representative and/or employee of the Defendants was working within the course and scope of said agency, representation and/or employment with the knowledge, consent, ratification, and authorization of the Defendants and their directors, officers and/or managing agents.

CASE FACTS

- 25. Glyphosate is a broad-spectrum, non-selective herbicide used in a wide variety of herbicidal products around the world.
- 26. Plants treated with glyphosate translocate the systemic herbicide to their roots, shoot regions and fruit, where it interferes with the plant's ability to form aromatic amino acids necessary for protein synthesis.
- 27. Treated plants generally die within two to three days. Because plants absorb glyphosate, it cannot be completely removed by washing or peeling produce or by milling, baking, or brewing grains.

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28. For nearly 40 years, farms across the world have used Roundup® without knowing of the dangers its use poses. That is because when Monsanto first introduced Roundup®, it touted glyphosate as a technological breakthrough: it could kill almost every weed without causing harm either to people or to the environment. Of course, history has shown that not to be true. According to the WHO, the main chemical ingredient of Roundup®—glyphosate—is a probable cause of cancer. Those most at risk are farm workers and other individuals with workplace exposure to Roundup®, such as workers in garden centers, nurseries, and landscapers. Agricultural workers are, once again, victims of corporate greed.

29. Monsanto assured the public that Roundup® was harmless. In order to prove this, Monsanto championed falsified data and attacked legitimate studies that revealed its dangers. Monsanto led a prolonged campaign of misinformation to convince government agencies, farmers and the general population that Roundup® was safe.

The Discovery of Glyphosate and Development of Roundup®

30. The herbicidal properties of glyphosate were discovered in 1970 by Monsanto chemist John Franz. The first glyphosate-based herbicide was introduced to the market in the mid-1970s under the brand name Roundup®. From the outset, Monsanto marketed Roundup® as a "safe" general-purpose herbicide for widespread commercial and consumer use. It still markets Roundup® as safe today.

Registration of Herbicides under Federal Law

31. The manufacture, formulation and distribution of herbicides, such as Roundup®, are regulated under the Federal Insecticide, Fungicide, and Rodenticide Act ("FIFRA" or "Act"), 7

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U.S.C. § 136 et seq. FIFRA requires that all pesticides be registered with the Environmental Protection Agency ("EPA" or "Agency") prior to their distribution, sale, or use, except as described by the Act. 7 U.S.C. § 136a(a)

- 32. Because pesticides are toxic to plants, animals, and humans, at least to some degree, the EPA requires as part of the registration process, among other things, a variety of tests to evaluate the potential for exposure to pesticides, toxicity to people and other potential non-target organisms, and other adverse effects on the environment. Registration by the EPA, however, is not an assurance or finding of safety. The determination the Agency must make in registering or re-registering a product is not that the product is "safe," but rather that use of the product in accordance with its label directions "will not generally cause unreasonable adverse effects on the environment." 7 U.S.C. § 136a(c)(5)(D).
- 33. FIFRA defines "unreasonable adverse effects on the environment" to mean "any unreasonable risk to man or the environment, taking into account the economic, social, and environmental costs and benefits of the use of any pesticide." 7 U.S.C. § 136(bb). FIFRA thus requires EPA to make a risk/benefit analysis in determining whether a registration should be granted or allowed to continue to be sold in commerce.
- 34. The EPA and the State of California registered Roundup® for distribution, sale, and manufacture in the United States and the State of California.

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the conduct of tests required for registration and the laboratory practices that must be followed in conducting these tests. The data produced by the registrant must be submitted to the EPA for review and evaluation. The government is not required, nor is it able, however, to perform the product tests that are required of the manufacturer.

36. The evaluation of each pesticide product distributed, sold, or manufactured is completed at the time the product is initially registered. The data necessary for registration of a pesticide has changed over time. The EPA is now in the process of re-evaluating all pesticide products through a Congressionally-mandated process called "re-registration." 7 U.S.C. § 136a-1. In order to reevaluate these pesticides, the EPA is demanding the completion of additional tests and the submission of data for the EPA's review and evaluation.

FIFRA generally requires that the registrant, Monsanto in the case of Roundup®,

conducts the health and safety testing of pesticide products. The EPA has protocols governing

37. In the case of glyphosate, and therefore Roundup®, the EPA had planned on releasing its preliminary risk assessment —in relation to the reregistration process—no later than July 2015. The EPA completed its review of glyphosate in early 2015, but it delayed releasing the risk assessment pending further review in light of the WHO's health-related findings.

Scientific Fraud Underlying the Marketing and Sale of Glyphosate/Roundup

38. Based on early studies that glyphosate could cause cancer in laboratory animals, the EPA originally classified glyphosate as *possibly carcinogenic to humans* (Group C) in 1985. After

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pressure from Monsanto, including contrary studies it provided to the EPA, the EPA changed its classification to evidence of non-carcinogenicity in humans (Group E) in 1991. In so classifying glyphosate, however, the EPA made clear that the designation did not mean the chemical does not cause cancer: "It should be emphasized, however, that designation of an agent in Group E is based on the available evidence at the time of evaluation and should not be interpreted as a definitive conclusion that the agent will not be a carcinogen under any circumstances."

- 39. On two occasions, the EPA found that the laboratories hired by Monsanto to test the toxicity of its Roundup® products for registration purposes committed fraud.
- 40. In the first instance, Monsanto, in seeking initial registration of Roundup® by EPA, hired Industrial Bio-Test Laboratories ("IBT") to perform and evaluate pesticide toxicology studies relating to Roundup®. IBT performed about 30 tests on glyphosate and glyphosate-containing products, including nine of the 15 residue studies needed to register Roundup®.
- In 1976, the United States Food and Drug Administration ("FDA") performed an inspection of Industrial Bio-Test Industries ("IBT") that revealed discrepancies between the raw data and the final report relating to the toxicological impacts of glyphosate. The EPA subsequently audited IBT; it too found the toxicology studies conducted for the Roundup® herbicide to be invalid. An EPA reviewer stated, after finding "routine falsification of data" at IBT, that it was "hard to believe the scientific integrity of the studies when they said they took specimens of the uterus from male rabbits."

42. Three top executives of IBT were convicted of fraud in 1983.

43. In the second incident of data falsification, Monsanto hired Craven Laboratories in 1991 to perform pesticide and herbicide studies, including for Roundup®. In that same year, the owner of Craven Laboratories and three of its employees were indicted, and later convicted, of fraudulent laboratory practices in the testing of pesticides and herbicides.

44. Despite the falsity of the tests that underlie its registration, within a few years of its launch, Monsanto was marketing Roundup® in 115 countries.

The Importance of Roundup® to Monsanto's Market Dominance Profits

- 45. The success of Roundup® was key to Monsanto's continued reputation and dominance in the marketplace. Largely due to the success of Roundup® sales, Monsanto's agriculture division was out-performing its chemicals division's operating income, and that gap increased yearly. But with its patent for glyphosate expiring in the United States in the year 2000, Monsanto needed a strategy to maintain its Roundup® market dominance and to ward off impending competition.
- A6. In response, Monsanto began the development and sale of genetically engineered Roundup Ready® seeds in 1996. Since Roundup Ready® crops are resistant to glyphosate; farmers can spray Roundup® onto their fields during the growing season without harming the crop. This allowed Monsanto to expand its market for Roundup® even further; by 2000, Monsanto's biotechnology seeds were planted on more than 80 million acres worldwide and

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27 28 nearly 70% of American soybeans were planted from Roundup Ready® seeds. It also secured Monsanto's dominant share of the glyphosate/Roundup® market through a marketing strategy that coupled proprietary Roundup Ready® seeds with continued sales of its Roundup® herbicide.

47. Through a three-pronged strategy of increased production, decreased prices and by coupling with Roundup Ready® seeds, Roundup® became Monsanto's most profitable product. In 2000, Roundup® accounted for almost \$2.8 billion in sales, outselling other herbicides by a margin of five to one, and accounting for close to half of Monsanto's revenue. Today, glyphosate remains one of the world's largest herbicides by sales volume.

Monsanto has known for decades that it falsely advertises the safety of Roundup®

- 48. In 1996, the New York Attorney General ("NYAG") filed a lawsuit against Monsanto based on its false and misleading advertising of Roundup ® products. Specifically, the lawsuit challenged Monsanto's general representations that its spray-on glyphosate-based herbicides, including Roundup®, were "safer than table salt" and "practically non-toxic" to mammals, birds, and fish. Among the representations the NYAG found deceptive and misleading about the human and environmental safety of Roundup® are the following:
- a) Remember that environmentally friendly Roundup herbicide is biodegradable. It won't build up in the soil so you can use Roundup with confidence along customers' driveways, sidewalks and fences ...
- b) And remember that Roundup is biodegradable and won't build up in the soil. That will give you the environmental confidence you need to use Roundup everywhere you've got a weed, brush, edging or trimming problem. COMPLAINT FOR DAMAGES AND DEMAND FOR JURY TRIAL

c) Roundup biodegrades into naturally occurring elements.

- d) Remember that versatile Roundup herbicide stays where you put it. That means there's no washing or leaching to harm customers' shrubs or other desirable vegetation.
- e) This non-residual herbicide will not wash or leach in the soil. It ... stays where you apply it.
- f) You can apply Accord with "confidence because it will stay where you put it" it bonds tightly to soil particles, preventing leaching. Then, soon after application, soil microorganisms biodegrade Accord into natural products.
 - g) Glyphosate is less toxic to rats than table salt following acute oral ingestion.
- h) Glyphosate's safety margin is much greater than required. It has over a 1,000-fold safety margin in food and over a 700-fold safety margin for workers who manufacture it or use it.
- i) You can feel good about using herbicides by Monsanto. They carry a toxicity category rating of 'practically non-toxic' as it pertains to mammals, birds and fish.
- j) "Roundup can be used where kids and pets will play and breaks down into natural material." This ad depicts a person with his head in the ground and a pet dog standing in an area which has been treated with Roundup.
- 49. On November 19, 1996, Monsanto entered into an Assurance of Discontinuance with NYAG, in which Monsanto agreed, among other things, "to cease and desist from publishing or broadcasting any advertisements [in New York] that represent, directly or by implication" that:
- a) its glyphosate-containing pesticide products or any component thereof are safe, non-toxic, harmless or free from risk.

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b) its glyphosate-containing pesticide products or any component thereof

manufactured, formulated, distributed or sold by Monsanto are biodegradable

* * *

c) its glyphosate-containing pesticide products or any component thereof stay where they are applied under all circumstances and will not move through the environment by any means.

* * *

d) its glyphosate-containing pesticide products or any component thereof are "good" for the environment or are "known for their environmental characteristics."

* * *

- e) glyphosate-containing pesticide products or any component thereof are safer or less toxic than common consumer products other than herbicides;
- f) its glyphosate-containing products or any component thereof might be classified as "practically non-toxic.
- 50. Monsanto did not alter its advertising in the same manner in any state other than New York, and on information and belief still has not done so today.
- 51. In 2009, France's highest court ruled that Monsanto had not told the truth about the safety of Roundup®. The French court affirmed an earlier judgement that Monsanto had falsely advertised its herbicide Roundup® as "biodegradable" and that it "left the soil clean."

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Classifications and Assessments of Glyphosate

- The IARC process for the classification of glyphosate followed the stringent procedures for the evaluation of a chemical agent. Over time, the IARC Monograph program has reviewed 980 agents. Of those reviewed, it has determined 116 agents to be Group 1 (Known Human Carcinogens); 73 agents to be Group 2A (Probable Human Carcinogens); 287 agents to be Group 2B (Possible Human Carcinogens); 503 agents to be Group 3 (Not Classified); and one agent to be Probably Not Carcinogenic.
- The established procedure for IARC Monograph evaluations is described in the IARC Programme's Preamble. Evaluations are performed by panels of international experts, selected on the basis of their expertise and the absence of actual or apparent conflicts of interest.
- 54. One year before the Monograph meeting, the meeting is announced and there is a call both for data and for experts. Eight months before the Monograph meeting, the Working Group membership is selected and the sections of the Monograph are developed by the Working Group members. One month prior to the Monograph meeting, the call for data is closed and the various draft sections are distributed among Working Group members for review and comment. Finally, at the Monograph meeting, the Working Group finalizes review of all literature, evaluates the evidence in each category, and completes the overall evaluation. Within two weeks after the Monograph meeting, the summary of the Working Group findings are published in Lancet Oncology, and within a year after the meeting, the final Monograph is finalized and published.

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55. In assessing an agent, the IARC Working Group reviews the following information: (a) human, experimental, and mechanistic data; (b) all pertinent epidemiological studies and cancer bioassays; and (c) representative mechanistic data. The studies must be publicly available and have sufficient detail for meaningful review, and reviewers cannot be associated with the underlying study.

- 56. In March 2015, IARC reassessed glyphosate. The summary published in *The Lancet Oncology* reported that glyphosate is a Group 2A agent and probably carcinogenic in humans.
- 57. On July 29, 2015, IARC issued its Monograph for glyphosate, Monograph 112. For Volume 112, the volume that assessed glyphosate, a Working Group of 17 experts from 11 countries met at IARC from March 3–10, 2015, to assess the carcinogenicity of certain herbicides, including glyphosate. The March meeting culminated nearly a one-year review and preparation by the IARC Secretariat and the Working Group, including a comprehensive review of the latest available scientific evidence. According to published procedures, the Working Group considered "reports that have been published or accepted for publication in the openly available scientific literature" as well as "data from governmental reports that are publicly available."
- 58. The studies considered the following exposure groups: occupational exposure of farmers and tree nursery workers in the United States, forestry workers in Canada and Finland and COMPLAINT FOR DAMAGES AND DEMAND FOR JURY TRIAL

municipal weed-control workers in the United Kingdom; and para-occupational exposure in farming families.

59. Glyphosate was identified as the second-most used household herbicide in the United States for weed control between 2001 and 2007 and the most heavily used herbicide in the world in 2012.

60. Exposure pathways are identified as air (especially during spraying), water, and food.

Community exposure to glyphosate is widespread and found in soil, air, surface water, and groundwater, as well as in food.

- 61. The assessment of the IARC Working Group identified several case control studies of occupational exposure in the United States, Canada, and Sweden. These studies show a human health concern from agricultural and other work-related exposure to glyphosate.
- 62. The IARC Working Group found an increased risk between exposure to glyphosate and non-Hodgkin lymphoma ("NHL") and several subtypes of NHL, and the increased risk persisted after adjustment for other pesticides.

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- 63. The IARC Working Group also found that glyphosate caused DNA and chromosomal damage in human cells. One study in community residents reported increases in blood markers of chromosomal damage (micronuclei) after glyphosate formulations were sprayed.
- 64. In male CD-1 mice, glyphosate induced a positive trend in the incidence of a rare tumor, renal tubule carcinoma. A second study reported a positive trend for haemangiosarcoma in male mice. Glyphosate increased pancreatic islet-cell adenoma in male rats in two studies. A glyphosate formulation promoted skin tumors in an initiation-promotion study in mice.
- 65. The IARC Working Group also noted that glyphosate has been detected in the urine of agricultural workers, indicating absorption. Soil microbes degrade glyphosate to aminomethylphosphoric acid (AMPA). Blood AMPA detection after exposure suggests intestinal microbial metabolism in humans.
- 66. The IARC Working Group further found that glyphosate and glyphosate formulations induced DNA and chromosomal damage in mammals, and in human and animal cells in utero.
- 67. The IARC Working Group also noted genotoxic, hormonal, and enzymatic effects in mammals exposed to glyphosate. Essentially, glyphosate inhibits the biosynthesis of aromatic amino acids, which leads to several metabolic disturbances, including the inhibition of protein and secondary product biosynthesis and general metabolic disruption.

68. The IARC Working Group also reviewed an Agricultural Health Study, consisting of a prospective cohort of 57,311 licensed pesticide applicators in Iowa and North Carolina. While this study differed from others in that it was based on a self-administered questionnaire, the results support an association between glyphosate exposure and Multiple Myeloma, Hairy Cell Leukemia (HCL), and Chronic Lymphocytic Leukemia (CLL), in addition to several other cancers.

Other Earlier Findings About Glyphosate's Dangers to Human Health

69. The EPA has a technical fact sheet, as part of its Drinking Water and Health, National Primary Drinking Water Regulations publication, relating to glyphosate. This technical fact sheet predates the IARC March 20, 2015, evaluation. The fact sheet describes the release patterns for glyphosate as follows:

Release Patterns

- 70. Glyphosate is released to the environment in its use as a herbicide for controlling woody and herbaceous weeds on forestry, right-of-way, cropped and non-cropped sites. These sites may be around water and in wetlands.
- 71. It may also be released to the environment during its manufacture, formulation, transport, storage, disposal and cleanup, and from spills. Since glyphosate is not a listed chemical in the Toxics Release Inventory, data on releases during its manufacture and handling are not available.
- 72. Occupational workers and home gardeners may be exposed to glyphosate by inhalation and dermal contact during spraying, mixing, and cleanup. They may also be exposed by touching COMPLAINT FOR DAMAGES AND DEMAND FOR JURY TRIAL

soil and plants to which glyphosate was applied. Occupational exposure may also occur during glyphosate's manufacture, transport storage, and disposal.

73. In 1995, the Northwest Coalition for Alternatives to Pesticides reported that in California, the state with the most comprehensive program for reporting of pesticide-caused illness, glyphosate was the third most commonly-reported cause of pesticide illness among agricultural workers.

Recent Worldwide Bans on Roundup®/Glyphosate

- 74. Several countries around the world have instituted bans on the sale of Roundup® and other glyphosate-containing herbicides, both before and since IARC first announced its assessment for glyphosate in March 2015, and more countries undoubtedly will follow suit in light of the as the dangers of the use of Roundup® are more widely known. The Netherlands issued a ban on all glyphosate-based herbicides in April 2014, including Roundup®, which takes effect by the end of 2015. In issuing the ban, the Dutch Parliament member who introduced the successful legislation stated: "Agricultural pesticides in user-friendly packaging are sold in abundance to private persons. In garden centers, Roundup® is promoted as harmless, but unsuspecting customers have no idea what the risks of this product are. Especially children are sensitive to toxic substances and should therefore not be exposed to it."
- 75. The Brazilian Public Prosecutor in the Federal District requested that the Brazilian Justice Department suspend the use of glyphosate.

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81. Wilbur-Ellis had superior knowledge compared to Roundup users and consumers, including regarding the carcinogenic properties of the product, yet failed to accompany its sales and/or marketing of Roundup with any warnings or precautions for that grave danger. On information and belief, Wilbur-Ellis was one of the distributors providing Roundup and other glyphosate-containing products actually used by the Plaintiffs.

CLAIM ONE

STRICT LIABILITY - DESIGN DEFECT

(Against All Defendants)

- 82. Plaintiffs incorporate by reference each and every allegation set forth in the preceding paragraphs as if fully stated herein.
- 83. Plaintiffs brings this strict liability claim against Defendants for defective design.
- At all times relevant to this litigation, Defendants engaged in the business of testing, developing, designing, manufacturing, marketing, selling, distributing, and promoting Roundup® products, which are defective and unreasonably dangerous to consumers, including Plaintiffs, thereby placing Roundup® products into the stream of commerce. These actions were under the ultimate control and supervision of Defendants. At all times relevant to this litigation, Defendants designed, researched, developed, manufactured, produced, tested, assembled, labeled, advertised, promoted, marketed, sold, and distributed the Roundup® products that Plaintiffs were exposed to, as described above.

85. At all times relevant to this litigation, Defendants' Roundup® products were manufactured, designed, and labeled in an unsafe, defective, and inherently dangerous manner that was dangerous for use by or exposure to the public, and, in particular, the Plaintiffs.

86. At all times relevant to this litigation, Defendants' Roundup® products reached the intended consumers, handlers, and users or other persons coming into contact with these products in California and throughout the United States, including Plaintiffs, without substantial change in their condition as designed, manufactured, sold, distributed, labeled, and marketed by Defendants.

87. Defendants' Roundup® products, as researched, tested, developed, designed, licensed, manufactured, packaged, labeled, distributed, sold, and marketed by Defendants were defective in design and formulation in that when they left the hands of the Defendants' manufacturers and/or suppliers, they were unreasonably dangerous and dangerous to an extent beyond that which an ordinary consumer would contemplate.

88. Defendants' Roundup® products, as researched, tested, developed, designed, licensed, manufactured, packaged, labeled, distributed, sold, and marketed by Defendants were defective in design and formulation in that when they left the hands of Defendants' manufacturers and/or suppliers, the foreseeable risks exceeded the alleged benefits associated with their design and formulation.

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- 89. At all times relevant to this action, Defendants knew or had reason to know that i Roundup® products were defective and were inherently dangerous and unsafe when used in the manner instructed and provided by Defendants.
- 90. Therefore, at all times relevant to this litigation, Defendants' Roundup® products, as researched, tested, developed, designed, licensed, manufactured, packaged, labeled, distributed, sold and marketed by Defendants were defective in design and formulation, in one or more of the following ways:
- a. When placed in the stream of commerce, Defendants' Roundup® products were defective in design and formulation, and, consequently, dangerous to an extent beyond that which an ordinary consumer would contemplate.
- b. When placed in the stream of commerce, Defendants' Roundup® products were unreasonably dangerous in that they were hazardous and posed a grave risk of cancer and other serious illnesses when used in a reasonably anticipated manner.
- c. When placed in the stream of commerce, Defendants' Roundup® products contained unreasonably dangerous design defects and were not reasonably safe when used in a reasonably anticipated or intended manner.
- d. Defendants did not sufficiently test, investigate, or study its Roundup® products and, specifically, the active ingredient glyphosate.
- e. Exposure to Roundup® and glyphosate-containing products presents a risk of harmful side effects that outweigh any potential utility stemming from the use of the herbicide.

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- f. Defendants knew or should have known at the time of marketing its Roundup® products that exposure to Roundup® and specifically, its active ingredient glyphosate, could result in cancer and other severe illnesses and injuries.
- g. Defendants did not conduct adequate post-marketing surveillance of its Roundup® products.

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- h. Defendants could have employed safer alternative designs and formulations.
- 91. Plaintiffs were exposed to Defendants' Roundup® products without knowledge of its dangerous characteristics.
- 92. At all times relevant to this litigation, Plaintiffs were exposed to the use of Defendants' Roundup® products in an intended or reasonably foreseeable manner without knowledge of their dangerous characteristics.
- 93. Plaintiffs could not have reasonably discovered the defects and risks associated with Roundup® or glyphosate-containing products before or at the time of exposure.
- 94. The harm caused by Defendants' Roundup® products far outweighed their benefit, rendering Defendants' products dangerous to an extent beyond that which an ordinary consumer would contemplate. Defendants' Roundup® products were and are more dangerous than alternative products and Defendants could have designed its Roundup® products to make them less dangerous. Indeed, at the time that Defendants designed its Roundup® products, the state of

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the industry's scientific knowledge was such that a less risky design or formulation was attainable.

- 95. At the time Roundup® products left Defendants' control, there was a practical, technically feasible and safer alternative design that would have prevented the harm without substantially impairing the reasonably anticipated or intended function of Defendants' herbicides.
- 96. Defendants' defective design of its Roundup® products was willful, wanton, fraudulent, malicious, and conducted with reckless disregard for the health and safety of users of the Roundup® products, including the Plaintiffs herein.
- 97. Therefore, as a result of the unreasonably dangerous condition of its Roundup® products, Defendants are strictly liable to Plaintiffs.
- 98. The defects in Defendants' Roundup® products were substantial and contributing factors in causing Plaintiffs' grave injuries, and, but for Defendants' misconduct and omissions, Plaintiffs would not have sustained their injuries.
- 99. Defendants' conduct, as described above, was reckless. Defendants risked the lives of consumers and users of its products, including Plaintiffs, with knowledge of the safety problems associated with Roundup® and glyphosate-containing products, and suppressed this knowledge

from the general public. Defendants made conscious decisions not to redesign, warn or inform the unsuspecting public. Defendants' reckless conduct warrants an award of punitive damages.

100. As a direct and proximate result of Defendants placing defective Roundup® products into the stream of commerce, Plaintiffs have suffered and continue to suffer grave injuries, and has endured physical pain and discomfort, as well as economic hardship, including considerable financial expenses for medical care and treatment. Plaintiffs will continue to incur these expenses in the future.

101. WHEREFORE, Plaintiffs respectfully request that this Court enter judgment in Plaintiffs' favor for compensatory and punitive damages, together with interest, costs herein incurred, attorneys' fees and all such other and further relief as this Court deems just and proper. Plaintiffs also demands a jury trial on the issues contained herein.

CLAIM TWO

STRICT LIABILITY - FAILURE TO WARN

(Against All Defendants)

- 102. Plaintiffs incorporate by reference each and every allegation set forth in the preceding paragraphs as if fully stated herein.
- 103. Plaintiffs bring this strict liability claim against Defendants for failure to warn.

104. At all times relevant to this litigation, Defendants engaged in the business of testing, developing, designing, manufacturing, marketing, selling, distributing, promoting and applying Roundup® products, which are defective and unreasonably dangerous to consumers, including Plaintiffs, because they do not contain adequate warnings or instructions concerning the dangerous characteristics of Roundup® and specifically, the active ingredient glyphosate. These actions were under the ultimate control and supervision of Defendants.

105. Monsanto researched, developed, designed, tested, manufactured, inspected, labeled, distributed, marketed, promoted, sold, and otherwise released into the stream of commerce its Roundup® products, and in the course of same, directly advertised or marketed the products to consumers and end users, including the Plaintiffs, and persons responsible for consumers (such as employers), and therefore had a duty to warn of the risks associated with the use of Roundup® and glyphosate-containing products.

106. At all times relevant to this litigation, Defendants had a duty to properly test, develop, design, manufacture, inspect, package, label, market, promote, sell, distribute, maintain supply, provide proper warnings, and take such steps as necessary to ensure that Roundup® products did not cause users and consumers to suffer from unreasonable and dangerous risks. Defendants had a continuing duty to warn the Plaintiffs of the dangers associated with Roundup® use and exposure. Defendants, as manufacturer, seller, or distributor of chemical herbicides, are held to the knowledge of an expert in the field.

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107. At the time of manufacture, Defendants could have provided the warnings or instructions regarding the full and complete risks of Roundup® and glyphosate-containing products because they knew or should have known of the unreasonable risks of harm associated with the use of and/or exposure to such products.

At all times relevant to this litigation, Defendants failed to investigate, study, test, or promote the safety or to minimize the dangers to users and consumers of this product and to those who would foreseeably use or be harmed by Roundup, including Plaintiffs.

109. Despite the fact that Defendants knew or should have known that Roundup® posed a grave risk of harm, they failed to exercise reasonable care to warn of the dangerous risks associated with use and exposure. The dangerous propensities of its products and the carcinogenic characteristics of glyphosate, as described above, were known to Defendants, or scientifically knowable to Defendants through appropriate research and testing by known methods, at the time it distributed, supplied or sold the product, and not known to end users and consumers, such as Plaintiffs.

110. Defendants knew or should have known that these products created significant risks of serious bodily harm to consumers, as alleged herein, and Defendants failed to adequately warn consumers and reasonably foreseeable users of the risks of exposure to its products. Defendants have wrongfully concealed information concerning the dangerous nature of Roundup® and its

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active ingredient glyphosate, and further made false and/or misleading statements concerning the safety of Roundup® and glyphosate.

At all times relevant to this litigation, Defendants' Roundup® products reached the 111. intended consumers, handlers, and users or other persons coming into contact with these products in California and throughout the United States, including Plaintiffs, without substantial change in their condition as designed, manufactured, sold, distributed, labeled, marketed and sprayed/applied by Defendants.

- 112. Plaintiffs were exposed to Roundup® products, as described above, without knowledge of their dangerous characteristics.
- At all times relevant to this litigation, Plaintiffs were exposed to the use of Defendants' 113. Roundup® products in their intended or reasonably foreseeable manner without knowledge of their dangerous characteristics.
- Plaintiffs could not have reasonably discovered the defects and risks associated with Roundup® or glyphosate-containing products prior to or at the time of Plaintiffs' exposure. Plaintiffs relied upon the skill, superior knowledge, and judgment of Defendants.
- 115. Defendants knew or should have known that the minimal warnings disseminated with or accompanying the application of Roundup® products were inadequate, but they failed to COMPLAINT FOR DAMAGES AND DEMAND FOR JURY TRIAL

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communicate adequate information on the dangers and safe use/exposure and failed to communicate warnings and instructions that were appropriate and adequate to render the products safe for their ordinary, intended and reasonably foreseeable uses, including agricultural and horticultural applications.

The information that Defendants did provide or communicate failed to contain relevant

warnings, hazards, and precautions that would have enabled those exposed such as Plaintiffs to utilize the products safely and with adequate protection. Instead, Defendants disseminated information that was inaccurate, false, and misleading and which failed to communicate accurately or adequately the comparative severity, duration, and extent of the risk of injuries with use of and/or exposure to Roundup® and glyphosate; continued to aggressively promote the efficacy of its products, even after it knew or should have known of the unreasonable risks from use or exposure; and concealed, downplayed, or otherwise suppressed, through aggressive marketing and promotion, any information or research about the risks and dangers of exposure to Roundup® and glyphosate.

117. To this day, Defendants have failed to adequately and accurately warn of the true risks of Plaintiffs' injuries associated with the use of and exposure to Roundup® and its active ingredient glyphosate, a probable carcinogen.

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118. As a result of their inadequate warnings, Roundup® products were defective and unreasonably dangerous when they left the possession and/or control of Defendants, were sold or distributed by Defendants, were applied by Defendants, and when Plaintiffs became exposed.

- 119. Defendants are liable to Plaintiffs for injuries caused by negligent or willful failure, as described above, to provide adequate warnings or other clinically relevant information and data regarding the appropriate use of their products and the risks associated with the use of or exposure to Roundup® and glyphosate.
- 120. The defects in these Roundup® products were substantial and contributing factors in causing Plaintiffs' injuries, and, but for Defendants' misconduct and omissions, Plaintiffs would not have sustained their injuries.
- 121. Had Defendants provided adequate warnings and instructions and properly disclosed and disseminated the risks associated with Roundup® products and application, Plaintiffs could have avoided the risk of developing injuries as alleged herein and the company who employed Plaintiffs could have obtained alternative herbicides.
- 122. As a direct and proximate result of Defendants placing defective Roundup® products into the stream of commerce and exposing Plaintiffs to them, Plaintiffs have suffered and continue to suffer severe injuries, and has endured physical pain and discomfort, as well as economic

will continue to incur these expenses in the future.

hardship, including considerable financial expenses for medical care and treatment. Plaintiffs

123. WHEREFORE, Plaintiffs respectfully request that this Court enter judgment in Plaintiffs' favor for compensatory and punitive damages, together with interest, costs herein incurred, attorneys' fees and all such other and further relief as this Court deems just and proper. Plaintiffs also demand a jury trial on the issues contained herein.

CLAIM THREE

NEGLIGENCE

(Against All Defendants)

- 124. Plaintiffs incorporate by reference each and every allegation set forth in the preceding paragraphs as if fully stated herein.
- 125. Defendants, directly or indirectly, caused Roundup® products to be sold, distributed, packaged, labeled, marketed, promoted, and/or used by Plaintiffs.
- 126. At all times relevant to this litigation, Defendants had a duty to exercise reasonable care in the design, research, manufacture, marketing, advertisement, supply, promotion, packaging, sale, and distribution of Roundup® products, including the duty to take all reasonable steps necessary to manufacture, promote, and/or sell a product that was not unreasonably dangerous to consumers and users of the product.

COMPLAINT FOR DAMAGES AND DEMAND FOR JURY TRIAL

 127. At all times relevant to this litigation, Defendants had a duty to exercise reasonable care in the marketing, advertisement, and sale of the Roundup® products. Defendants' duty of care owed to consumers and the general public included providing accurate, true, and correct information concerning the risks of using Roundup® and appropriate, complete, and accurate warnings concerning the potential adverse effects of exposure to Roundup®, and, in particular, its active ingredient glyphosate.

- 128. At all times relevant to this litigation, Defendants knew or, in the exercise of reasonable care, should have known of the hazards and dangers of Roundup® and specifically, the carcinogenic properties of the chemical glyphosate.
- 129. Accordingly, at all times relevant to this litigation, Defendants knew or, in the exercise of reasonable care, should have known that use of or exposure to Roundup® products could cause or be associated with Plaintiffs' injuries and thus created a dangerous and unreasonable risk of injury to the users of these products, including Plaintiffs.
- 130. Defendants also knew or, in the exercise of reasonable care, should have known that users and consumers of Roundup® were unaware of the risks and the magnitude of the risks associated with use of and/or exposure to Roundup® and glyphosate-containing products.

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 131. As such, Defendants breached their duty of reasonable care and failed to exercise ordinary care in the design, research, development, manufacture, testing, marketing, supply, promotion, advertisement, packaging, sale, and distribution of its Roundup® products, in that Defendants manufactured and produced defective herbicides containing the chemical glyphosate, knew or had reason to know of the defects inherent in its products, knew or had reason to know that a user's or consumer's exposure to the products created a significant risk of harm and unreasonably dangerous side effects, and failed to prevent or adequately warn of these risks and injuries.

- 132. Despite ability and means to investigate, study, and test products and to provide adequate warnings, Defendants have failed to do so. Indeed, Defendants wrongfully concealed information and has further made false and/or misleading statements concerning the safety and/or exposure to Roundup® and glyphosate.
- 133. Defendants' negligence included, but are not limited to:
- a. Manufacturing, producing, promoting, formulating, creating, developing, designing, selling, and/or distributing Roundup® products without thorough and adequate pre- and post-market testing;
- b. Manufacturing, producing, promoting, formulating, creating, developing, designing, selling, and/or distributing Roundup® while negligently and/or intentionally concealing and failing to disclose the results of trials, tests, and studies of exposure to glyphosate, and, consequently, the risk of serious harm associated with human use of and exposure to Roundup®;

- c. Failing to undertake sufficient studies and conduct necessary tests to determine whether or not Roundup® products and glyphosate-containing products were safe for their intended use in agriculture and horticulture;
- d. Failing to use reasonable and prudent care in the design, research, manufacture, and development of Roundup® products so as to avoid the risk of serious harm associated with the prevalent use of Roundup®/glyphosate as an herbicide;
- e. Failing to design and manufacture Roundup® products so as to ensure they were at least as safe and effective as other herbicides on the market;
- f. Failing to provide adequate instructions, guidelines, and safety precautions to those persons who Defendants could reasonably foresee would use and be exposed to its Roundup® products;
- g. Failing to disclose to Plaintiffs, users/consumers, and the general public that use of and exposure to Roundup® presented severe risks of cancer and other grave illnesses;
- h. Failing to warn Plaintiffs, consumers, and the general public that the product's risk of harm was unreasonable and that there were safer and effective alternative herbicides available to Plaintiffs and other consumers;
- i. Systematically suppressing or downplaying contrary evidence about the risks, incidence, and prevalence of the side effects of Roundup® and glyphosate-containing products;
- j. Representing that its Roundup® products were safe for their intended use when, in fact,

 Defendants knew or should have known that the products were not safe for their intended

 purpose;

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k. Declining to make or propose any changes to Roundup® products' labeling or other promotional materials that would alert the consumers and the general public of the risks of Roundup® and glyphosate;

- 1. Advertising, marketing, and recommending the use of the Roundup® products, while concealing and failing to disclose or warn of the dangers known by Defendants to be associated with or caused by the use of or exposure to Roundup® and glyphosate;
- m. Continuing to disseminate information to its consumers, which indicate or imply that Defendants' Roundup® products are not unsafe for use in the agricultural and horticultural industries; and
- n. Continuing the manufacture and sale of its products with the knowledge that the products were unreasonably unsafe and dangerous.
- 134. Defendants knew and/or should have known that it was foreseeable that consumers such as Plaintiffs would suffer injuries as a result of Defendants' failure to exercise ordinary care in the manufacturing, marketing, labeling, distribution, and sale of Roundup®.
- 135. Plaintiffs did not know the nature and extent of the injuries that could result from the intended use of and/or exposure to Roundup® or its active ingredient glyphosate.
- 136. Defendants' negligence was the proximate cause of the injuries, harm, and economic losses that Plaintiffs suffered, and will continue to suffer, as described herein.

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Defendants' conduct, as described above, was reckless. Defendants regularly risk the 137. lives of consumers and users of their products, including Plaintiffs, with full knowledge of the dangers of its products. Defendants made conscious decisions not to redesign, re-label, warn, or inform the unsuspecting public, including Plaintiffs. Defendants' reckless conduct therefore warrants an award of punitive damages.

138. As a proximate result of Defendants' wrongful acts and omissions in placing defective Roundup® products into the stream of commerce without adequate warnings of the hazardous and carcinogenic nature of glyphosate, Plaintiffs have suffered and continues to suffer severe and permanent physical and emotional injuries. Plaintiffs have endured pain and suffering, have suffered economic losses (including significant expenses for medical care and treatment) and will continue to incur these expenses in the future.

139. WHEREFORE, Plaintiffs respectfully requests that this Court enter judgment in Plaintiffs' favor for compensatory and punitive damages, together with interest, costs herein incurred, attorneys' fees and all such other and further relief as this Court deems just and proper. Plaintiffs also demands a jury trial on the issues contained herein.

CLAIM FOUR

BREACH OF IMPLIED WARRANTIES

(Against All Defendants)

140. Plaintiffs incorporate by reference each and every allegation set forth in the preceding paragraphs as if fully stated herein.

COMPLAINT FOR DAMAGES AND DEMAND FOR JURY TRIAL

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- 141. At all times relevant to this litigation, Defendants engaged in the business of testing, developing, designing, manufacturing, marketing, selling, distributing, and promoting Roundup® products, which are defective and unreasonably dangerous to consumers, including Plaintiffs, thereby placing Roundup® products into the stream of commerce. These actions were under the ultimate control and supervision of Defendants.
- 142. Before the time that Plaintiffs were exposed to the use of the aforementioned Roundup® products, Defendants impliedly warranted to consumers and those exposed—including Plaintiffs—that Roundup® products were of merchantable quality and safe and fit for the use for which they were intended; specifically, as horticultural herbicides.
- 143. Defendants, however, failed to disclose that Roundup® has dangerous propensities when used as intended and that the use of and/or exposure to Roundup® and glyphosate-containing products carries an increased risk of developing severe injuries, including Plaintiffs' injuries.
- 144. Plaintiffs reasonably relied upon the skill, superior knowledge and judgment of

 Defendants and upon their implied warranties that the Roundup® products were of merchantable
 quality and fit for their intended purpose or use.
- 145. Upon information and belief, Plaintiffs were at all relevant times in privity with Defendants.

146. Plaintiffs are the intended third-party beneficiaries of implied warranties made by Defendants to the purchasers and/or users of their horticultural herbicides, and as such Plaintiffs are entitled to assert this claim.

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147. The Roundup® products were expected to reach and did in fact reach consumers and/or users, including Plaintiffs, without substantial change in the condition in which they were manufactured and sold by Defendants.

At all times relevant to this litigation, Defendants were aware that consumers and users of their products, including Plaintiffs, would use Roundup® products as marketed by Defendants, which is to say that Plaintiffs were foreseeable users of Roundup®.

149. Defendants intended that Roundup® products be used in the manner in which Plaintiffs were exposed to it and Defendants impliedly warranted each product to be of merchantable quality, safe, and fit for this use, despite the fact that Roundup® was not adequately tested and/or researched.

150. In reliance upon Defendants' implied warranty, Plaintiffs used or were exposed to Roundup® as instructed and labeled and in the foreseeable manner intended, recommended, promoted and marketed by Defendants.

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1	151. Plaintiffs could not have reasonably discovered or known of the risks of serious injury			
2	associated with Roundup® or glyphosate.			
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5	152. Defendants breached their implied warranty to Plaintiffs in that Roundup® products were			
6	not of merchantable quality, safe, or fit for their intended use, and/or adequately tested.			
7	Roundup® has dangerous propensities when used as intended and can cause serious injuries,			
8	including those injuries complained of herein.			
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12	153. The harm caused by Roundup® products far outweighed their benefit, rendering the			
13	products more dangerous than an ordinary consumer or user would expect and more dangerous			
14	than alternative products.			
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17	154. As a direct and proximate result of Defendants' wrongful acts and omissions Plaintiffs			
18	have suffered severe and permanent physical and emotional injuries, including but not limited to			
19	their diagnoses of non-Hodgkin lymphoma. Plaintiffs have endured pain and suffering, have			
20	suffered economic loss (including significant expenses for medical care and treatment) and will			
21	continue to incur these expenses in the future.			
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24	155. WHEREFORE, Plaintiffs respectfully requests that this Court enter judgment in Plaintiffs'			
25	favor for compensatory and punitive damages, together with interest, costs herein incurred,			
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ain and suffering, have are and treatment) and will enter judgment in Plaintiffs' costs herein incurred, attorneys' fees, and all such other and further relief as this Court deems just and proper. Plaintiffs also demand a jury trial on the issues contained herein. COMPLAINT FOR DAMAGES AND DEMAND FOR JURY TRIAL 43

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COUNT FIVE

PUNITIVE DAMAGES

- 156. Plaintiffs repeat and reiterate the allegations previously set forth herein.
- 157. At all times material hereto, the Defendants knew or should have known that the subject product was inherently dangerous with respect to its health risks
- 158. At all times material hereto, the Defendants attempted to misrepresent and did misrepresent facts concerning the safety of the subject product.
- 159. Defendants' misrepresentations included knowingly withholding material information from the public, including the Plaintiffs herein, concerning the safety of the subject product.
- 160. At all times material hereto, the Defendants knew and recklessly disregarded the fact that human exposure to Roundup can and does cause health hazard, including non Hodgkin lymphoma.
- 161. Notwithstanding the foregoing, the Defendants continued to aggressively market and apply the subject product without disclosing the aforesaid risks.
- 162. Defendants knew of the subject product's defective and unreasonably dangerous nature, as set forth herein, but continued to design, develop, manufacture, market, distribute, sell, and COMPLAINT FOR DAMAGES AND DEMAND FOR JURY TRIAL

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apply it so as to maximize sales and profits at the expense of the health and safety of the public, including the Plaintiffs herein, in conscious and/or negligent disregard of the foreseeable harm caused by Roundup.

163. The Defendants intentionally concealed and/or recklessly failed to disclose to the public, including the Plaintiffs herein, the potentially life threatening hazards of Roundup in order to ensure continued and increased sales.

164. The Defendants' intentional and/or reckless failure to disclose information deprived the Plaintiffs of necessary information to enable Plaintiffs to weigh the true risks of using or being exposed to the subject product against its benefits.

165. As a direct and proximate result of the Defendants' conscious and deliberate disregard for the rights and safety of consumers such as the Plaintiffs, Plaintiffs suffered severe and permanent physical injuries. The Plaintiffs have endured substantial pain and suffering and has undergone extensive medical and surgical procedures. Plaintiffs have incurred significant expenses for medical care and treatment, and will continue to incur such expenses in the future. The Plaintiffs have lost past earnings and have suffered a loss of earning capacity. The Plaintiffs have suffered and will continue to suffer economic loss, and has otherwise been physically, emotionally and economically injured. The Plaintiffs' injuries and damages are permanent and will continue into the future.

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The aforesaid conduct of the Defendants was committed with knowing, conscious, and 166. deliberate disregard for the rights and safety of consumers, including the Plaintiffs herein, thereby entitling the Plaintiffs to punitive damages in an amount appropriate to punish the Defendants and deter them from similar conduct in the future.

WHEREFORE, Plaintiffs demand judgment against Defendants for compensatory, treble, 167. and punitive damages, together with interest, costs of suit, attorneys' fees, and all such other relief as the Court deems proper.

PRAYER FOR RELIEF

WHEREFORE, Plaintiffs request that the Court enter judgment in their favor and against Defendants, awarding as follows:

- 1. Judgment for Plaintiffs and against Defendants;
- 2. For compensatory damages in an amount to be proven at trial;
- 3. For mental and physical suffering, according to proof;
- 4. For lost wages according to proof;
- 5. For disgorgement of profits, according to proof;
- 6. For punitive damages;
- 7. For default judgment as a sanction for the bad faith destruction of evidence, if any, and according to proof, if any;
- 8. For costs including reasonable attorneys' fees, court costs, and other litigation expenses; and

COMPLAINT FOR DAMAGES AND DEMAND FOR JURY TRIAL

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9. For any other relief the Court may deem just and proper.

DEMAND FOR JURY TRIAL

Plaintiffs hereby demand a jury trial on all claims so triable in this action.

Dated: June 2, 2017

Respectfully submitted,

THE MILLER FIRM, LLC

Curtis G. Hoke (SBN 282465)

THE MILLER FIRM LLC

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Attorneys for Plaintiffs

COMPLAINT FOR DAMAGES AND DEMAND FOR JURY TRIAL

EXHIBIT 2

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13	ALVA AND ALBERTA PILLIOD	JCCP No.: 4953	
14	Plaintiffs,	Case No.: RG17862702	
15 16 17 18 19 20 21 22 23 24	vs. MONSANTO COMPANY; WILBUR-ELLIS COMPANY, LLC; an WILBUR-ELLIS FEED, LLC, Defendants.	FIRST AMENDED COMPLAINT FOR DAMAGES AND DEMAND FOR JURY TRIAL 1. Strict Liability – Design Defect 2. Strict Liability – Failure to Warn 3. Negligence 4. Breach of Implied Warranty 5. Punitive Damage 6. Loss of Consortium JURY TRIAL DEMANDED	
25 26 27			
28	COMPLAINT FOR DAMAGES AND DEMAND FOR JURY TRIAL 1		

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COMPLAINT AND DEMAND FOR JURY TRIAL

The above-captioned Plaintiffs (hereinafter, "Plaintiffs"), by attorneys, THE MILLER FIRM, LLC, as and for the Complaint herein allege upon information and belief the following:

SUMMARY OF THE CASE

- 1. In 1970, defendant Monsanto Company, Inc. discovered the herbicidal properties of glyphosate and began marketing it in products in 1974 under the brand name Roundup®. Roundup® is a non-selective herbicide used to kill weeds that commonly compete with the growing of crops. By 2001, glyphosate had become the most-used active ingredient in American agriculture with 85–90 millions of pounds used annually. That number grew to 185 million pounds by 2007. As of 2013, glyphosate was the world's most widely used herbicide.
- 2. Monsanto is a multinational agricultural biotechnology corporation based in St. Louis, Missouri. It is the world's leading producer of glyphosate. As of 2009, Monsanto was the world's leading producer of seeds, accounting for 27% of the world seed market. The majority of these seeds are of the Roundup Ready® brand. The stated advantage of Roundup Ready® crops is that they substantially improve a farmer's ability to control weeds, since glyphosate can be sprayed in the fields during the growing season without harming their crops. In 2010, an estimated 70% of corn and cotton, and 90% of soybean fields in the United States were Roundup Ready®.
- 3. Monsanto's glyphosate products are registered in 130 countries and approved for use on over 100 different crops. They are ubiquitous in the environment. Numerous studies confirm that glyphosate is found in rivers, streams, and groundwater in agricultural areas where Roundup® is COMPLAINT FOR DAMAGES AND DEMAND FOR JURY TRIAL

used. It has been found in food, in the urine of agricultural workers, and even in the urine of urban dwellers who are not in direct contact with glyphosate.

- 4. On March 20, 2015, the International Agency for Research on Cancer ("IARC"), an agency of the World Health Organization ("WHO"), issued an evaluation of several herbicides, including glyphosate. That evaluation was based, in part, on studies of exposures to glyphosate in several countries around the world, and it traces the health implications from exposure to glyphosate since 2001.
- 5. On July 29, 2015, IARC issued the formal monograph relating to glyphosate. In that monograph, the IARC Working Group provides a thorough review of the numerous studies and data relating to glyphosate exposure in humans.
- 6. The IARC Working Group classified glyphosate as a Group 2A herbicide, which means that it is probably carcinogenic to humans. The IARC Working Group concluded that the cancers most associated with glyphosate exposure are non-Hodgkin lymphoma and other haematopoietic cancers, including lymphocytic lymphoma/chronic lymphocytic leukemia, B-cell lymphoma, and multiple myeloma.
- 7. The IARC evaluation is significant. It confirms what has been believed for years: that glyphosate is toxic to humans.
- 8. Nevertheless, Monsanto, since it began selling Roundup®, has represented it as safe to COMPLAINT FOR DAMAGES AND DEMAND FOR JURY TRIAL

humans and the environment. Indeed, Monsanto has repeatedly proclaimed and continues to proclaim to the world, and particularly to United States consumers, that glyphosate-based herbicides, including Roundup®, create no unreasonable risks to human health or to the environment.

JURISDICTION AND VENUE

- 9. The California Superior Court has jurisdiction over this action pursuant to California Constitution Article VI, Section 10, which grants the Superior Court "original jurisdiction in all causes except those given by statute to other trial courts." The Statutes under which this action is brought do not specify any other basis for jurisdiction.
- 10. The California Superior Court has jurisdiction over the Defendants because, based on information and belief, each is a California resident, a corporation and/or entity organized under the laws of the State of California, a foreign corporation or association authorized to do business in California and registered with the California Secretary of State or has sufficient minimum contacts in California, or otherwise intentionally avails itself of the California market so as to render the exercise of jurisdiction over it by the California courts consistent with traditional notions of fair play and substantial justice.
- 11. Venue is proper in this Court pursuant to California Code of Civil Procedure Section 395 in that the subject injury occurred in Alameda County.

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12. Furthermore the Defendants have purposefully availed themselves of the benefits and the protections of the laws within the State of California. Monsanto has had sufficient contact such that the exercise of jurisdiction would be consistent with the traditional notions of fair play and substantial justice.

13. Plaintiffs seek relief that is within the jurisdictional limits of this Court.

THE PARTIES

Plaintiffs

14. Plaintiff ALVA PILLIOD is a competent individual over the age of 18, a resident and citizen of the United States, and hereby submits to the jurisdiction of the Court and alleges that jurisdiction and venue in this court is proper. Mr. Pilliod currently resides in Livermore, County of Alameda, California. Mr. Pilliod used Monsanto's glyphosate-containing Roundup® product and suffered from severe physical, economy, and emotional injuries as a result of his use of Roundup® between 1975 and 2011, including but not limited to non-Hodgkin lymphoma, diagnosed in 2011 in California. Plaintiff ALBERTA PILLIOD is a competent individual over the age of 18, a resident and citizen of the United States, and hereby submits to the jurisdiction of the Court and alleges that jurisdiction and venue in this court is proper. Mrs. Pilliod currently resides in Livermore, County of Alameda, California. Mrs. Pilliod used the same Monsanto manufactured glyphosate-containing Roundup® products as Mr. Pilliod and suffered from severe physical, economic, and emotional injuries as a result of her use of Roundup® between 1975 and 2011, including but not limited to non-Hodgkin lymphoma, diagnosed in April, 2015 in

California. Mr. and Mrs. Pilliod are lawfully married spouses and each seek damages for loss of consortium herein.

15. Plaintiff CHARLES BAKER is a competent individual over the age of 18, a resident and

citizen of the United States, and hereby submits to the jurisdiction of the Court and alleges that jurisdiction and venue in this court is proper. Mr. Baker currently resides in Raymond, California. Mr. Baker used Monsanto's glyphosate containing Roundup® product at work and at home from 1995 to 2016 and suffered from severe physical, economy, and emotional injuries as a result of his use of Roundup®, including but not limited to non-Hodgkin lymphoma, diagnosed in June, 2015 in California.

16. Plaintiff JOHN NOVAK is a competent individual over the age of 18, a resident and eitizen of the United States, and hereby submits to the jurisdiction of the Court and alleges that jurisdiction and venue in this court is proper. Mr. Novak currently resides in Vista, California. Mr. Novak used Monsanto's glyphosate containing Roundup® product at work and at home from 2002 to 2010 and suffered from severe physical, economy, and emotional injuries as a result of his use of Roundup®, including but not limited to non-Hodgkin lymphoma, diagnosed in 2011 in California.

17. Plaintiff SHARON ROWLAND is a competent individual over the age of 18, a resident and citizen of the United States, and hereby submits to the jurisdiction of the Court and alleges that jurisdiction and venue in this court is proper. Ms. Rowland currently resides in Fontana, California. Ms. Rowland used Monsanto's glyphosate containing Roundup® product from 1985 to 2015 on her 10 acres and suffered from severe physical, economy, and emotional injuries as a result of his use of Roundup®, including but not limited to non Hodgkin lymphoma, diagnosed in June, 2015 in California.

COMPLAINT FOR DAMAGES AND DEMAND FOR JURY TRIAL

18. Plaintiff SHARON MCCLURG is a competent individual over the age of 18, a resident and citizen of the United States, and hereby submits to the jurisdiction of the Court and alleges that jurisdiction and venue in this court is proper. Ms. McClurg currently resides in Roseville, California. Ms. McClurg used Monsanto's glyphosate containing Roundup® product at home from 1974 to 2016 and suffered from severe physical, economy, and emotional injuries as a result of his use of Roundup®, including but not limited to non-Hodgkin lymphoma, diagnosed in 2003 in California.

and citizen of the United States, and hereby submits to the jurisdiction of the Court and alleges that jurisdiction and venue in this court is proper. Ms. Grubka currently resides in Martinez, California. Mr. Cole used Monsanto's glyphosate containing Roundup® product at home from 1986 to 2016 and suffered from severe physical, economy, and emotional injuries as a result of his use of Roundup®, including but not limited to non Hodgkin lymphoma, diagnosed in December 2016 in California.

Plaintiff MARJORIE GRUBKA is a competent individual over the age of 18, a resident

Defendants

15. Defendant Monsanto Company ("Monsanto") is a Delaware corporation with its headquarters and principal place of business in St. Louis, Missouri. At all times relevant to this complaint, Monsanto was the entity that discovered the herbicidal properties of glyphosate and the manufacturer of Roundup®. Monsanto has regularly transacted and conducted business within the state of California, and has derived substantial revenue from goods and products, including Roundup, used in the State of California. Monsanto expected or should have expected

their acts to have consequences within the State of California, and derived substantial revenue from interstate commerce.

16. Defendants Wilbur-Ellis Company LLC is a California limited liability corporation with its headquarters and principal place of business in San Francisco, California. At all times relevant to this complaint, Wilbur-Ellis Company, LLC sold and distributed Monsanto products including Roundup, within the State of California.

17. Defendants Wilbur-Ellis Feed LLC (with Wilbur-Ellis Company LLC, hereinafter "Wilbur-Ellis") is a California limited liability corporation with its headquarters and principal place of business in San Francisco, California. At all times relevant to this complaint, Wilbur-Ellis Feed, LLC sold and distributed Monsanto products including Roundup, within the State of California.

- 18. Defendants Monsanto Company, Wilbur-Ellis Company, LLC, and Wilbur-Ellis Feed, LLC are collectively referred to throughout this Complaint as "Defendants."
- 19. Plaintiffs are informed and believe, and based thereon allege, that in committing the acts alleged herein, each and every managing agent, agent, representative and/or employee of the Defendants was working within the course and scope of said agency, representation and/or employment with the knowledge, consent, ratification, and authorization of the Defendants and their directors, officers and/or managing agents.

CASE FACTS

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20. Glyphosate is a broad-spectrum, non-selective herbicide used in a wide variety of herbicidal products around the world.

21. Plants treated with glyphosate translocate the systemic herbicide to their roots, shoot regions and fruit, where it interferes with the plant's ability to form aromatic amino acids necessary for protein synthesis.

22. Treated plants generally die within two to three days. Because plants absorb glyphosate, it cannot be completely removed by washing or peeling produce or by milling, baking, or brewing grains.

23. For nearly 40 years, farms across the world have used Roundup® without knowing of the dangers its use poses. That is because when Monsanto first introduced Roundup®, it touted glyphosate as a technological breakthrough: it could kill almost every weed without causing harm either to people or to the environment. Of course, history has shown that not to be true. According to the WHO, the main chemical ingredient of Roundup®—glyphosate—is a probable cause of cancer. Those most at risk are farm workers and other individuals with workplace exposure to Roundup®, such as workers in garden centers, nurseries, and landscapers. Agricultural workers are, once again, victims of corporate greed.

24. Monsanto assured the public that Roundup® was harmless. In order to prove this,

Monsanto championed falsified data and attacked legitimate studies that revealed its dangers.

COMPLAINT FOR DAMAGES AND DEMAND FOR JURY TRIAL

Monsanto led a prolonged campaign of misinformation to convince government agencies, farmers and the general population that Roundup® was safe.

The Discovery of Glyphosate and Development of Roundup®

25. The herbicidal properties of glyphosate were discovered in 1970 by Monsanto chemist John Franz. The first glyphosate-based herbicide was introduced to the market in the mid-1970s under the brand name Roundup®. From the outset, Monsanto marketed Roundup® as a "safe" general-purpose herbicide for widespread commercial and consumer use. It still markets Roundup® as safe today.

Registration of Herbicides under Federal Law

- 26. The manufacture, formulation and distribution of herbicides, such as Roundup®, are regulated under the Federal Insecticide, Fungicide, and Rodenticide Act ("FIFRA" or "Act"), 7 U.S.C. § 136 *et seq.* FIFRA requires that all pesticides be registered with the Environmental Protection Agency ("EPA" or "Agency") prior to their distribution, sale, or use, except as described by the Act. 7 U.S.C. § 136a(a)
- 27. Because pesticides are toxic to plants, animals, and humans, at least to some degree, the EPA requires as part of the registration process, among other things, a variety of tests to evaluate the potential for exposure to pesticides, toxicity to people and other potential non-target organisms, and other adverse effects on the environment. Registration by the EPA, however, is not an assurance or finding of safety. The determination the Agency must make in registering or re-registering a product is not that the product is "safe," but rather that use of the product in accordance with its label directions "will not generally cause unreasonable adverse effects on the environment." 7 U.S.C. § 136a(c)(5)(D).

 COMPLAINT FOR DAMAGES AND DEMAND FOR JURY TRIAL

28. FIFRA defines "unreasonable adverse effects on the environment" to mean "any unreasonable risk to man or the environment, taking into account the economic, social, and environmental costs and benefits of the use of any pesticide." 7 U.S.C. § 136(bb). FIFRA thus requires EPA to make a risk/benefit analysis in determining whether a registration should be granted or allowed to continue to be sold in commerce.

29. The EPA and the State of California registered Roundup® for distribution, sale, and manufacture in the United States and the State of California.

30. FIFRA generally requires that the registrant, Monsanto in the case of Roundup®, conducts the health and safety testing of pesticide products. The EPA has protocols governing the conduct of tests required for registration and the laboratory practices that must be followed in conducting these tests. The data produced by the registrant must be submitted to the EPA for review and evaluation. The government is not required, nor is it able, however, to perform the product tests that are required of the manufacturer.

31. The evaluation of each pesticide product distributed, sold, or manufactured is completed at the time the product is initially registered. The data necessary for registration of a pesticide has changed over time. The EPA is now in the process of re-evaluating all pesticide products through a Congressionally-mandated process called "re-registration." 7 U.S.C. § 136a-1. In order to

reevaluate these pesticides, the EPA is demanding the completion of additional tests and the submission of data for the EPA's review and evaluation.

32. In the case of glyphosate, and therefore Roundup®, the EPA had planned on releasing its preliminary risk assessment —in relation to the reregistration process—no later than July 2015. The EPA completed its review of glyphosate in early 2015, but it delayed releasing the risk assessment pending further review in light of the WHO's health-related findings.

Scientific Fraud Underlying the Marketing and Sale of Glyphosate/Roundup

- 33. Based on early studies that glyphosate could cause cancer in laboratory animals, the EPA originally classified glyphosate as *possibly carcinogenic to humans* (Group C) in 1985. After pressure from Monsanto, including contrary studies it provided to the EPA, the EPA changed its classification to *evidence of non-carcinogenicity in humans* (Group E) in 1991. In so classifying glyphosate, however, the EPA made clear that the designation did not mean the chemical does not cause cancer: "It should be emphasized, however, that designation of an agent in Group E is based on the available evidence at the time of evaluation and should not be interpreted as a definitive conclusion that the agent will not be a carcinogen under any circumstances."
- 34. On two occasions, the EPA found that the laboratories hired by Monsanto to test the toxicity of its Roundup® products for registration purposes committed fraud.

35. In the first instance, Monsanto, in seeking initial registration of Roundup® by EPA, hired Industrial Bio-Test Laboratories ("IBT") to perform and evaluate pesticide toxicology studies relating to Roundup®. IBT performed about 30 tests on glyphosate and glyphosate-containing products, including nine of the 15 residue studies needed to register Roundup®.

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36. In 1976, the United States Food and Drug Administration ("FDA") performed an inspection of Industrial Bio-Test Industries ("IBT") that revealed discrepancies between the raw data and the final report relating to the toxicological impacts of glyphosate. The EPA subsequently audited IBT; it too found the toxicology studies conducted for the Roundup® herbicide to be invalid. An EPA reviewer stated, after finding "routine falsification of data" at IBT, that it was "hard to believe the scientific integrity of the studies when they said they took specimens of the uterus from male rabbits."

- 37. Three top executives of IBT were convicted of fraud in 1983.
- 38. In the second incident of data falsification, Monsanto hired Craven Laboratories in 1991 to perform pesticide and herbicide studies, including for Roundup®. In that same year, the owner of Craven Laboratories and three of its employees were indicted, and later convicted, of fraudulent laboratory practices in the testing of pesticides and herbicides.
- 39. Despite the falsity of the tests that underlie its registration, within a few years of its launch, Monsanto was marketing Roundup® in 115 countries.

The Importance of Roundup® to Monsanto's Market Dominance Profits

- 40. The success of Roundup® was key to Monsanto's continued reputation and dominance in the marketplace. Largely due to the success of Roundup® sales, Monsanto's agriculture division was out-performing its chemicals division's operating income, and that gap increased yearly. But with its patent for glyphosate expiring in the United States in the year 2000, Monsanto needed a strategy to maintain its Roundup® market dominance and to ward off impending competition.
- A1. In response, Monsanto began the development and sale of genetically engineered Roundup Ready® seeds in 1996. Since Roundup Ready® crops are resistant to glyphosate; farmers can spray Roundup® onto their fields during the growing season without harming the crop. This allowed Monsanto to expand its market for Roundup® even further; by 2000, Monsanto's biotechnology seeds were planted on more than 80 million acres worldwide and nearly 70% of American soybeans were planted from Roundup Ready® seeds. It also secured Monsanto's dominant share of the glyphosate/Roundup® market through a marketing strategy that coupled proprietary Roundup Ready® seeds with continued sales of its Roundup® herbicide.
- 42. Through a three-pronged strategy of increased production, decreased prices and by coupling with Roundup Ready® seeds, Roundup® became Monsanto's most profitable product. In 2000, Roundup® accounted for almost \$2.8 billion in sales, outselling other herbicides by a margin of five to one, and accounting for close to half of Monsanto's revenue. Today, glyphosate remains one of the world's largest herbicides by sales volume.

Monsanto has known for decades that it falsely advertises the safety of Roundup®

- 43. In 1996, the New York Attorney General ("NYAG") filed a lawsuit against Monsanto based on its false and misleading advertising of Roundup ® products. Specifically, the lawsuit challenged Monsanto's general representations that its spray-on glyphosate-based herbicides, including Roundup®, were "safer than table salt" and "practically non-toxic" to mammals, birds, and fish. Among the representations the NYAG found deceptive and misleading about the human and environmental safety of Roundup® are the following:
- a) Remember that environmentally friendly Roundup herbicide is biodegradable. It won't build up in the soil so you can use Roundup with confidence along customers' driveways, sidewalks and fences ...
- b) And remember that Roundup is biodegradable and won't build up in the soil.

 That will give you the environmental confidence you need to use Roundup everywhere you've got a weed, brush, edging or trimming problem.
 - c) Roundup biodegrades into naturally occurring elements.
- d) Remember that versatile Roundup herbicide stays where you put it. That means there's no washing or leaching to harm customers' shrubs or other desirable vegetation.
- e) This non-residual herbicide will not wash or leach in the soil. It ... stays where you apply it.
- f) You can apply Accord with "confidence because it will stay where you put it" it bonds tightly to soil particles, preventing leaching. Then, soon after application, soil microorganisms biodegrade Accord into natural products.
 - g) Glyphosate is less toxic to rats than table salt following acute oral ingestion. COMPLAINT FOR DAMAGES AND DEMAND FOR JURY TRIAL

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	h) Glyphosate's safety margin is much greater than required. It has over a 1,000-
fold safety 1	margin in food and over a 700-fold safety margin for workers who manufacture it o
use it.	

- i) You can feel good about using herbicides by Monsanto. They carry a toxicity category rating of 'practically non-toxic' as it pertains to mammals, birds and fish.
- j) "Roundup can be used where kids and pets will play and breaks down into natural material." This ad depicts a person with his head in the ground and a pet dog standing in an area which has been treated with Roundup.
- 44. On November 19, 1996, Monsanto entered into an Assurance of Discontinuance with NYAG, in which Monsanto agreed, among other things, "to cease and desist from publishing or broadcasting any advertisements [in New York] that represent, directly or by implication" that:
- a) its glyphosate-containing pesticide products or any component thereof are safe, non-toxic, harmless or free from risk.

* * *

b) its glyphosate-containing pesticide products or any component thereof manufactured, formulated, distributed or sold by Monsanto are biodegradable

* * *

c) its glyphosate-containing pesticide products or any component thereof stay where they are applied under all circumstances and will not move through the environment by any means.

* * :

COMPLAINT FOR DAMAGES AND DEMAND FOR JURY TRIAL

d) its glyphosate-containing pesticide products or any component thereof are "good" for the environment or are "known for their environmental characteristics."

* * *

- e) glyphosate-containing pesticide products or any component thereof are safer or less toxic than common consumer products other than herbicides;
- f) its glyphosate-containing products or any component thereof might be classified as "practically non-toxic.
- 45. Monsanto did not alter its advertising in the same manner in any state other than New York, and on information and belief still has not done so today.
- 46. In 2009, France's highest court ruled that Monsanto had not told the truth about the safety of Roundup®. The French court affirmed an earlier judgement that Monsanto had falsely advertised its herbicide Roundup® as "biodegradable" and that it "left the soil clean."

Classifications and Assessments of Glyphosate

47. The IARC process for the classification of glyphosate followed the stringent procedures for the evaluation of a chemical agent. Over time, the IARC Monograph program has reviewed 980 agents. Of those reviewed, it has determined 116 agents to be Group 1 (Known Human Carcinogens); 73 agents to be Group 2A (Probable Human Carcinogens); 287 agents to be Group 2B (Possible Human Carcinogens); 503 agents to be Group 3 (Not Classified); and one agent to be Probably Not Carcinogenic.

COMPLAINT FOR DAMAGES AND DEMAND FOR JURY TRIAL

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48. The established procedure for IARC Monograph evaluations is described in the IARC Programme's Preamble. Evaluations are performed by panels of international experts, selected on the basis of their expertise and the absence of actual or apparent conflicts of interest.

49. One year before the Monograph meeting, the meeting is announced and there is a call both for data and for experts. Eight months before the Monograph meeting, the Working Group membership is selected and the sections of the Monograph are developed by the Working Group members. One month prior to the Monograph meeting, the call for data is closed and the various draft sections are distributed among Working Group members for review and comment. Finally, at the Monograph meeting, the Working Group finalizes review of all literature, evaluates the evidence in each category, and completes the overall evaluation. Within two weeks after the Monograph meeting, the summary of the Working Group findings are published in Lancet Oncology, and within a year after the meeting, the final Monograph is finalized and published.

50. In assessing an agent, the IARC Working Group reviews the following information: (a) human, experimental, and mechanistic data; (b) all pertinent epidemiological studies and cancer bioassays; and (c) representative mechanistic data. The studies must be publicly available and have sufficient detail for meaningful review, and reviewers cannot be associated with the underlying study.

51. In March 2015, IARC reassessed glyphosate. The summary published in *The Lancet Oncology* reported that glyphosate is a Group 2A agent and probably carcinogenic in humans.

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52. On July 29, 2015, IARC issued its Monograph for glyphosate, Monograph 112. For Volume 112, the volume that assessed glyphosate, a Working Group of 17 experts from 11 countries met at IARC from March 3–10, 2015, to assess the carcinogenicity of certain herbicides, including glyphosate. The March meeting culminated nearly a one-year review and preparation by the IARC Secretariat and the Working Group, including a comprehensive review of the latest available scientific evidence. According to published procedures, the Working Group considered "reports that have been published or accepted for publication in the openly available scientific literature" as well as "data from governmental reports that are publicly available."

- 53. The studies considered the following exposure groups: occupational exposure of farmers and tree nursery workers in the United States, forestry workers in Canada and Finland and municipal weed-control workers in the United Kingdom; and para-occupational exposure in farming families.
- 54. Glyphosate was identified as the second-most used household herbicide in the United States for weed control between 2001 and 2007 and the most heavily used herbicide in the world in 2012.

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55. Exposure pathways are identified as air (especially during spraying), water, and food. Community exposure to glyphosate is widespread and found in soil, air, surface water, and groundwater, as well as in food.

- 56. The assessment of the IARC Working Group identified several case control studies of occupational exposure in the United States, Canada, and Sweden. These studies show a human health concern from agricultural and other work-related exposure to glyphosate.
- 57. The IARC Working Group found an increased risk between exposure to glyphosate and non-Hodgkin lymphoma ("NHL") and several subtypes of NHL, and the increased risk persisted after adjustment for other pesticides.
- 58. The IARC Working Group also found that glyphosate caused DNA and chromosomal damage in human cells. One study in community residents reported increases in blood markers of chromosomal damage (micronuclei) after glyphosate formulations were sprayed.
- 59. In male CD-1 mice, glyphosate induced a positive trend in the incidence of a rare tumor, renal tubule carcinoma. A second study reported a positive trend for haemangiosarcoma in male mice. Glyphosate increased pancreatic islet-cell adenoma in male rats in two studies. A glyphosate formulation promoted skin tumors in an initiation-promotion study in mice.

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60. The IARC Working Group also noted that glyphosate has been detected in the urine of agricultural workers, indicating absorption. Soil microbes degrade glyphosate to aminomethylphosphoric acid (AMPA). Blood AMPA detection after exposure suggests intestinal microbial metabolism in humans.

61. The IARC Working Group further found that glyphosate and glyphosate formulations induced DNA and chromosomal damage in mammals, and in human and animal cells in utero.

62. The IARC Working Group also noted genotoxic, hormonal, and enzymatic effects in mammals exposed to glyphosate. Essentially, glyphosate inhibits the biosynthesis of aromatic amino acids, which leads to several metabolic disturbances, including the inhibition of protein and secondary product biosynthesis and general metabolic disruption.

63. The IARC Working Group also reviewed an Agricultural Health Study, consisting of a prospective cohort of 57,311 licensed pesticide applicators in Iowa and North Carolina. While this study differed from others in that it was based on a self-administered questionnaire, the results support an association between glyphosate exposure and Multiple Myeloma, Hairy Cell Leukemia (HCL), and Chronic Lymphocytic Leukemia (CLL), in addition to several other cancers.

Other Earlier Findings About Glyphosate's Dangers to Human Health

64. The EPA has a technical fact sheet, as part of its Drinking Water and Health, National Primary Drinking Water Regulations publication, relating to glyphosate. This technical fact sheet predates the IARC March 20, 2015, evaluation. The fact sheet describes the release patterns for glyphosate as follows:

Release Patterns

- 65. Glyphosate is released to the environment in its use as a herbicide for controlling woody and herbaceous weeds on forestry, right-of-way, cropped and non-cropped sites. These sites may be around water and in wetlands.
- 66. It may also be released to the environment during its manufacture, formulation, transport, storage, disposal and cleanup, and from spills. Since glyphosate is not a listed chemical in the Toxics Release Inventory, data on releases during its manufacture and handling are not available.
- 67. Occupational workers and home gardeners may be exposed to glyphosate by inhalation and dermal contact during spraying, mixing, and cleanup. They may also be exposed by touching soil and plants to which glyphosate was applied. Occupational exposure may also occur during glyphosate's manufacture, transport storage, and disposal.
- 68. In 1995, the Northwest Coalition for Alternatives to Pesticides reported that in California, the state with the most comprehensive program for reporting of pesticide-caused illness, glyphosate was the third most commonly-reported cause of pesticide illness among agricultural workers.

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Recent Worldwide Bans on Roundup®/Glyphosate

- 69. Several countries around the world have instituted bans on the sale of Roundup® and other glyphosate-containing herbicides, both before and since IARC first announced its assessment for glyphosate in March 2015, and more countries undoubtedly will follow suit in light of the as the dangers of the use of Roundup® are more widely known. The Netherlands issued a ban on all glyphosate-based herbicides in April 2014, including Roundup®, which takes effect by the end of 2015. In issuing the ban, the Dutch Parliament member who introduced the successful legislation stated: "Agricultural pesticides in user-friendly packaging are sold in abundance to private persons. In garden centers, Roundup® is promoted as harmless, but unsuspecting customers have no idea what the risks of this product are. Especially children are sensitive to toxic substances and should therefore not be exposed to it."
- 70. The Brazilian Public Prosecutor in the Federal District requested that the Brazilian Justice Department suspend the use of glyphosate.
- 71. France banned the private sale of Roundup® and glyphosate following the IARC assessment for Glyphosate.
- 72. Bermuda banned both the private and commercial sale of glyphosates, including Roundup®. The Bermuda government explained its ban as follows: "Following a recent scientific study carried out by a leading cancer agency, the importation of weed spray 'Roundup' has been suspended."

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73. The Sri Lankan government banned the private and commercial use of glyphosates, particularly out of concern that Glyphosate has been linked to fatal kidney disease in agricultural workers.

74. The government of Columbia announced its ban on using Roundup® and glyphosate to destroy illegal plantations of coca, the raw ingredient for cocaine, because of the WHO's finding that glyphosate is probably carcinogenic.

75. On information and belief, Wilbur-Ellis was, at all relevant times, engaged in the distribution of Roundup, Roundup-ready crops and other glyphosate-containing products from Monsanto to retailers and commercial/agricultural users in California.

76. Wilbur-Ellis had superior knowledge compared to Roundup users and consumers, including regarding the carcinogenic properties of the product, yet failed to accompany its sales and/or marketing of Roundup with any warnings or precautions for that grave danger. On information and belief, Wilbur-Ellis was one of the distributors providing Roundup and other glyphosate-containing products actually used by the Plaintiffs.

CLAIM ONE

STRICT LIABILITY - DESIGN DEFECT

(Against All Defendants)

COMPLAINT FOR DAMAGES AND DEMAND FOR JURY TRIAL

77. Plaintiffs incorporate by reference each and every allegation set forth in the preceding paragraphs as if fully stated herein.

78. Plaintiffs brings this strict liability claim against Defendants for defective design.

79. At all times relevant to this litigation, Defendants engaged in the business of testing, developing, designing, manufacturing, marketing, selling, distributing, and promoting Roundup® products, which are defective and unreasonably dangerous to consumers, including Plaintiffs, thereby placing Roundup® products into the stream of commerce. These actions were under the ultimate control and supervision of Defendants. At all times relevant to this litigation, Defendants designed, researched, developed, manufactured, produced, tested, assembled, labeled, advertised, promoted, marketed, sold, and distributed the Roundup® products that Plaintiffs were exposed to, as described above.

80. At all times relevant to this litigation, Defendants' Roundup® products were manufactured, designed, and labeled in an unsafe, defective, and inherently dangerous manner that was dangerous for use by or exposure to the public, and, in particular, the Plaintiffs.

81. At all times relevant to this litigation, Defendants' Roundup® products reached the intended consumers, handlers, and users or other persons coming into contact with these products in California and throughout the United States, including Plaintiffs, without substantial change in their condition as designed, manufactured, sold, distributed, labeled, and marketed by Defendants.

82. Defendants' Roundup® products, as researched, tested, developed, designed, licensed, manufactured, packaged, labeled, distributed, sold, and marketed by Defendants were defective in design and formulation in that when they left the hands of the Defendants' manufacturers and/or suppliers, they were unreasonably dangerous and dangerous to an extent beyond that which an ordinary consumer would contemplate.

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- 83. Defendants' Roundup® products, as researched, tested, developed, designed, licensed, manufactured, packaged, labeled, distributed, sold, and marketed by Defendants were defective in design and formulation in that when they left the hands of Defendants' manufacturers and/or suppliers, the foreseeable risks exceeded the alleged benefits associated with their design and formulation.
- 84. At all times relevant to this action, Defendants knew or had reason to know that i Roundup® products were defective and were inherently dangerous and unsafe when used in the manner instructed and provided by Defendants.
- 85. Therefore, at all times relevant to this litigation, Defendants' Roundup® products, as researched, tested, developed, designed, licensed, manufactured, packaged, labeled, distributed, sold and marketed by Defendants were defective in design and formulation, in one or more of the following ways:

- a. When placed in the stream of commerce, Defendants' Roundup® products were defective in design and formulation, and, consequently, dangerous to an extent beyond that which an ordinary consumer would contemplate.
- b. When placed in the stream of commerce, Defendants' Roundup® products were unreasonably dangerous in that they were hazardous and posed a grave risk of cancer and other serious illnesses when used in a reasonably anticipated manner.
- c. When placed in the stream of commerce, Defendants' Roundup® products contained unreasonably dangerous design defects and were not reasonably safe when used in a reasonably anticipated or intended manner.
- d. Defendants did not sufficiently test, investigate, or study its Roundup® products and, specifically, the active ingredient glyphosate.
- e. Exposure to Roundup® and glyphosate-containing products presents a risk of harmful side effects that outweigh any potential utility stemming from the use of the herbicide.
- f. Defendants knew or should have known at the time of marketing its Roundup® products that exposure to Roundup® and specifically, its active ingredient glyphosate, could result in cancer and other severe illnesses and injuries.
- g. Defendants did not conduct adequate post-marketing surveillance of its Roundup® products.
 - h. Defendants could have employed safer alternative designs and formulations.

86. Plaintiffs were exposed to Defendants' Roundup® products without knowledge of its dangerous characteristics.

87. At all times relevant to this litigation, Plaintiffs were exposed to the use of Defendants' Roundup® products in an intended or reasonably foreseeable manner without knowledge of their dangerous characteristics.

88. Plaintiffs could not have reasonably discovered the defects and risks associated with Roundup® or glyphosate-containing products before or at the time of exposure.

89. The harm caused by Defendants' Roundup® products far outweighed their benefit, rendering Defendants' products dangerous to an extent beyond that which an ordinary consumer would contemplate. Defendants' Roundup® products were and are more dangerous than alternative products and Defendants could have designed its Roundup® products to make them less dangerous. Indeed, at the time that Defendants designed its Roundup® products, the state of the industry's scientific knowledge was such that a less risky design or formulation was attainable.

90. At the time Roundup® products left Defendants' control, there was a practical, technically feasible and safer alternative design that would have prevented the harm without substantially impairing the reasonably anticipated or intended function of Defendants' herbicides.

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91. Defendants' defective design of its Roundup® products was willful, wanton, fraudulent, malicious, and conducted with reckless disregard for the health and safety of users of the Roundup® products, including the Plaintiffs herein.

- 92. Therefore, as a result of the unreasonably dangerous condition of its Roundup® products, Defendants are strictly liable to Plaintiffs.
- 93. The defects in Defendants' Roundup® products were substantial and contributing factors in causing Plaintiffs' grave injuries, and, but for Defendants' misconduct and omissions, Plaintiffs would not have sustained their injuries.
- 94. Defendants' conduct, as described above, was reckless. Defendants risked the lives of consumers and users of its products, including Plaintiffs, with knowledge of the safety problems associated with Roundup® and glyphosate-containing products, and suppressed this knowledge from the general public. Defendants made conscious decisions not to redesign, warn or inform the unsuspecting public. Defendants' reckless conduct warrants an award of punitive damages.
- 95. As a direct and proximate result of Defendants placing defective Roundup® products into the stream of commerce, Plaintiffs have suffered and continue to suffer grave injuries, and has endured physical pain and discomfort, as well as economic hardship, including considerable financial expenses for medical care and treatment. Plaintiffs will continue to incur these expenses in the future.

96. WHEREFORE, Plaintiffs respectfully request that this Court enter judgment in Plaintiffs' favor for compensatory and punitive damages, together with interest, costs herein incurred, attorneys' fees and all such other and further relief as this Court deems just and proper. Plaintiffs also demands a jury trial on the issues contained herein.

CLAIM TWO

STRICT LIABILITY - FAILURE TO WARN

(Against All Defendants)

- 97. Plaintiffs incorporate by reference each and every allegation set forth in the preceding paragraphs as if fully stated herein.
- 98. Plaintiffs bring this strict liability claim against Defendants for failure to warn.
- 99. At all times relevant to this litigation, Defendants engaged in the business of testing, developing, designing, manufacturing, marketing, selling, distributing, promoting and applying Roundup® products, which are defective and unreasonably dangerous to consumers, including Plaintiffs, because they do not contain adequate warnings or instructions concerning the dangerous characteristics of Roundup® and specifically, the active ingredient glyphosate. These actions were under the ultimate control and supervision of Defendants.

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100. Monsanto researched, developed, designed, tested, manufactured, inspected, labeled, distributed, marketed, promoted, sold, and otherwise released into the stream of commerce its Roundup® products, and in the course of same, directly advertised or marketed the products to consumers and end users, including the Plaintiffs, and persons responsible for consumers (such as employers), and therefore had a duty to warn of the risks associated with the use of Roundup® and glyphosate-containing products.

101. At all times relevant to this litigation, Defendants had a duty to properly test, develop, design, manufacture, inspect, package, label, market, promote, sell, distribute, maintain supply, provide proper warnings, and take such steps as necessary to ensure that Roundup® products did not cause users and consumers to suffer from unreasonable and dangerous risks. Defendants had a continuing duty to warn the Plaintiffs of the dangers associated with Roundup® use and exposure. Defendants, as manufacturer, seller, or distributor of chemical herbicides, are held to the knowledge of an expert in the field.

102. At the time of manufacture, Defendants could have provided the warnings or instructions regarding the full and complete risks of Roundup® and glyphosate-containing products because they knew or should have known of the unreasonable risks of harm associated with the use of and/or exposure to such products.

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At all times relevant to this litigation, Defendants failed to investigate, study, test, or promote the safety or to minimize the dangers to users and consumers of this product and to those who would foreseeably use or be harmed by Roundup, including Plaintiffs.

104. Despite the fact that Defendants knew or should have known that Roundup® posed a grave risk of harm, they failed to exercise reasonable care to warn of the dangerous risks associated with use and exposure. The dangerous propensities of its products and the carcinogenic characteristics of glyphosate, as described above, were known to Defendants, or scientifically knowable to Defendants through appropriate research and testing by known methods, at the time it distributed, supplied or sold the product, and not known to end users and consumers, such as Plaintiffs.

105. Defendants knew or should have known that these products created significant risks of serious bodily harm to consumers, as alleged herein, and Defendants failed to adequately warn consumers and reasonably foreseeable users of the risks of exposure to its products. Defendants have wrongfully concealed information concerning the dangerous nature of Roundup® and its active ingredient glyphosate, and further made false and/or misleading statements concerning the safety of Roundup® and glyphosate.

At all times relevant to this litigation, Defendants' Roundup® products reached the intended consumers, handlers, and users or other persons coming into contact with these products in California and throughout the United States, including Plaintiffs, without substantial

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change in their condition as designed, manufactured, sold, distributed, labeled, marketed and sprayed/applied by Defendants.

107. Plaintiffs were exposed to Roundup® products, as described above, without knowledge of their dangerous characteristics.

108. At all times relevant to this litigation, Plaintiffs were exposed to the use of Defendants' Roundup® products in their intended or reasonably foreseeable manner without knowledge of their dangerous characteristics.

109. Plaintiffs could not have reasonably discovered the defects and risks associated with Roundup® or glyphosate-containing products prior to or at the time of Plaintiffs' exposure. Plaintiffs relied upon the skill, superior knowledge, and judgment of Defendants.

110. Defendants knew or should have known that the minimal warnings disseminated with or accompanying the application of Roundup® products were inadequate, but they failed to communicate adequate information on the dangers and safe use/exposure and failed to communicate warnings and instructions that were appropriate and adequate to render the products safe for their ordinary, intended and reasonably foreseeable uses, including agricultural and horticultural applications.

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111. The information that Defendants did provide or communicate failed to contain relevant warnings, hazards, and precautions that would have enabled those exposed such as Plaintiffs to utilize the products safely and with adequate protection. Instead, Defendants disseminated information that was inaccurate, false, and misleading and which failed to communicate accurately or adequately the comparative severity, duration, and extent of the risk of injuries with use of and/or exposure to Roundup® and glyphosate; continued to aggressively promote the efficacy of its products, even after it knew or should have known of the unreasonable risks from use or exposure; and concealed, downplayed, or otherwise suppressed, through aggressive marketing and promotion, any information or research about the risks and dangers of exposure to Roundup® and glyphosate.

112. To this day, Defendants have failed to adequately and accurately warn of the true risks of Plaintiffs' injuries associated with the use of and exposure to Roundup® and its active ingredient glyphosate, a probable carcinogen.

113. As a result of their inadequate warnings, Roundup® products were defective and unreasonably dangerous when they left the possession and/or control of Defendants, were sold or distributed by Defendants, were applied by Defendants, and when Plaintiffs became exposed.

114. Defendants are liable to Plaintiffs for injuries caused by negligent or willful failure, as described above, to provide adequate warnings or other clinically relevant information and data

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regarding the appropriate use of their products and the risks associated with the use of or exposure to Roundup® and glyphosate.

115. The defects in these Roundup® products were substantial and contributing factors in causing Plaintiffs' injuries, and, but for Defendants' misconduct and omissions, Plaintiffs would not have sustained their injuries.

116. Had Defendants provided adequate warnings and instructions and properly disclosed and disseminated the risks associated with Roundup® products and application, Plaintiffs could have avoided the risk of developing injuries as alleged herein and the company who employed Plaintiffs could have obtained alternative herbicides.

117. As a direct and proximate result of Defendants placing defective Roundup® products into the stream of commerce and exposing Plaintiffs to them, Plaintiffs have suffered and continue to suffer severe injuries, and has endured physical pain and discomfort, as well as economic hardship, including considerable financial expenses for medical care and treatment. Plaintiffs will continue to incur these expenses in the future.

118. WHEREFORE, Plaintiffs respectfully request that this Court enter judgment in Plaintiffs' favor for compensatory and punitive damages, together with interest, costs herein incurred, attorneys' fees and all such other and further relief as this Court deems just and proper. Plaintiffs also demand a jury trial on the issues contained herein.

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CLAIM THREE

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NEGLIGENCE

(Against All Defendants)

119. Plaintiffs incorporate by reference each and every allegation set forth in the preceding paragraphs as if fully stated herein.

120. Defendants, directly or indirectly, caused Roundup® products to be sold, distributed, packaged, labeled, marketed, promoted, and/or used by Plaintiffs.

121. At all times relevant to this litigation, Defendants had a duty to exercise reasonable care in the design, research, manufacture, marketing, advertisement, supply, promotion, packaging, sale, and distribution of Roundup® products, including the duty to take all reasonable steps necessary to manufacture, promote, and/or sell a product that was not unreasonably dangerous to consumers and users of the product.

At all times relevant to this litigation, Defendants had a duty to exercise reasonable care in the marketing, advertisement, and sale of the Roundup® products. Defendants' duty of care owed to consumers and the general public included providing accurate, true, and correct information concerning the risks of using Roundup® and appropriate, complete, and accurate warnings concerning the potential adverse effects of exposure to Roundup®, and, in particular, its active ingredient glyphosate.

COMPLAINT FOR DAMAGES AND DEMAND FOR JURY TRIAL

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123. At all times relevant to this litigation, Defendants knew or, in the exercise of reasonable care, should have known of the hazards and dangers of Roundup® and specifically, the carcinogenic properties of the chemical glyphosate.

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124. Accordingly, at all times relevant to this litigation, Defendants knew or, in the exercise of reasonable care, should have known that use of or exposure to Roundup® products could cause or be associated with Plaintiffs' injuries and thus created a dangerous and unreasonable risk of injury to the users of these products, including Plaintiffs.

Defendants also knew or, in the exercise of reasonable care, should have known that 125. users and consumers of Roundup® were unaware of the risks and the magnitude of the risks associated with use of and/or exposure to Roundup® and glyphosate-containing products.

126. As such, Defendants breached their duty of reasonable care and failed to exercise ordinary care in the design, research, development, manufacture, testing, marketing, supply, promotion, advertisement, packaging, sale, and distribution of its Roundup® products, in that Defendants manufactured and produced defective herbicides containing the chemical glyphosate, knew or had reason to know of the defects inherent in its products, knew or had reason to know that a user's or consumer's exposure to the products created a significant risk of harm and unreasonably dangerous side effects, and failed to prevent or adequately warn of these risks and injuries.

127. Despite ability and means to investigate, study, and test products and to provide adequate warnings, Defendants have failed to do so. Indeed, Defendants wrongfully concealed information and has further made false and/or misleading statements concerning the safety and/or exposure to Roundup® and glyphosate.

128. Defendants' negligence included, but are not limited to:

- a. Manufacturing, producing, promoting, formulating, creating, developing, designing, selling, and/or distributing Roundup® products without thorough and adequate pre- and post-market testing;
- b. Manufacturing, producing, promoting, formulating, creating, developing, designing, selling, and/or distributing Roundup® while negligently and/or intentionally concealing and failing to disclose the results of trials, tests, and studies of exposure to glyphosate, and, consequently, the risk of serious harm associated with human use of and exposure to Roundup®;
- c. Failing to undertake sufficient studies and conduct necessary tests to determine whether or not Roundup® products and glyphosate-containing products were safe for their intended use in agriculture and horticulture;
- d. Failing to use reasonable and prudent care in the design, research, manufacture, and development of Roundup® products so as to avoid the risk of serious harm associated with the prevalent use of Roundup®/glyphosate as an herbicide;
- e. Failing to design and manufacture Roundup® products so as to ensure they were at least as safe and effective as other herbicides on the market;

f. Failing to provide adequate instructions, guidelines, and safety precautions to those persons who Defendants could reasonably foresee would use and be exposed to its Roundup® products;

- g. Failing to disclose to Plaintiffs, users/consumers, and the general public that use of and exposure to Roundup® presented severe risks of cancer and other grave illnesses;
- h. Failing to warn Plaintiffs, consumers, and the general public that the product's risk of harm was unreasonable and that there were safer and effective alternative herbicides available to Plaintiffs and other consumers;
- i. Systematically suppressing or downplaying contrary evidence about the risks, incidence, and prevalence of the side effects of Roundup® and glyphosate-containing products;
- j. Representing that its Roundup® products were safe for their intended use when, in fact,

 Defendants knew or should have known that the products were not safe for their intended

 purpose;
- k. Declining to make or propose any changes to Roundup® products' labeling or other promotional materials that would alert the consumers and the general public of the risks of Roundup® and glyphosate;
- 1. Advertising, marketing, and recommending the use of the Roundup® products, while concealing and failing to disclose or warn of the dangers known by Defendants to be associated with or caused by the use of or exposure to Roundup® and glyphosate;
- m. Continuing to disseminate information to its consumers, which indicate or imply that Defendants' Roundup® products are not unsafe for use in the agricultural and horticultural industries; and

n. Continuing the manufacture and sale of its products with the knowledge that the products were unreasonably unsafe and dangerous.

- 129. Defendants knew and/or should have known that it was foreseeable that consumers such as Plaintiffs would suffer injuries as a result of Defendants' failure to exercise ordinary care in the manufacturing, marketing, labeling, distribution, and sale of Roundup®.
- 130. Plaintiffs did not know the nature and extent of the injuries that could result from the intended use of and/or exposure to Roundup® or its active ingredient glyphosate.
- 131. Defendants' negligence was the proximate cause of the injuries, harm, and economic losses that Plaintiffs suffered, and will continue to suffer, as described herein.
- 132. Defendants' conduct, as described above, was reckless. Defendants regularly risk the lives of consumers and users of their products, including Plaintiffs, with full knowledge of the dangers of its products. Defendants made conscious decisions not to redesign, re-label, warn, or inform the unsuspecting public, including Plaintiffs. Defendants' reckless conduct therefore warrants an award of punitive damages.
- 133. As a proximate result of Defendants' wrongful acts and omissions in placing defective Roundup® products into the stream of commerce without adequate warnings of the hazardous and carcinogenic nature of glyphosate, Plaintiffs have suffered and continues to suffer severe and permanent physical and emotional injuries. Plaintiffs have endured pain and suffering, have COMPLAINT FOR DAMAGES AND DEMAND FOR JURY TRIAL

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suffered economic losses (including significant expenses for medical care and treatment) and will continue to incur these expenses in the future.

134. WHEREFORE, Plaintiffs respectfully requests that this Court enter judgment in Plaintiffs' favor for compensatory and punitive damages, together with interest, costs herein incurred, attorneys' fees and all such other and further relief as this Court deems just and proper. Plaintiffs also demands a jury trial on the issues contained herein.

CLAIM FOUR

BREACH OF IMPLIED WARRANTIES

(Against All Defendants)

- 135. Plaintiffs incorporate by reference each and every allegation set forth in the preceding paragraphs as if fully stated herein.
- At all times relevant to this litigation, Defendants engaged in the business of testing, developing, designing, manufacturing, marketing, selling, distributing, and promoting Roundup® products, which are defective and unreasonably dangerous to consumers, including Plaintiffs, thereby placing Roundup® products into the stream of commerce. These actions were under the ultimate control and supervision of Defendants.
- 137. Before the time that Plaintiffs were exposed to the use of the aforementioned Roundup® products, Defendants impliedly warranted to consumers and those exposed—including

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manufactured and sold by Defendants.

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COMPLAINT FOR DAMAGES AND DEMAND FOR JURY TRIAL

users, including Plaintiffs, without substantial change in the condition in which they were

The Roundup® products were expected to reach and did in fact reach consumers and/or

148. The harm caused by Roundup® products far outweighed their benefit, rendering the products more dangerous than an ordinary consumer or user would expect and more dangerous than alternative products.

149. As a direct and proximate result of Defendants' wrongful acts and omissions Plaintiffs have suffered severe and permanent physical and emotional injuries, including but not limited to their diagnoses of non-Hodgkin lymphoma. Plaintiffs have endured pain and suffering, have suffered economic loss (including significant expenses for medical care and treatment) and will continue to incur these expenses in the future.

150. WHEREFORE, Plaintiffs respectfully requests that this Court enter judgment in Plaintiffs' favor for compensatory and punitive damages, together with interest, costs herein incurred, attorneys' fees, and all such other and further relief as this Court deems just and proper. Plaintiffs also demand a jury trial on the issues contained herein.

COUNT FIVE

PUNITIVE DAMAGES

- 151. Plaintiffs repeat and reiterate the allegations previously set forth herein.
- 152. At all times material hereto, the Defendants knew or should have known that the subject product was inherently dangerous with respect to its health risks

153. At all times material hereto, the Defendants attempted to misrepresent and did misrepresent facts concerning the safety of the subject product.

- 154. Defendants' misrepresentations included knowingly withholding material information from the public, including the Plaintiffs herein, concerning the safety of the subject product.
- 155. At all times material hereto, the Defendants knew and recklessly disregarded the fact that human exposure to Roundup can and does cause health hazard, including non Hodgkin lymphoma.
- 156. Notwithstanding the foregoing, the Defendants continued to aggressively market and apply the subject product without disclosing the aforesaid risks.
- 157. Defendants knew of the subject product's defective and unreasonably dangerous nature, as set forth herein, but continued to design, develop, manufacture, market, distribute, sell, and apply it so as to maximize sales and profits at the expense of the health and safety of the public, including the Plaintiffs herein, in conscious and/or negligent disregard of the foreseeable harm caused by Roundup.
- 158. The Defendants intentionally concealed and/or recklessly failed to disclose to the public, including the Plaintiffs herein, the potentially life threatening hazards of Roundup in order to ensure continued and increased sales.

159. The Defendants' intentional and/or reckless failure to disclose information deprived the Plaintiffs of necessary information to enable Plaintiffs to weigh the true risks of using or being exposed to the subject product against its benefits.

160. As a direct and proximate result of the Defendants' conscious and deliberate disregard for the rights and safety of consumers such as the Plaintiffs, Plaintiffs suffered severe and permanent physical injuries. The Plaintiffs have endured substantial pain and suffering and has undergone extensive medical and surgical procedures. Plaintiffs have incurred significant expenses for medical care and treatment, and will continue to incur such expenses in the future. The Plaintiffs have lost past earnings and have suffered a loss of earning capacity. The Plaintiffs have suffered and will continue to suffer economic loss, and has otherwise been physically, emotionally and economically injured. The Plaintiffs' injuries and damages are permanent and will continue into the future.

161. The aforesaid conduct of the Defendants was committed with knowing, conscious, and deliberate disregard for the rights and safety of consumers, including the Plaintiffs herein, thereby entitling the Plaintiffs to punitive damages in an amount appropriate to punish the Defendants and deter them from similar conduct in the future.

162. WHEREFORE, Plaintiffs demand judgment against Defendants for compensatory, treble, and punitive damages, together with interest, costs of suit, attorneys' fees, and all such other relief as the Court deems proper.

COUNT SIX

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LOSS OF CONSORTIUM

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(Against All Defendants)

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163. Plaintiffs repeat and reiterate the allegations previously set forth herein.

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164. Plaintiffs ALVA PILLIOD and ALBERTA PILLIOD each allege causes of action for Loss of Consortium herein.

165. Plaintiffs ALVA PILLIOD and ALBERTA PILLIOD are currently lawfully married spouses and were married to one another at the time of their respective injuries. Plaintiff ALVA PILLIOD was entitled to Alberta Pilliod's comfort, care, affection, companionship, services, society, advice, guidance, counsel, and consortium and was deprived of such due to Defendants' conduct. Likewise, Plaintiff ALBERTA PILLIOD was entitled to Alva Pilliod's comfort, care, affection, companionship, services, society, advice, guidance, counsel, and consortium and was deprived of such due to Defendants' conduct.

166. Plaintiffs ALVA PILLIOD and ALBERTA PILLIOD each demand judgment against all Defendants for compensatory and punitive damages in excess of the jurisdictional minimum of this Court, together with interest, costs of suit, attorneys' fees, and all such other relief, as this Court deems proper.

COMPLAINT FOR DAMAGES AND DEMAND FOR JURY TRIAL

1 PRAYER FOR RELIEF 2 WHEREFORE, Plaintiffs request that the Court enter judgment in their favor and against 3 Defendants, awarding as follows: 4 1. Judgment for Plaintiffs and against Defendants; 5 2. For compensatory damages in an amount to be proven at trial; 6 7 3. For mental and physical suffering, according to proof; 8 4. For lost wages according to proof; 9 10 5. For disgorgement of profits, according to proof; 11 6. For punitive damages; 12 7. For default judgment as a sanction for the bad faith destruction of evidence, if 13 14 any, and according to proof, if any; 15 8. For costs including reasonable attorneys' fees, court costs, and other litigation 16 expenses; and 17 18 9. For any other relief the Court may deem just and proper. 19 **DEMAND FOR JURY TRIAL** 20 Plaintiffs hereby demand a jury trial on all claims so triable in this action. 21 Dated: August 30, 2018 22 Respectfully submitted, 23 24 THE MILLER FIRM, LLC 25 By: /s/ Curtis G. Hoke 26 Curtis G. Hoke (SBN 282465) 27 THE MILLER FIRM LLC 28 COMPLAINT FOR DAMAGES AND DEMAND FOR JURY TRIAL

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28	COMPLAINT FOR DAMAGES AND DEMAND FOR JURY TRIAL 49		

EXHIBIT 3

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1	Curtis G. Hoke (SBN 282465)				
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6	Attorneys for Plaintiffs				
7					
8	SUPERIOR COURT OF THE STATE OF CALIFORNIA				
9	IN AND FOR THE COUNTY OF ALAMEDA				
10					
11	(UNLIMITED JURISDICTION)				
12					
13	ALVA AND ALBERTA PILLIOD	JCCP No.: 4953			
14	Plaintiffs,	Case No.: RG17862702			
15	vs.				
16		FIRST AMENDED COMPLAINT FOR DAMAGES AND DEMAND FOR JURY			
17	MONSANTO COMPANY; WILBUR-ELLIS COMPANY, LLC; and	TRIAL			
18	WILBUR-ELLIS FEED, LLC,	BY FAX			
19 20	Defendants.	1. Strict Liability – Design Defect			
20		2. Strict Liability – Failure to Warn3. Negligence			
22		4. Breach of Implied Warranty5. Punitive Damage			
23		6. Loss of Consortium			
24		JURY TRIAL DEMANDED			
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28	COMPLAINT FOR DAMAGES AND DEMAND FOR JURY TRIAL				
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COMPLAINT AND DEMAND FOR JURY TRIAL

The above-captioned Plaintiffs (hereinafter, "Plaintiffs"), by attorneys, THE MILLER FIRM, LLC, as and for the Complaint herein allege upon information and belief the following:

SUMMARY OF THE CASE

- 1. In 1970, defendant Monsanto Company, Inc. discovered the herbicidal properties of glyphosate and began marketing it in products in 1974 under the brand name Roundup®. Roundup® is a non-selective herbicide used to kill weeds that commonly compete with the growing of crops. By 2001, glyphosate had become the most-used active ingredient in American agriculture with 85–90 millions of pounds used annually. That number grew to 185 million pounds by 2007. As of 2013, glyphosate was the world's most widely used herbicide.
- 2. Monsanto is a multinational agricultural biotechnology corporation based in St. Louis, Missouri. It is the world's leading producer of glyphosate. As of 2009, Monsanto was the world's leading producer of seeds, accounting for 27% of the world seed market. The majority of these seeds are of the Roundup Ready® brand. The stated advantage of Roundup Ready® crops is that they substantially improve a farmer's ability to control weeds, since glyphosate can be sprayed in the fields during the growing season without harming their crops. In 2010, an estimated 70% of corn and cotton, and 90% of soybean fields in the United States were Roundup Ready®.
- 3. Monsanto's glyphosate products are registered in 130 countries and approved for use on over 100 different crops. They are ubiquitous in the environment. Numerous studies confirm that glyphosate is found in rivers, streams, and groundwater in agricultural areas where Roundup® is COMPLAINT FOR DAMAGES AND DEMAND FOR JURY TRIAL

used. It has been found in food, in the urine of agricultural workers, and even in the urine of urban dwellers who are not in direct contact with glyphosate.

- 4. On March 20, 2015, the International Agency for Research on Cancer ("IARC"), an agency of the World Health Organization ("WHO"), issued an evaluation of several herbicides, including glyphosate. That evaluation was based, in part, on studies of exposures to glyphosate in several countries around the world, and it traces the health implications from exposure to glyphosate since 2001.
- 5. On July 29, 2015, IARC issued the formal monograph relating to glyphosate. In that monograph, the IARC Working Group provides a thorough review of the numerous studies and data relating to glyphosate exposure in humans.
- 6. The IARC Working Group classified glyphosate as a Group 2A herbicide, which means that it is probably carcinogenic to humans. The IARC Working Group concluded that the cancers most associated with glyphosate exposure are non-Hodgkin lymphoma and other haematopoietic cancers, including lymphocytic lymphoma/chronic lymphocytic leukemia, B-cell lymphoma, and multiple myeloma.
- 7. The IARC evaluation is significant. It confirms what has been believed for years: that glyphosate is toxic to humans.
- 8. Nevertheless, Monsanto, since it began selling Roundup®, has represented it as safe to COMPLAINT FOR DAMAGES AND DEMAND FOR JURY TRIAL

humans and the environment. Indeed, Monsanto has repeatedly proclaimed and continues to proclaim to the world, and particularly to United States consumers, that glyphosate-based herbicides, including Roundup®, create no unreasonable risks to human health or to the environment.

JURISDICTION AND VENUE

- 9. The California Superior Court has jurisdiction over this action pursuant to California Constitution Article VI, Section 10, which grants the Superior Court "original jurisdiction in all causes except those given by statute to other trial courts." The Statutes under which this action is brought do not specify any other basis for jurisdiction.
- 10. The California Superior Court has jurisdiction over the Defendants because, based on information and belief, each is a California resident, a corporation and/or entity organized under the laws of the State of California, a foreign corporation or association authorized to do business in California and registered with the California Secretary of State or has sufficient minimum contacts in California, or otherwise intentionally avails itself of the California market so as to render the exercise of jurisdiction over it by the California courts consistent with traditional notions of fair play and substantial justice.
- 11. Venue is proper in this Court pursuant to California Code of Civil Procedure Section 395 in that the subject injury occurred in Alameda County.

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12. Furthermore the Defendants have purposefully availed themselves of the benefits and the protections of the laws within the State of California. Monsanto has had sufficient contact such that the exercise of jurisdiction would be consistent with the traditional notions of fair play and substantial justice.

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13. Plaintiffs seek relief that is within the jurisdictional limits of this Court.

THE PARTIES

Plaintiffs

14. Plaintiff ALVA PILLIOD is a competent individual over the age of 18, a resident and citizen of the United States, and hereby submits to the jurisdiction of the Court and alleges that jurisdiction and venue in this court is proper. Mr. Pilliod currently resides in Livermore, County of Alameda, California. Mr. Pilliod used Monsanto's glyphosate-containing Roundup® product and suffered from severe physical, economy, and emotional injuries as a result of his use of Roundup® between 1975 and 2011, including but not limited to non-Hodgkin lymphoma, diagnosed in 2011 in California. Plaintiff ALBERTA PILLIOD is a competent individual over the age of 18, a resident and citizen of the United States, and hereby submits to the jurisdiction of the Court and alleges that jurisdiction and venue in this court is proper. Mrs. Pilliod currently resides in Livermore, County of Alameda, California. Mrs. Pilliod used the same Monsanto manufactured glyphosate-containing Roundup® products as Mr. Pilliod and suffered from severe physical, economic, and emotional injuries as a result of her use of Roundup® between 1975 and 2011, including but not limited to non-Hodgkin lymphoma, diagnosed in April, 2015 in

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California. Mr. and Mrs. Pilliod are lawfully married spouses and each seek damages for loss of consortium herein.

Defendants

- 15. Defendant Monsanto Company ("Monsanto") is a Delaware corporation with its headquarters and principal place of business in St. Louis, Missouri. At all times relevant to this complaint, Monsanto was the entity that discovered the herbicidal properties of glyphosate and the manufacturer of Roundup®. Monsanto has regularly transacted and conducted business within the state of California, and has derived substantial revenue from goods and products, including Roundup, used in the State of California. Monsanto expected or should have expected their acts to have consequences within the State of California, and derived substantial revenue from interstate commerce.
- 16. Defendants Wilbur-Ellis Company LLC is a California limited liability corporation with its headquarters and principal place of business in San Francisco, California. At all times relevant to this complaint, Wilbur-Ellis Company, LLC sold and distributed Monsanto products including Roundup, within the State of California.
- 17. Defendants Wilbur-Ellis Feed LLC (with Wilbur-Ellis Company LLC, hereinafter "Wilbur-Ellis") is a California limited liability corporation with its headquarters and principal place of business in San Francisco, California. At all times relevant to this complaint, Wilbur-Ellis Feed, LLC sold and distributed Monsanto products including Roundup, within the State of California.

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22. Treated plants generally die within two to three days. Because plants absorb glyphosate, it cannot be completely removed by washing or peeling produce or by milling, baking, or brewing grains.

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23. For nearly 40 years, farms across the world have used Roundup® without knowing of the dangers its use poses. That is because when Monsanto first introduced Roundup®, it touted glyphosate as a technological breakthrough: it could kill almost every weed without causing harm either to people or to the environment. Of course, history has shown that not to be true. According to the WHO, the main chemical ingredient of Roundup®—glyphosate—is a probable cause of cancer. Those most at risk are farm workers and other individuals with workplace exposure to Roundup®, such as workers in garden centers, nurseries, and landscapers. Agricultural workers are, once again, victims of corporate greed.

24. Monsanto assured the public that Roundup® was harmless. In order to prove this, Monsanto championed falsified data and attacked legitimate studies that revealed its dangers. Monsanto led a prolonged campaign of misinformation to convince government agencies, farmers and the general population that Roundup® was safe.

The Discovery of Glyphosate and Development of Roundup®

25. The herbicidal properties of glyphosate were discovered in 1970 by Monsanto chemist John Franz. The first glyphosate-based herbicide was introduced to the market in the mid-1970s under the brand name Roundup®. From the outset, Monsanto marketed Roundup® as a "safe" general-purpose herbicide for widespread commercial and consumer use. It still markets Roundup® as safe today.

Registration of Herbicides under Federal Law

26. The manufacture, formulation and distribution of herbicides, such as Roundup®, are regulated under the Federal Insecticide, Fungicide, and Rodenticide Act ("FIFRA" or "Act"), 7 U.S.C. § 136 et seq. FIFRA requires that all pesticides be registered with the Environmental COMPLAINT FOR DAMAGES AND DEMAND FOR JURY TRIAL

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Protection Agency ("EPA" or "Agency") prior to their distribution, sale, or use, except as described by the Act. 7 U.S.C. § 136a(a)

27. Because pesticides are toxic to plants, animals, and humans, at least to some degree, the EPA requires as part of the registration process, among other things, a variety of tests to evaluate the potential for exposure to pesticides, toxicity to people and other potential non-target organisms, and other adverse effects on the environment. Registration by the EPA, however, is not an assurance or finding of safety. The determination the Agency must make in registering or re-registering a product is not that the product is "safe," but rather that use of the product in accordance with its label directions "will not generally cause unreasonable adverse effects on the environment." 7 U.S.C. § 136a(c)(5)(D).

28. FIFRA defines "unreasonable adverse effects on the environment" to mean "any unreasonable risk to man or the environment, taking into account the economic, social, and environmental costs and benefits of the use of any pesticide." 7 U.S.C. § 136(bb). FIFRA thus requires EPA to make a risk/benefit analysis in determining whether a registration should be granted or allowed to continue to be sold in commerce.

29. The EPA and the State of California registered Roundup® for distribution, sale, and manufacture in the United States and the State of California.

30. FIFRA generally requires that the registrant, Monsanto in the case of Roundup®, conducts the health and safety testing of pesticide products. The EPA has protocols governing the conduct of tests required for registration and the laboratory practices that must be followed in conducting these tests. The data produced by the registrant must be submitted to the EPA for review and evaluation. The government is not required, nor is it able, however, to perform the product tests that are required of the manufacturer.

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31. The evaluation of each pesticide product distributed, sold, or manufactured is completed at the time the product is initially registered. The data necessary for registration of a pesticide has changed over time. The EPA is now in the process of re-evaluating all pesticide products through a Congressionally-mandated process called "re-registration." 7 U.S.C. § 136a-1. In order to reevaluate these pesticides, the EPA is demanding the completion of additional tests and the submission of data for the EPA's review and evaluation.

32. In the case of glyphosate, and therefore Roundup®, the EPA had planned on releasing its preliminary risk assessment —in relation to the reregistration process—no later than July 2015. The EPA completed its review of glyphosate in early 2015, but it delayed releasing the risk assessment pending further review in light of the WHO's health-related findings.

Scientific Fraud Underlying the Marketing and Sale of Glyphosate/Roundup

33. Based on early studies that glyphosate could cause cancer in laboratory animals, the EPA originally classified glyphosate as *possibly carcinogenic to humans* (Group C) in 1985. After

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pressure from Monsanto, including contrary studies it provided to the EPA, the EPA changed its classification to *evidence of non-carcinogenicity in humans* (Group E) in 1991. In so classifying glyphosate, however, the EPA made clear that the designation did not mean the chemical does not cause cancer: "It should be emphasized, however, that designation of an agent in Group E is based on the available evidence at the time of evaluation and should not be interpreted as a definitive conclusion that the agent will not be a carcinogen under any circumstances."

34. On two occasions, the EPA found that the laboratories hired by Monsanto to test the toxicity of its Roundup® products for registration purposes committed fraud.

- 35. In the first instance, Monsanto, in seeking initial registration of Roundup® by EPA, hired Industrial Bio-Test Laboratories ("IBT") to perform and evaluate pesticide toxicology studies relating to Roundup®. IBT performed about 30 tests on glyphosate and glyphosate-containing products, including nine of the 15 residue studies needed to register Roundup®.
- 36. In 1976, the United States Food and Drug Administration ("FDA") performed an inspection of Industrial Bio-Test Industries ("IBT") that revealed discrepancies between the raw data and the final report relating to the toxicological impacts of glyphosate. The EPA subsequently audited IBT; it too found the toxicology studies conducted for the Roundup® herbicide to be invalid. An EPA reviewer stated, after finding "routine falsification of data" at IBT, that it was "hard to believe the scientific integrity of the studies when they said they took specimens of the uterus from male rabbits."

37. Three top executives of IBT were convicted of fraud in 1983.

38. In the second incident of data falsification, Monsanto hired Craven Laboratories in 1991 to perform pesticide and herbicide studies, including for Roundup®. In that same year, the owner of Craven Laboratories and three of its employees were indicted, and later convicted, of fraudulent laboratory practices in the testing of pesticides and herbicides.

39. Despite the falsity of the tests that underlie its registration, within a few years of its launch, Monsanto was marketing Roundup® in 115 countries.

The Importance of Roundup® to Monsanto's Market Dominance Profits

- 40. The success of Roundup® was key to Monsanto's continued reputation and dominance in the marketplace. Largely due to the success of Roundup® sales, Monsanto's agriculture division was out-performing its chemicals division's operating income, and that gap increased yearly. But with its patent for glyphosate expiring in the United States in the year 2000, Monsanto needed a strategy to maintain its Roundup® market dominance and to ward off impending competition.
- A1. In response, Monsanto began the development and sale of genetically engineered Roundup Ready® seeds in 1996. Since Roundup Ready® crops are resistant to glyphosate; farmers can spray Roundup® onto their fields during the growing season without harming the crop. This allowed Monsanto to expand its market for Roundup® even further; by 2000, Monsanto's biotechnology seeds were planted on more than 80 million acres worldwide and

nearly 70% of American soybeans were planted from Roundup Ready® seeds. It also secured Monsanto's dominant share of the glyphosate/Roundup® market through a marketing strategy that coupled proprietary Roundup Ready® seeds with continued sales of its Roundup® herbicide.

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42. Through a three-pronged strategy of increased production, decreased prices and by coupling with Roundup Ready® seeds, Roundup® became Monsanto's most profitable product. In 2000, Roundup® accounted for almost \$2.8 billion in sales, outselling other herbicides by a margin of five to one, and accounting for close to half of Monsanto's revenue. Today, glyphosate remains one of the world's largest herbicides by sales volume.

Monsanto has known for decades that it falsely advertises the safety of Roundup®

- 43. In 1996, the New York Attorney General ("NYAG") filed a lawsuit against Monsanto based on its false and misleading advertising of Roundup ® products. Specifically, the lawsuit challenged Monsanto's general representations that its spray-on glyphosate-based herbicides, including Roundup®, were "safer than table salt" and "practically non-toxic" to mammals, birds, and fish. Among the representations the NYAG found deceptive and misleading about the human and environmental safety of Roundup® are the following:
- a) Remember that environmentally friendly Roundup herbicide is biodegradable. It won't build up in the soil so you can use Roundup with confidence along customers' driveways, sidewalks and fences ...

- b) And remember that Roundup is biodegradable and won't build up in the soil. That will give you the environmental confidence you need to use Roundup everywhere you've got a weed, brush, edging or trimming problem.
 - c) Roundup biodegrades into naturally occurring elements.
- d) Remember that versatile Roundup herbicide stays where you put it. That means there's no washing or leaching to harm customers' shrubs or other desirable vegetation.
- e) This non-residual herbicide will not wash or leach in the soil. It ... stays where you apply it.
- f) You can apply Accord with "confidence because it will stay where you put it" it bonds tightly to soil particles, preventing leaching. Then, soon after application, soil microorganisms biodegrade Accord into natural products.
 - g) Glyphosate is less toxic to rats than table salt following acute oral ingestion.
- h) Glyphosate's safety margin is much greater than required. It has over a 1,000-fold safety margin in food and over a 700-fold safety margin for workers who manufacture it or use it.
- i) You can feel good about using herbicides by Monsanto. They carry a toxicity category rating of 'practically non-toxic' as it pertains to mammals, birds and fish.
- j) "Roundup can be used where kids and pets will play and breaks down into natural material." This ad depicts a person with his head in the ground and a pet dog standing in an area which has been treated with Roundup.

1	44. On November 19, 1996, Monsanto entered into an Assurance of Discontinuance with
2	NYAG, in which Monsanto agreed, among other things, "to cease and desist from publishing or
3	broadcasting any advertisements [in New York] that represent, directly or by implication" that:
4	a) its glyphosate-containing pesticide products or any component thereof are safe,
5	
6	non-toxic, harmless or free from risk.
7	* * *
8	b) its glyphosate-containing pesticide products or any component thereof
9	manufactured, formulated, distributed or sold by Monsanto are biodegradable
10	* * *
11	
12	c) its glyphosate-containing pesticide products or any component thereof stay
13	where they are applied under all circumstances and will not move through the environment by
14	any means.
15	* * *
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17	d) its glyphosate-containing pesticide products or any component thereof are
18	"good" for the environment or are "known for their environmental characteristics."
19	* * *
20	e) glyphosate-containing pesticide products or any component thereof are safer or
21	
22	less toxic than common consumer products other than herbicides;
23	f) its glyphosate-containing products or any component thereof might be
24	classified as "practically non-toxic.
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26	45. Monsanto did not alter its advertising in the same manner in any state other than New
27	Vork and on information and belief still has not done so today
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e products or any component thereof onsanto are biodegradable e products or any component thereof stay will not move through the environment by e products or any component thereof are environmental characteristics." roducts or any component thereof are safer or an herbicides; or any component thereof might be same manner in any state other than New and on information and belief still has not done so today. COMPLAINT FOR DAMAGES AND DEMAND FOR JURY TRIAL

46. In 2009, France's highest court ruled that Monsanto had not told the truth about the safety of Roundup®. The French court affirmed an earlier judgement that Monsanto had falsely advertised its herbicide Roundup® as "biodegradable" and that it "left the soil clean."

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Classifications and Assessments of Glyphosate

- 47. The IARC process for the classification of glyphosate followed the stringent procedures for the evaluation of a chemical agent. Over time, the IARC Monograph program has reviewed 980 agents. Of those reviewed, it has determined 116 agents to be Group 1 (Known Human Carcinogens); 73 agents to be Group 2A (Probable Human Carcinogens); 287 agents to be Group 2B (Possible Human Carcinogens); 503 agents to be Group 3 (Not Classified); and one agent to be Probably Not Carcinogenic.
- 48. The established procedure for IARC Monograph evaluations is described in the IARC Programme's Preamble. Evaluations are performed by panels of international experts, selected on the basis of their expertise and the absence of actual or apparent conflicts of interest.
- 49. One year before the Monograph meeting, the meeting is announced and there is a call both for data and for experts. Eight months before the Monograph meeting, the Working Group membership is selected and the sections of the Monograph are developed by the Working Group members. One month prior to the Monograph meeting, the call for data is closed and the various

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draft sections are distributed among Working Group members for review and comment. Finally, at the Monograph meeting, the Working Group finalizes review of all literature, evaluates the evidence in each category, and completes the overall evaluation. Within two weeks after the Monograph meeting, the summary of the Working Group findings are published in Lancet Oncology, and within a year after the meeting, the final Monograph is finalized and published.

50. In assessing an agent, the IARC Working Group reviews the following information: (a) human, experimental, and mechanistic data; (b) all pertinent epidemiological studies and cancer bioassays; and (c) representative mechanistic data. The studies must be publicly available and have sufficient detail for meaningful review, and reviewers cannot be associated with the underlying study.

- 51. In March 2015, IARC reassessed glyphosate. The summary published in *The Lancet Oncology* reported that glyphosate is a Group 2A agent and probably carcinogenic in humans.
- 52. On July 29, 2015, IARC issued its Monograph for glyphosate, Monograph 112. For Volume 112, the volume that assessed glyphosate, a Working Group of 17 experts from 11 countries met at IARC from March 3–10, 2015, to assess the carcinogenicity of certain herbicides, including glyphosate. The March meeting culminated nearly a one-year review and preparation by the IARC Secretariat and the Working Group, including a comprehensive review of the latest available scientific evidence. According to published procedures, the Working Group considered "reports that have been published or accepted for publication in the openly COMPLAINT FOR DAMAGES AND DEMAND FOR JURY TRIAL

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- 57. The IARC Working Group found an increased risk between exposure to glyphosate and non-Hodgkin lymphoma ("NHL") and several subtypes of NHL, and the increased risk persisted after adjustment for other pesticides.
- 58. The IARC Working Group also found that glyphosate caused DNA and chromosomal damage in human cells. One study in community residents reported increases in blood markers of chromosomal damage (micronuclei) after glyphosate formulations were sprayed.
- 59. In male CD-1 mice, glyphosate induced a positive trend in the incidence of a rare tumor, renal tubule carcinoma. A second study reported a positive trend for haemangiosarcoma in male mice. Glyphosate increased pancreatic islet-cell adenoma in male rats in two studies. A glyphosate formulation promoted skin tumors in an initiation-promotion study in mice.
- 60. The IARC Working Group also noted that glyphosate has been detected in the urine of agricultural workers, indicating absorption. Soil microbes degrade glyphosate to aminomethylphosphoric acid (AMPA). Blood AMPA detection after exposure suggests intestinal microbial metabolism in humans.
- 61. The IARC Working Group further found that glyphosate and glyphosate formulations induced DNA and chromosomal damage in mammals, and in human and animal cells in utero.

62. The IARC Working Group also noted genotoxic, hormonal, and enzymatic effects in mammals exposed to glyphosate. Essentially, glyphosate inhibits the biosynthesis of aromatic amino acids, which leads to several metabolic disturbances, including the inhibition of protein and secondary product biosynthesis and general metabolic disruption.

63. The IARC Working Group also reviewed an Agricultural Health Study, consisting of a prospective cohort of 57,311 licensed pesticide applicators in Iowa and North Carolina. While this study differed from others in that it was based on a self-administered questionnaire, the results support an association between glyphosate exposure and Multiple Myeloma, Hairy Cell Leukemia (HCL), and Chronic Lymphocytic Leukemia (CLL), in addition to several other cancers.

Other Earlier Findings About Glyphosate's Dangers to Human Health

64. The EPA has a technical fact sheet, as part of its Drinking Water and Health, National Primary Drinking Water Regulations publication, relating to glyphosate. This technical fact sheet predates the IARC March 20, 2015, evaluation. The fact sheet describes the release patterns for glyphosate as follows:

Release Patterns

65. Glyphosate is released to the environment in its use as a herbicide for controlling woody and herbaceous weeds on forestry, right-of-way, cropped and non-cropped sites. These sites may be around water and in wetlands.

- 66. It may also be released to the environment during its manufacture, formulation, transport, storage, disposal and cleanup, and from spills. Since glyphosate is not a listed chemical in the Toxics Release Inventory, data on releases during its manufacture and handling are not available.
- 67. Occupational workers and home gardeners may be exposed to glyphosate by inhalation and dermal contact during spraying, mixing, and cleanup. They may also be exposed by touching soil and plants to which glyphosate was applied. Occupational exposure may also occur during glyphosate's manufacture, transport storage, and disposal.
- 68. In 1995, the Northwest Coalition for Alternatives to Pesticides reported that in California, the state with the most comprehensive program for reporting of pesticide-caused illness, glyphosate was the third most commonly-reported cause of pesticide illness among agricultural workers.

Recent Worldwide Bans on Roundup®/Glyphosate

69. Several countries around the world have instituted bans on the sale of Roundup® and other glyphosate-containing herbicides, both before and since IARC first announced its assessment for glyphosate in March 2015, and more countries undoubtedly will follow suit in light of the as the dangers of the use of Roundup® are more widely known. The Netherlands issued a ban on all glyphosate-based herbicides in April 2014, including Roundup®, which takes effect by the end of 2015. In issuing the ban, the Dutch Parliament member who introduced the successful legislation stated: "Agricultural pesticides in user-friendly packaging are sold in abundance to private persons. In garden centers, Roundup® is promoted as harmless, but COMPLAINT FOR DAMAGES AND DEMAND FOR JURY TRIAL

75. On information and belief, Wilbur-Ellis was, at all relevant times, engaged in the distribution of Roundup, Roundup-ready crops and other glyphosate-containing products from Monsanto to retailers and commercial/agricultural users in California.

76. Wilbur-Ellis had superior knowledge compared to Roundup users and consumers, including regarding the carcinogenic properties of the product, yet failed to accompany its sales and/or marketing of Roundup with any warnings or precautions for that grave danger. On information and belief, Wilbur-Ellis was one of the distributors providing Roundup and other glyphosate-containing products actually used by the Plaintiffs.

CLAIM ONE

STRICT LIABILITY - DESIGN DEFECT

(Against All Defendants)

- 77. Plaintiffs incorporate by reference each and every allegation set forth in the preceding paragraphs as if fully stated herein.
- 78. Plaintiffs brings this strict liability claim against Defendants for defective design.
- 79. At all times relevant to this litigation, Defendants engaged in the business of testing, developing, designing, manufacturing, marketing, selling, distributing, and promoting Roundup® products, which are defective and unreasonably dangerous to consumers, including Plaintiffs, thereby placing Roundup® products into the stream of commerce. These actions were

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under the ultimate control and supervision of Defendants. At all times relevant to this litigation, Defendants designed, researched, developed, manufactured, produced, tested, assembled, labeled, advertised, promoted, marketed, sold, and distributed the Roundup® products that Plaintiffs were exposed to, as described above.

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- 80. At all times relevant to this litigation, Defendants' Roundup® products were manufactured, designed, and labeled in an unsafe, defective, and inherently dangerous manner that was dangerous for use by or exposure to the public, and, in particular, the Plaintiffs.
- 81. At all times relevant to this litigation, Defendants' Roundup® products reached the intended consumers, handlers, and users or other persons coming into contact with these products in California and throughout the United States, including Plaintiffs, without substantial change in their condition as designed, manufactured, sold, distributed, labeled, and marketed by Defendants.
- 82. Defendants' Roundup® products, as researched, tested, developed, designed, licensed, manufactured, packaged, labeled, distributed, sold, and marketed by Defendants were defective in design and formulation in that when they left the hands of the Defendants' manufacturers and/or suppliers, they were unreasonably dangerous and dangerous to an extent beyond that which an ordinary consumer would contemplate.
- 83. Defendants' Roundup® products, as researched, tested, developed, designed, licensed, manufactured, packaged, labeled, distributed, sold, and marketed by Defendants were defective

in design and formulation in that when they left the hands of Defendants' manufacturers and/or suppliers, the foreseeable risks exceeded the alleged benefits associated with their design and formulation.

- 84. At all times relevant to this action, Defendants knew or had reason to know that i Roundup® products were defective and were inherently dangerous and unsafe when used in the manner instructed and provided by Defendants.
- 85. Therefore, at all times relevant to this litigation, Defendants' Roundup® products, as researched, tested, developed, designed, licensed, manufactured, packaged, labeled, distributed, sold and marketed by Defendants were defective in design and formulation, in one or more of the following ways:
- a. When placed in the stream of commerce, Defendants' Roundup® products were defective in design and formulation, and, consequently, dangerous to an extent beyond that which an ordinary consumer would contemplate.
- b. When placed in the stream of commerce, Defendants' Roundup® products were unreasonably dangerous in that they were hazardous and posed a grave risk of cancer and other serious illnesses when used in a reasonably anticipated manner.
- c. When placed in the stream of commerce, Defendants' Roundup® products contained unreasonably dangerous design defects and were not reasonably safe when used in a reasonably anticipated or intended manner.
- d. Defendants did not sufficiently test, investigate, or study its Roundup® products and, specifically, the active ingredient glyphosate.

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would contemplate. Defendants' Roundup® products were and are more dangerous than alternative products and Defendants could have designed its Roundup® products to make them less dangerous. Indeed, at the time that Defendants designed its Roundup® products, the state of the industry's scientific knowledge was such that a less risky design or formulation was attainable.

90. At the time Roundup® products left Defendants' control, there was a practical, technically feasible and safer alternative design that would have prevented the harm without substantially impairing the reasonably anticipated or intended function of Defendants' herbicides.

- 91. Defendants' defective design of its Roundup® products was willful, wanton, fraudulent, malicious, and conducted with reckless disregard for the health and safety of users of the Roundup® products, including the Plaintiffs herein.
- 92. Therefore, as a result of the unreasonably dangerous condition of its Roundup® products, Defendants are strictly liable to Plaintiffs.
- 93. The defects in Defendants' Roundup® products were substantial and contributing factors in causing Plaintiffs' grave injuries, and, but for Defendants' misconduct and omissions, Plaintiffs would not have sustained their injuries.

94. Defendants' conduct, as described above, was reckless. Defendants risked the lives of consumers and users of its products, including Plaintiffs, with knowledge of the safety problems associated with Roundup® and glyphosate-containing products, and suppressed this knowledge from the general public. Defendants made conscious decisions not to redesign, warn or inform the unsuspecting public. Defendants' reckless conduct warrants an award of punitive damages.

95. As a direct and proximate result of Defendants placing defective Roundup® products into the stream of commerce, Plaintiffs have suffered and continue to suffer grave injuries, and has endured physical pain and discomfort, as well as economic hardship, including considerable financial expenses for medical care and treatment. Plaintiffs will continue to incur these expenses in the future.

96. WHEREFORE, Plaintiffs respectfully request that this Court enter judgment in Plaintiffs' favor for compensatory and punitive damages, together with interest, costs herein incurred, attorneys' fees and all such other and further relief as this Court deems just and proper. Plaintiffs also demands a jury trial on the issues contained herein.

CLAIM TWO

STRICT LIABILITY - FAILURE TO WARN

(Against All Defendants)

97. Plaintiffs incorporate by reference each and every allegation set forth in the preceding paragraphs as if fully stated herein.

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98. Plaintiffs bring this strict liability claim against Defendants for failure to warn.

99. At all times relevant to this litigation, Defendants engaged in the business of testing, developing, designing, manufacturing, marketing, selling, distributing, promoting and applying Roundup® products, which are defective and unreasonably dangerous to consumers, including Plaintiffs, because they do not contain adequate warnings or instructions concerning the dangerous characteristics of Roundup® and specifically, the active ingredient glyphosate. These actions were under the ultimate control and supervision of Defendants.

100. Monsanto researched, developed, designed, tested, manufactured, inspected, labeled, distributed, marketed, promoted, sold, and otherwise released into the stream of commerce its Roundup® products, and in the course of same, directly advertised or marketed the products to consumers and end users, including the Plaintiffs, and persons responsible for consumers (such as employers), and therefore had a duty to warn of the risks associated with the use of Roundup® and glyphosate-containing products.

101. At all times relevant to this litigation, Defendants had a duty to properly test, develop, design, manufacture, inspect, package, label, market, promote, sell, distribute, maintain supply, provide proper warnings, and take such steps as necessary to ensure that Roundup® products did not cause users and consumers to suffer from unreasonable and dangerous risks. Defendants had a continuing duty to warn the Plaintiffs of the dangers associated with Roundup® use and COMPLAINT FOR DAMAGES AND DEMAND FOR JURY TRIAL

exposure. Defendants, as manufacturer, seller, or distributor of chemical herbicides, are held to the knowledge of an expert in the field.

102. At the time of manufacture, Defendants could have provided the warnings or instructions regarding the full and complete risks of Roundup® and glyphosate-containing products because they knew or should have known of the unreasonable risks of harm associated with the use of and/or exposure to such products.

103. At all times relevant to this litigation, Defendants failed to investigate, study, test, or promote the safety or to minimize the dangers to users and consumers of this product and to those who would foreseeably use or be harmed by Roundup, including Plaintiffs.

104. Despite the fact that Defendants knew or should have known that Roundup® posed a grave risk of harm, they failed to exercise reasonable care to warn of the dangerous risks associated with use and exposure. The dangerous propensities of its products and the carcinogenic characteristics of glyphosate, as described above, were known to Defendants, or scientifically knowable to Defendants through appropriate research and testing by known methods, at the time it distributed, supplied or sold the product, and not known to end users and consumers, such as Plaintiffs.

105. Defendants knew or should have known that these products created significant risks of serious bodily harm to consumers, as alleged herein, and Defendants failed to adequately warn COMPLAINT FOR DAMAGES AND DEMAND FOR JURY TRIAL

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consumers and reasonably foreseeable users of the risks of exposure to its products. Defendants have wrongfully concealed information concerning the dangerous nature of Roundup® and its active ingredient glyphosate, and further made false and/or misleading statements concerning the safety of Roundup® and glyphosate.

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106. At all times relevant to this litigation, Defendants' Roundup® products reached the intended consumers, handlers, and users or other persons coming into contact with these products in California and throughout the United States, including Plaintiffs, without substantial change in their condition as designed, manufactured, sold, distributed, labeled, marketed and sprayed/applied by Defendants.

107. Plaintiffs were exposed to Roundup® products, as described above, without knowledge of their dangerous characteristics.

108. At all times relevant to this litigation, Plaintiffs were exposed to the use of Defendants' Roundup® products in their intended or reasonably foreseeable manner without knowledge of their dangerous characteristics.

109. Plaintiffs could not have reasonably discovered the defects and risks associated with Roundup® or glyphosate-containing products prior to or at the time of Plaintiffs' exposure. Plaintiffs relied upon the skill, superior knowledge, and judgment of Defendants.

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Defendants knew or should have known that the minimal warnings disseminated with or accompanying the application of Roundup® products were inadequate, but they failed to communicate adequate information on the dangers and safe use/exposure and failed to communicate warnings and instructions that were appropriate and adequate to render the products safe for their ordinary, intended and reasonably foreseeable uses, including agricultural and horticultural applications.

The information that Defendants did provide or communicate failed to contain relevant 111. warnings, hazards, and precautions that would have enabled those exposed such as Plaintiffs to utilize the products safely and with adequate protection. Instead, Defendants disseminated information that was inaccurate, false, and misleading and which failed to communicate accurately or adequately the comparative severity, duration, and extent of the risk of injuries with use of and/or exposure to Roundup® and glyphosate; continued to aggressively promote the efficacy of its products, even after it knew or should have known of the unreasonable risks from use or exposure; and concealed, downplayed, or otherwise suppressed, through aggressive marketing and promotion, any information or research about the risks and dangers of exposure to Roundup® and glyphosate.

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glyphosate, a probable carcinogen.

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Plaintiffs' injuries associated with the use of and exposure to Roundup® and its active ingredient

To this day, Defendants have failed to adequately and accurately warn of the true risks of

113. As a result of their inadequate warnings, Roundup® products were defective and unreasonably dangerous when they left the possession and/or control of Defendants, were sold or distributed by Defendants, were applied by Defendants, and when Plaintiffs became exposed.

114. Defendants are liable to Plaintiffs for injuries caused by negligent or willful failure, as described above, to provide adequate warnings or other clinically relevant information and data regarding the appropriate use of their products and the risks associated with the use of or exposure to Roundup® and glyphosate.

115. The defects in these Roundup® products were substantial and contributing factors in causing Plaintiffs' injuries, and, but for Defendants' misconduct and omissions, Plaintiffs would not have sustained their injuries.

116. Had Defendants provided adequate warnings and instructions and properly disclosed and disseminated the risks associated with Roundup® products and application, Plaintiffs could have avoided the risk of developing injuries as alleged herein and the company who employed Plaintiffs could have obtained alternative herbicides.

117. As a direct and proximate result of Defendants placing defective Roundup® products into the stream of commerce and exposing Plaintiffs to them, Plaintiffs have suffered and continue to suffer severe injuries, and has endured physical pain and discomfort, as well as economic

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hardship, including considerable financial expenses for medical care and treatment. Plaintiffs will continue to incur these expenses in the future.

WHEREFORE, Plaintiffs respectfully request that this Court enter judgment in Plaintiffs' 118. favor for compensatory and punitive damages, together with interest, costs herein incurred, attorneys' fees and all such other and further relief as this Court deems just and proper. Plaintiffs also demand a jury trial on the issues contained herein.

CLAIM THREE

NEGLIGENCE

(Against All Defendants)

- 119. Plaintiffs incorporate by reference each and every allegation set forth in the preceding paragraphs as if fully stated herein.
- 120. Defendants, directly or indirectly, caused Roundup® products to be sold, distributed, packaged, labeled, marketed, promoted, and/or used by Plaintiffs.
- At all times relevant to this litigation, Defendants had a duty to exercise reasonable care 121. in the design, research, manufacture, marketing, advertisement, supply, promotion, packaging, sale, and distribution of Roundup® products, including the duty to take all reasonable steps necessary to manufacture, promote, and/or sell a product that was not unreasonably dangerous to consumers and users of the product.

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122. At all times relevant to this litigation, Defendants had a duty to exercise reasonable care in the marketing, advertisement, and sale of the Roundup® products. Defendants' duty of care owed to consumers and the general public included providing accurate, true, and correct information concerning the risks of using Roundup® and appropriate, complete, and accurate warnings concerning the potential adverse effects of exposure to Roundup®, and, in particular, its active ingredient glyphosate.

123. At all times relevant to this litigation, Defendants knew or, in the exercise of reasonable care, should have known of the hazards and dangers of Roundup® and specifically, the carcinogenic properties of the chemical glyphosate.

124. Accordingly, at all times relevant to this litigation, Defendants knew or, in the exercise of reasonable care, should have known that use of or exposure to Roundup® products could cause or be associated with Plaintiffs' injuries and thus created a dangerous and unreasonable risk of injury to the users of these products, including Plaintiffs.

125. Defendants also knew or, in the exercise of reasonable care, should have known that users and consumers of Roundup® were unaware of the risks and the magnitude of the risks associated with use of and/or exposure to Roundup® and glyphosate-containing products.

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126. As such, Defendants breached their duty of reasonable care and failed to exercise ordinary care in the design, research, development, manufacture, testing, marketing, supply, promotion, advertisement, packaging, sale, and distribution of its Roundup® products, in that Defendants manufactured and produced defective herbicides containing the chemical glyphosate, knew or had reason to know of the defects inherent in its products, knew or had reason to know that a user's or consumer's exposure to the products created a significant risk of harm and unreasonably dangerous side effects, and failed to prevent or adequately warn of these risks and injuries.

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127. Despite ability and means to investigate, study, and test products and to provide adequate warnings, Defendants have failed to do so. Indeed, Defendants wrongfully concealed information and has further made false and/or misleading statements concerning the safety and/or exposure to Roundup® and glyphosate.

- 128. Defendants' negligence included, but are not limited to:
- a. Manufacturing, producing, promoting, formulating, creating, developing, designing, selling, and/or distributing Roundup® products without thorough and adequate pre- and post-market testing;
- b. Manufacturing, producing, promoting, formulating, creating, developing, designing, selling, and/or distributing Roundup® while negligently and/or intentionally concealing and failing to disclose the results of trials, tests, and studies of exposure to glyphosate, and, consequently, the risk of serious harm associated with human use of and exposure to Roundup®;

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c. Failing to undertake sufficient studies and conduct necessary tests to determine
whether or not Roundup® products and glyphosate-containing products were safe for their
intended use in agriculture and horticulture;

- d. Failing to use reasonable and prudent care in the design, research, manufacture, and development of Roundup® products so as to avoid the risk of serious harm associated with the prevalent use of Roundup®/glyphosate as an herbicide;
- e. Failing to design and manufacture Roundup® products so as to ensure they were at least as safe and effective as other herbicides on the market;
- f. Failing to provide adequate instructions, guidelines, and safety precautions to those persons who Defendants could reasonably foresee would use and be exposed to its Roundup® products;
- g. Failing to disclose to Plaintiffs, users/consumers, and the general public that use of and exposure to Roundup® presented severe risks of cancer and other grave illnesses;
- h. Failing to warn Plaintiffs, consumers, and the general public that the product's risk of harm was unreasonable and that there were safer and effective alternative herbicides available to Plaintiffs and other consumers;
- i. Systematically suppressing or downplaying contrary evidence about the risks, incidence, and prevalence of the side effects of Roundup® and glyphosate-containing products;
- j. Representing that its Roundup® products were safe for their intended use when, in fact, Defendants knew or should have known that the products were not safe for their intended purpose;

k. Declining to make or propose any changes to Roundup® products' labeling or other promotional materials that would alert the consumers and the general public of the risks of Roundup® and glyphosate;

- 1. Advertising, marketing, and recommending the use of the Roundup® products, while concealing and failing to disclose or warn of the dangers known by Defendants to be associated with or caused by the use of or exposure to Roundup® and glyphosate;
- m. Continuing to disseminate information to its consumers, which indicate or imply that Defendants' Roundup® products are not unsafe for use in the agricultural and horticultural industries; and
- n. Continuing the manufacture and sale of its products with the knowledge that the products were unreasonably unsafe and dangerous.
- 129. Defendants knew and/or should have known that it was foreseeable that consumers such as Plaintiffs would suffer injuries as a result of Defendants' failure to exercise ordinary care in the manufacturing, marketing, labeling, distribution, and sale of Roundup®.
- 130. Plaintiffs did not know the nature and extent of the injuries that could result from the intended use of and/or exposure to Roundup® or its active ingredient glyphosate.
- 131. Defendants' negligence was the proximate cause of the injuries, harm, and economic losses that Plaintiffs suffered, and will continue to suffer, as described herein.

Defendants' conduct, as described above, was reckless. Defendants regularly risk the lives of consumers and users of their products, including Plaintiffs, with full knowledge of the dangers of its products. Defendants made conscious decisions not to redesign, re-label, warn, or inform the unsuspecting public, including Plaintiffs. Defendants' reckless conduct therefore warrants an award of punitive damages.

133.

As a proximate result of Defendants' wrongful acts and omissions in placing defective Roundup® products into the stream of commerce without adequate warnings of the hazardous and carcinogenic nature of glyphosate, Plaintiffs have suffered and continues to suffer severe and permanent physical and emotional injuries. Plaintiffs have endured pain and suffering, have suffered economic losses (including significant expenses for medical care and treatment) and will continue to incur these expenses in the future.

WHEREFORE, Plaintiffs respectfully requests that this Court enter judgment in 134. Plaintiffs' favor for compensatory and punitive damages, together with interest, costs herein incurred, attorneys' fees and all such other and further relief as this Court deems just and proper. Plaintiffs also demands a jury trial on the issues contained herein.

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CLAIM FOUR

BREACH OF IMPLIED WARRANTIES

(Against All Defendants)

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135. Plaintiffs incorporate by reference each and every allegation set forth in the preceding paragraphs as if fully stated herein.

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136. At all times relevant to this litigation, Defendants engaged in the business of testing, developing, designing, manufacturing, marketing, selling, distributing, and promoting Roundup® products, which are defective and unreasonably dangerous to consumers, including Plaintiffs, thereby placing Roundup® products into the stream of commerce. These actions were under the ultimate control and supervision of Defendants.

137. Before the time that Plaintiffs were exposed to the use of the aforementioned Roundup® products, Defendants impliedly warranted to consumers and those exposed—including Plaintiffs—that Roundup® products were of merchantable quality and safe and fit for the use for which they were intended; specifically, as horticultural herbicides.

138. Defendants, however, failed to disclose that Roundup® has dangerous propensities when used as intended and that the use of and/or exposure to Roundup® and glyphosate-containing products carries an increased risk of developing severe injuries, including Plaintiffs' injuries.

139. Plaintiffs reasonably relied upon the skill, superior knowledge and judgment of Defendants and upon their implied warranties that the Roundup® products were of merchantable quality and fit for their intended purpose or use.

140. Upon information and belief, Plaintiffs were at all relevant times in privity with Defendants.

141.

Defendants to the purchasers and/or users of their horticultural herbicides, and as such Plaintiffs are entitled to assert this claim.

Plaintiffs are the intended third-party beneficiaries of implied warranties made by

142. The Roundup® products were expected to reach and did in fact reach consumers and/or users, including Plaintiffs, without substantial change in the condition in which they were manufactured and sold by Defendants.

At all times relevant to this litigation, Defendants were aware that consumers and users of their products, including Plaintiffs, would use Roundup® products as marketed by Defendants, which is to say that Plaintiffs were foreseeable users of Roundup®.

144. Defendants intended that Roundup® products be used in the manner in which Plaintiffs were exposed to it and Defendants impliedly warranted each product to be of merchantable quality, safe, and fit for this use, despite the fact that Roundup® was not adequately tested and/or researched.

145. In reliance upon Defendants' implied warranty, Plaintiffs used or were exposed to Roundup® as instructed and labeled and in the foreseeable manner intended, recommended, promoted and marketed by Defendants.

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146. Plaintiffs could not have reasonably discovered or known of the risks of serious injury associated with Roundup® or glyphosate.

147. Defendants breached their implied warranty to Plaintiffs in that Roundup® products were not of merchantable quality, safe, or fit for their intended use, and/or adequately tested. Roundup® has dangerous propensities when used as intended and can cause serious injuries, including those injuries complained of herein.

148. The harm caused by Roundup® products far outweighed their benefit, rendering the products more dangerous than an ordinary consumer or user would expect and more dangerous than alternative products.

149. As a direct and proximate result of Defendants' wrongful acts and omissions Plaintiffs have suffered severe and permanent physical and emotional injuries, including but not limited to their diagnoses of non-Hodgkin lymphoma. Plaintiffs have endured pain and suffering, have suffered economic loss (including significant expenses for medical care and treatment) and will continue to incur these expenses in the future.

WHEREFORE, Plaintiffs respectfully requests that this Court enter judgment in 150. Plaintiffs' favor for compensatory and punitive damages, together with interest, costs herein incurred, attorneys' fees, and all such other and further relief as this Court deems just and proper. Plaintiffs also demand a jury trial on the issues contained herein.

COMPLAINT FOR DAMAGES AND DEMAND FOR JURY TRIAL

COUNT FIVE

PUNITIVE DAMAGES

- 151. Plaintiffs repeat and reiterate the allegations previously set forth herein.
- 152. At all times material hereto, the Defendants knew or should have known that the subject product was inherently dangerous with respect to its health risks
- 153. At all times material hereto, the Defendants attempted to misrepresent and did misrepresent facts concerning the safety of the subject product.
- 154. Defendants' misrepresentations included knowingly withholding material information from the public, including the Plaintiffs herein, concerning the safety of the subject product.
- 155. At all times material hereto, the Defendants knew and recklessly disregarded the fact that human exposure to Roundup can and does cause health hazard, including non Hodgkin lymphoma.
- 156. Notwithstanding the foregoing, the Defendants continued to aggressively market and apply the subject product without disclosing the aforesaid risks.
- 157. Defendants knew of the subject product's defective and unreasonably dangerous nature, as set forth herein, but continued to design, develop, manufacture, market, distribute, sell, and COMPLAINT FOR DAMAGES AND DEMAND FOR JURY TRIAL

apply it so as to maximize sales and profits at the expense of the health and safety of the public, including the Plaintiffs herein, in conscious and/or negligent disregard of the foreseeable harm caused by Roundup.

158. The Defendants intentionally concealed and/or recklessly failed to disclose to the public, including the Plaintiffs herein, the potentially life threatening hazards of Roundup in order to ensure continued and increased sales.

159. The Defendants' intentional and/or reckless failure to disclose information deprived the Plaintiffs of necessary information to enable Plaintiffs to weigh the true risks of using or being exposed to the subject product against its benefits.

160. As a direct and proximate result of the Defendants' conscious and deliberate disregard for the rights and safety of consumers such as the Plaintiffs, Plaintiffs suffered severe and permanent physical injuries. The Plaintiffs have endured substantial pain and suffering and has undergone extensive medical and surgical procedures. Plaintiffs have incurred significant expenses for medical care and treatment, and will continue to incur such expenses in the future. The Plaintiffs have lost past earnings and have suffered a loss of earning capacity. The Plaintiffs have suffered and will continue to suffer economic loss, and has otherwise been physically, emotionally and economically injured. The Plaintiffs' injuries and damages are permanent and will continue into the future.

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1	[DDODOGED] ODDED		
1	[PROPOSED] ORDER		
2	T1:	00 a.m., in Department 17 of th	
3	Court, located at 1221 Oak Street, Oakland, CA 94612. Upon reading and considering Plaintiffs' Motion		
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6	for Leave to File a First Amended Complaint and GOOD CAUSE APPEARING THEREFORE, IT IS		
7	HEREBY ORDERED THAT:		
8	1. Plaintiffs' Motion for Leave to File an Amended Complaint is hereby GRANTED.		
9	2. Plaintiffs' proposed First Amended Complaint (Hoke Decl., Exhibit 3) is deemed filed and		
10	served as of the date of this Order.		
11	3. Plaintiffs' First Amended Complaint shall be deemed to relate back to June 2, 2017, the date		
12	when Plaintiffs filed their original Complaint.		
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14	IT IS SO ORDERED		
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16	DATED:		
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18	The Honorable Ioan JUDGE OF THE SU		
19		TERIOR COURT	
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28	[PROPOSED] ORDER		
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FILED BY FAX ALAMEDA COUNTY 1 Curtis G. Hoke (State Bar No. 282465) August 30, 2018 THE MILLER FIRM, LLC CLERK OF 2 108 Railroad Ave. THE SUPERIOR COURT By Dajuana Turner, Deputy Orange, VA 22960 3 Telephone: (540) 672-4224 CASE NUMBER: RG17862702 Facsimile: (540) 672-3055 4 choke@millerfirmllc.com 5 Attorneys for Plaintiffs 6 7 SUPERIOR COURT OF THE STATE OF CALIFORNIA 8 FOR THE COUNTY OF ALAMEDA 9 10 COORDINATION PROCEEDING SPECIAL JCCP NO. 4953 11 **TITLE (RULE 3.550)** 12 ROUNDUP PRODUCTS CASES Case No.: RG17862702 13 14 THIS DOCUMENT RELATES TO: NOTICE OF MOTION AND 15 PLAINTIFFS' MOTION FOR LEAVE Pilliod, et al. v. Monsanto Company, et al. TO FILE A FIRST AMENDED Alameda Superior Court Case No.: 16 **COMPLAINT; MEMORANDUM OF** RG17862702 POINTS AND AUTHORITIES IN 17 SUPPORT OF MOTION 18 19 **BY FAX** 20 21 22 Hon. Judge Ioana Petrou 23 Hearing Date: October 9, 2018 Time: 9:00 a.m. 24 Department: 17 25 Reservation No.: R-1993571 26 27 28 NOTICE OF MOTION AND PLAINTIFFS' MOTION FOR LEAVE TO FILE AMENDED COMPLAINT;

NOTICE OF MOTION AND PLAINTIFFS' MOTION FOR LEAVE TO FILE AMENDED COMPLAINT; MEMORANDUM OF POINTS AND AUTHORITIES IN SUPPORT OF MOTION

TO ALL PARTIES AND THEIR COUNSELS OF RECORD:

PLEASE TAKE NOTICE that on October 9, 2018, at 9:00 a.m., or as soon thereafter as the matter can be heard in Department 17 of this Court, The Honorable Judge Ioana Petrou presiding, located at 1221 Oak Street, Oakland, CA 94612, Plaintiffs will move and hereby do respectfully move this Court for an Order permitting plaintiffs Alva Pilliod and Alberta Pilliod to file a First Amended Complaint pursuant to California Code of Civil Procedure Sections 452, 473 and 576.

This Motion is based primarily on newly-uncovered information that Mrs. Pilliod – in addition to Mr. Pilliod – has also been diagnosed with non-Hodgkin's lymphoma. Additionally, Plaintiffs' proposed First Amended Complaint would add individual Loss of Consortium Claims held by both Mr. Pilliod and Mrs. Pilliod and would remove unnecessary allegations (Complaint ¶15-19) relating to co-plaintiffs Charles Baker, John Novak, Sharon Rowland, Sharon McClurg, and Marjorie Grubka, who have since filed their own separate individualized Complaints. This motion is based on this Notice of Motion and Motion, the accompanying Memorandum of Points and Authorities, Declaration of Curtis G. Hoke, the records and file herein, and on such evidence and argument as may be presented at the hearing of this motion.

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DATED: August 30, 2018

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Respectfully submitted,

THE MILLER FIRM, LLC

By: /s/ Curtis Hoke

Curtis G. Hoke (SBN 282465)

THE MILLER FIRM, LLC

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Orange, VA 22960

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Attorneys for Plaintiffs

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NOTICE OF MOTION AND PLAINTIFFS' MOTION FOR LEAVE TO FILE AMENDED COMPLAINT; MEMORANDUM OF POINTS AND AUTHORITIES IN SUPPORT OF MOTION

MEMORANDUM OF POINTS AND AUTHORITIES

I. INTRODUCTION

Plaintiffs Alva Pilliod and Alberta Pilliod (respectively referred to as "Mr. Pilliod" and "Mrs. Pilliod"), both of which are named plaintiffs in *Pilliod, et al. v. Monsanto Company, et al.*, commenced this action on June 2, 2017 seeking damages relating to Mr. Pilliod's diagnosis of non-Hodgkin's lymphoma. Subsequently, on or about July 26, 2018, The Miller Firm learned that Mrs. Pilliod had also been diagnosed with non-Hodgkin's lymphoma. Plaintiffs therefore now respectfully move this Court for an order granting leave for them to file a First Amended Complaint (*see*, Hoke Decl., **Exhibit 3**) alleging that Mrs. Pilliod has also been diagnosed with non-Hodgkin's lymphoma and suffered injuries therefrom as a result of Defendants' conduct. Plaintiffs' proposed First Amended complaint (*see*, Hoke Decl., **Exhibit 3**) would also add a Loss of Consortium claim held by each spouse and remove now-unnecessary allegations (Complaint ¶15-19) relating to co-plaintiffs Charles Baker, John Novak, Sharon Rowland, Sharon McClurg, and Marjorie Grubka, who have since filed their own separate individual Complaints.

II. ARGUMENT

A. Motions To Amend Should Be Liberally Granted.

A court may, in the furtherance of justice, allow a party to amend any pleading on any terms as may be proper. *See*, Cal. Code Civ. Proc. § 452, § 473(a), and §576. These statutory provisions give courts the power to "permit amendments in furtherance of justice has received a very liberal interpretation by the courts of this state." *See*, *Klopstock v. Superior Court in & for City & Cty. of San Francisco*, 17 Cal. 2d 13, 19 (1941); *Nestle v. City of Santa Monica*, 6 Cal. 3d

920, 939 (1972) [the general rule of liberal construction of pleadings (Code Civ. Proc., § 452) and of liberal allowance of amendments should prevail].

"[T]hat the trial courts are to liberally permit such amendments, at any stage of the proceeding, has been established policy of this state since 1901." See, Hirsa v. Superior Court, 118 Cal. App. 3d 486, 488–89 (Cal. Ct. App. 1981). In Arthur L. Sachs, Inc. v. City of Oceanside, 151 Cal. App. 3d 315, 319 (Cal. Ct. App. 1984), the California Court of Appeal determined on the eve or trial, that it was error to deny the amendment of a cross-complaint to add an additional theory of recovery where the delay in seeking the amendment was attributable to the opposing party's failure to comply with discovery requests.

California's policy liberally favouring leave to amend is so strong that it is an abuse of discretion to deny an amendment unless the adverse party can show meaningful prejudice, such as the running of the statute of limitations, trial delay, the loss of critical evidence, or added preparation costs. *See*, *Atkinson v. Elk Corp.*, 109 Cal. App. 4th 739, 761 (2003); *Solit v. Tokai Bank, Ltd. New York Branch*, 68 Cal. App. 4th 1435, 1448 (1999). Absent a showing of such prejudice, delay alone is not grounds for denial of a motion to amend. *See*, *Kittredge Sports Co. v. Superior Court*, 213 Cal. App. 3d 1045, 1048 (Cal. Ct. App. 1989); *Higgins v. Del Faro*, 123 Cal. App. 3d 558, 563 (Cal. Ct. App. 1981).

A trial court may deny an otherwise proper amendment if there was an unwarranted delay in bringing the motion to amend. *See*, *Atkinson*, 109 Cal.App.4th at 761. However to deny leave to amend based on an unreasonable delay the opposing party must have been misled or prejudiced by the delay. *Kittredge*, 213 Cal.App.3d at 1048.

Here, on or about July 26, 2018, via a telephone conversation with Mrs. Alberta Pilliod, Plaintiffs' counsel discovered that Mrs. Pilliod – in addition to her husband, Alva Pilliod – was

also diagnosed with non-Hodgkin's lymphoma. Plaintiff's counsel had previously believed that only Mr. Pilliod was diagnosed with non-Hodgkin's lymphoma, and believed that Mrs. Pilliod was diagnosed with brain cancer, which Plaintiffs' counsel did not know to be non-Hodgkin's lymphoma. In August, 2018, The Miller Firm received medical records showing that Mrs. Pilliod's brain cancer is actually a form of non-Hodgkin's lymphoma called diffuse large B-cell CNS lymphoma. See, Decl. of Curtis G. Hoke at ¶ 2. On or about August 15, 2018, Plaintiffs' counsel met and conferred with Monsanto's counsel in order to determine whether Monsanto would stipulate to Plaintiffs' filing of their proposed First Amended Complaint. Monsanto objected to the proposed amendments, forcing Plaintiffs to file this Motion. See, Decl. of Curtis G. Hoke at ¶3.

Due to Mrs. Pilliod's diagnosis of non-Hodgkin's lymphoma, first known to The Miller Firm on or about July 26, 2018, Plaintiffs now seek leave to file their First Amended Complaint, appended as **Exhibit 3** to the Declaration of Curtis G. Hoke, filed concurrently herewith.

B. Defendants Will Not be Prejudiced By Plaintiffs' Proposed First Amended Complaint.

A defendant cannot make an adequate showing of "prejudice" simply because a proposed amended complaint expands the scope of litigation. Indeed, California law is clear that if a plaintiff wants to expand the scope of a complaint to add additional legitimate legal theories, it is proper to do so if discovery is still open. *See, Morgan v. Superior Court of Cal. In & For Los Angeles County*, 172 Cal. App. 2d 527, 530–31 (1959) (noting that there is no sound reason why a plaintiff having two causes of action for should be limited to proving but one of them). Therefore, Defendants cannot legitimately claim they would be "prejudiced" by Plaintiffs' proposed First Amended Complaint.

C. Plaintiffs' First Amended Complaint Should "Relate Back" To June 2, 2017, The Filing Date Of Plaintiffs' Original Complaint.

Even when a plaintiff seeks to add new legal theories or causes of action, an amended complaint relates back to the date of the filing of the original complaint so long as recovery sought in both pleadings is based upon the same general set of facts. *See, Smeltzley v. Nicholson Mfg. Co.*, 18 Cal. 3d 932, 938–39 (1977); *see, also, Kittredge*, 213 Cal.App.3d at 1048; *Hirsa*, 118 Cal.App.3d at 489.

In the California Supreme Court case *Grudt v. City of Los Angeles*, 2 Cal. 3d 575 (1970), that plaintiff sued the city under the *respondeat superior* theory alleging that police officers had intentionally killed her husband. The amended complaint charged that the city was liable for its own negligence in retaining officers known to be dangerous. Even though the plaintiff's amended complaint added a significant new dimension to the original, the court determined that the amendment was based "on the same general set of facts" as the original complaint" and allowed the amendment to be made. Furthermore, in *Smeltzley*, for example, the Court of Appeal found that the relation back precedent rests on the fundamental policy that cases should be decided on their merits. *Smeltzley*, 18 Cal.3d at 939.

Here, Mr. and Mrs. Pilliod filed their original Complaint on June 2, 2017. Mrs. Pilliod was already a named plaintiff therein and considered a party to the action. Plaintiffs now seek to clarify Mrs. Pilliod's claims due to her counsel's newly-learned information that she was also diagnosed with non-Hodgkin's lymphoma. Specifically, Plaintiffs' First Amended Complaint, among other things, alleges that Mrs. Pilliod used the same Monsanto manufactured glyphosate-containing Roundup® products as Mr. Pilliod from 1975 through 2011 (over 35 years) and suffered from severe physical, economic, and emotional injuries as

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a result of her use and exposure to Roundup®, including but not limited to non-Hodgkin's lymphoma. See, Decl. of Curtis G. Hoke, **Exhibit 3** (proposed First Amended Complaint) at ¶14. Further, Plaintiffs' First Amended Complaint retains, word-for-word, the exact same general set of factual allegations relating to Monsanto's conduct giving rise to both Mr. and Mrs. Pilliod's injuries – namely, inter alia, allegations that 1) Monsanto knew or should have known that glyphosate/Roundup is carcinogenic to humans, 2) Monsanto engaged in scientific fraud underlying the marketing and sale of glyphosate/Roundup, 3) that Roundup was important to Monsanto's market dominance profits, 4) that Monsanto has known for decades that it falsely advertised the safety of Roundup, and 5) Monsanto failed to adequately warn of the known or knowable risks of glyphosate/Roundup. Compare, proposed First Amended Complaint to Complaint at \P 38 – 51. Further, Counts I through V remain unchanged from Plaintiffs' original Complaint to the proposed First Amended Complaint. Compare, proposed First Amended Complaint to Complaint at \P 82 – 167. Indeed, Plaintiffs' motion only seeks to 1) clarify that Mrs. Pilliod was also diagnosed with non-Hodgkin's and that she used and was exposed to the same Roundup products that Mr. Pilliod was exposed to and suffered injuries therefrom due to Defendants' conduct, 2) add appropriate claims for Loss of Consortium held by each plaintiff, and 3) remove unnecessary allegations (Complaint ¶15-19) relating to co-plaintiffs originally filed in Plaintiffs' Complaint who have since filed their own separate individual Complaints.

Plaintiffs' Proposed First Amended Complaint Will Not Delay The Trial D. Because No Trial Date Has Been Set.

This honorable Court has not yet set a trial date for this action. See, Decl. of Curtis G. Hoke at ¶4. It is beyond cavil that Plaintiffs' proposed First Amended Complaint cannot delay a trial that has not yet been set.

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E. The Amended Complaint Will Not Require Additional Preparation Costs.

Here, Mrs. Alberta Pilliod is already a named plaintiff in Plaintiffs' original Complaint. She therefore is already subject to discovery requests, depositions, trial preparation and trial under either the original Complaint or Plaintiffs' proposed First Amended Complaint.

In order to further limit costs, Plaintiffs' counsel has already prepared a proposed First Amended Complaint containing the proposed amendments discussed herein and has appended it to the concurrently-filed Declaration of Curtis G. Hoke as **Exhibit 3**. Further, Plaintiffs concurrently-filed [Proposed] Order – if Plaintiffs' motion *sub judice* is granted – contemplates that Plaintiffs' proposed First Amended Complaint would be deemed filed and served on the date of the Order, therefore obviating the need for any expenditure of any additional time or resources.

III. CONCLUSION

For the foregoing reasons, Plaintiffs respectfully request that this honorable Court grant their Motion for Leave to File a First Amended Complaint.

DATED: August 30, 2018

Respectfully submitted,

THE MILLER FIRM, LLC

By: /s/ Curtis Hoke Curtis G. Hoke (SBN 282465)

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APPENDIX OF PROPOSED CHANGES

Pursuant to California Rules of Court, Rule 3.1324(a)(2)-(3), Plaintiffs propose the following amendments to the Complaint.

A. Add the following to Complaint ¶14:

"Plaintiff ALBERTA PILLIOD is a competent individual over the age of 18, a resident and citizen of the United States, and hereby submits to the jurisdiction of the Court and alleges that jurisdiction and venue in this court is proper. Mrs. Pilliod currently resides in Livermore, County of Alameda, California. Mrs. Pilliod used the same Monsanto manufactured glyphosate-containing Roundup® products as Mr. Pilliod and suffered from severe physical, economic, and emotional injuries as a result of her use of Roundup® between 1975 and 2011, including but not limited to non-Hodgkin lymphoma, diagnosed in April, 2015 in California. Mr. and Mrs. Pilliod are lawfully married spouses and each seek damages for loss of consortium herein."

- B. Add a cause of action for Loss of Consortium (Count VI) held by both Alva Pilliod and Alberta Pilliod. *See*, Decl. of Curtis G. Hoke, Exhibit 3 at ¶¶163-166.
- C. Remove unnecessary allegations (Complaint ¶15-19) relating to co-plaintiffs Charles Baker,

 John Novak, Sharon Rowland, Sharon McClurg, and Marjorie Grubka, who have since filed
 their own separate Complaints.

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