

1 Michael J. Miller (appearance *pro hac vice*)
2 Timothy Litzenburg (appearance *pro hac vice*)
3 Curtis G. Hoke (State Bar No. 282465)
4 **THE MILLER FIRM, LLC**
5 108 Railroad Ave.
6 Orange, VA 22960
7 Phone: (540) 672-4224
8 Fax: (540) 672-3055
9 mmiller@millerfirmllc.com
10 tlitzenburg@millerfirmllc.com
11 choke@millerfirmllc.com

12 *Attorneys for Plaintiff*
13 **DEWAYNE JOHNSON**

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SUPERIOR COURT OF THE STATE OF CALIFORNIA
FOR THE COUNTY OF SAN FRANCISCO

DEWAYNE JOHNSON,
Plaintiff,
v.
MONSANTO COMPANY
Defendants.

Case No. CGC-16-550128

**NOTICE OF MOTION AND PLAINTIFF'S
MOTION *IN LIMINE* NO. 12 TO EXCLUDE
ANY ARGUMENT AND TESTIMONY
THAT EPA REGISTRATION PRECLUDED
MONSANTO FROM WARNING OF THE
RISK OF NON-HODGKIN'S LYMPHOMA**

Trial Judge: TBD

Hearing Date: TBD

Time: TBD

Department: TBD

Trial Date: June 18, 2018

[Filed concurrently with Declaration of Curtis
Hoke and [Proposed] Order]

ELECTRONICALLY
FILED
Superior Court of California,
County of San Francisco
05/24/2018
Clerk of the Court
BY: SANDRA SCHIRO
Deputy Clerk

1 **TO ALL PARTIES AND THEIR ATTORNEYS OF RECORD:**

2 **PLEASE TAKE NOTICE** that, at a date and time set by the trial judge assigned to this matter
3 of the above-entitled Court located at 400 McAllister St. San Francisco, CA 94102-4515, Plaintiff
4 Dewayne Johnson will and hereby does move *in limine* to exclude any argument and testimony that
5 EPA registration precluded Monsanto from warning of the risk of Non-Hodgkin's lymphoma.

6 This motion *in limine* has been brought pursuant to Evid. Code §§ 210, 350, and 352 and is
7 based on the grounds that Monsanto should not be permitted to argue to the jury that it was precluded
8 from warning Plaintiff of the risks of its Glyphosate-based herbicides in light of this Court's Order
9 dismissing its affirmative defenses on federal preemption. Any argument, suggestion, or testimony
10 that Monsanto could not provide warnings beyond what was included in the EPA approved labeling is
11 an incorrect statement of the law and is entirely irrelevant to Plaintiff's warnings-based claims. Any
12 probative value of the evidence is substantially outweighed by the likelihood that the introduction of
13 such evidence would mislead and confuse the jury.

14 This motion *in limine* is based on this notice of motion, the motion and accompanying
15 memorandum of points and authorities, the Court's May 17, 2018 Order denying Monsanto's motion
16 for summary judgment and granting plaintiff's motion for summary adjudication, the concurrently filed
17 Declaration of Curtis Hoke, the concurrently filed proposed order, all pleadings and papers on file in
18 this matter, and such further oral and documentary evidence and papers as the Court may consider at
19 the time of the hearing.

20 Respectfully Submitted,

21 Dated: May 24, 2018

THE MILLER FIRM, LLC

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23 By: /s/ Curtis G. Hoke

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choke@millerfirmllc.com

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MEMORANDUM OF POINTS AND AUTHORITIES

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I. ARGUMENT

On May 17, 2018, this Court granted Plaintiff’s motions for summary adjudication of Monsanto’s affirmative defenses relating to federal preemption.¹ The Court found that Plaintiff’s failure-to-warn claims were not preempted because California product liability law “is equivalent to, and fully consistent with, FIFRA’s misbranding provisions.” *See*, Order Granting P’s Mot. for Sum. Adj. (“Preemption Order”) (May 17, 2018) at 39; *quoting Bates v. Dow Agrosiences LLC*, 554 U.S. 431, 444 (2005). In expressly rejecting Monsanto’s arguments in support of preemption, this Court concluded that:

Monsanto’s express preemption argument depends on the premise that Monsanto is immune from FIFRA liability so long as it uses a label that has been approved by the EPA or is otherwise consistent with the EPA’s factual findings. **That’s not true.**

Preemption Order at 40 (emphasis added); see also p. 42 (“EPA approval of a pesticide is *not* a defense for the commission of any offense under FIFRA...”)(emphasis in original). The Court again confirmed the legal principle that EPA registration “does not mean that the product complies with California law” when it refused Monsanto’s proposed jury instructions on the role of the EPA in regulating pesticides. See Order on Deposition Designations and Certain Proposed Jury Instructions (“Jury Inst. Order”) at 7.

Despite the Court’s previous Orders, Plaintiff anticipates that Monsanto will seek to introduce argument and testimony that: (1) Monsanto was precluded from warning consumers of risks not contained within the product labeling; and (2) Monsanto cannot be held liable for failing to warn of the risk of Non-Hodgkin’s lymphoma because the EPA approved labeling for glyphosate-based herbicides (GBHs) without requiring a cancer warning.² Plaintiff seeks to exclude these inaccurate, misleading, and prejudicial statements, including the following arguments and testimony:

¹ The Court also denied Monsanto’s Motion for Summary Judgment on the basis of preemption.

² Evidence that GBH’s are registered with the EPA amounts to prima facie evidence that the labeling complies with California law. See Jury Inst. Order at 7. EPA’s registration of GBH’s may be admissible on the question of whether Monsanto’s warnings were adequate, however, Monsanto is not entitled to mislead the jury into thinking that Monsanto was foreclosed from warning of additional risks associated with its product because of EPA registration.

- 1 • “Once the EPA has. . . approved an herbicide label, the manufacturer is required by FIFRA to
2 use only the labeling that has been approved by the EPA.” Monsanto’s Memorandum in
3 Support of Summary Judgment (“Def’s Mot. for Sum. J.”) at 11.
- 4 • “EPA’s labeling decisions have the force of federal law, and manufacturers that disregard those
5 federal regulatory decisions are subject to legal sanctions under FIFRA.” Def’s Mot. for Sum.
6 J. at 11.
- 7 • “Any label that deviates from EPA’s approved safety labeling may be deemed “misbranded”
8 by the Agency.” Exhibit A; Expert Report of John Fowle, III, Ph.D., DA BT (“Fowle Rep.”) at
9 49.
- 10 • “EPA takes label approval very seriously, and it does not allow companies the freedom to
11 choose to place a warning on the label that the product might cause cancer when EPA has
12 determined that it does not.” Fowle Rep. at 49.
- 13 • “EPA has the proper processes and procedures and requirements in place to make sure that those
14 – that the public is notified of how to properly use a pesticide, and that stems from the FIFRA,
15 the requirements in FIFRA, that EPA registers pesticides.” Deposition of John R. Fowle, III,
16 Ph.D., DABT (“Fowle Dep.”) at 50.

17 Under California law, Monsanto is responsible for adequately warning consumers of the risks
18 associated with GBH’s. *Webb v. Special Elec. Co.* (2016) 63 Cal.4th 167, 181 (“manufacturers have a
19 duty to warn consumers about the hazards inherent in their products). It is not the responsibility of the
20 EPA. Under both strict liability and negligent failure to warn, the jury must determine whether the
21 *manufacturer* knew (or should have known) of the dangerous propensities of its product and whether
22 the manufacturer adequately warned of the known risk. Any argument or testimony implying that
23 EPA’s registration of GBH’s relieves Monsanto of its legal duty is contrary to California law. As the
24 preemption affirmative defense is no longer in the case any such arguments are irrelevant and highly
25 prejudicial.

26 Indeed, Monsanto’s arguments and testimony rely on the premise that EPA’s labeling decisions
27 have “the force of law.” Def’s Mot. for Sum. J. at 11. This premise has already been rejected by the
28 Court. In finding that conflict preemption is not applicable to this case, the Court cited to *Hardeman*

1 v. *Monsanto Company*, 216 F. Supp. 3d 1038 (N.D. Cal. 2016) for the principle that “EPA’s approval
2 of Roundup’s label would preempt conflicting state law if it had the force of law under *Wyeth*, **but**
3 **finding no indication that EPA’s approval of Roundup’s label had the force of law.”** Preemption
4 Order at 42. Thus, evidence that Monsanto was restricted from warning Plaintiff of Non-Hodgkin’s
5 lymphoma because of EPA registration of GBHs misstates the applicable law.

6 Finally, any argument or testimony that Monsanto could face legal sanctions for warning
7 consumers of the risk of Non-Hodgkin’s lymphoma will unnecessarily confuse the jury as to the
8 appropriate standard to be applied in this case. This evidence would be unfairly prejudicial as there is
9 no evidence that the EPA would have considered labeling “misbranded” or initiated sanctions
10 proceedings against a manufacturer for providing *additional* safety warnings to consumers. The Court,
11 counsel, and jurors would become mired in issues surrounding the interpretation of EPA regulations.
12 The probative value of this speculative testimony would be substantially outweighed by risk of undue
13 consumption of time and confusion of the issues.

14 **II. CONCLUSION**

15 Any argument, suggestion, or testimony that Monsanto is not permitted to warn of risks outside
16 of the EPA approved labeling is inaccurate and prejudicial. Based on the foregoing, Plaintiff Dewayne
17 Johnson respectfully requests that the Court enter an Order granting this motion *in limine* and excluding
18 any argument or testimony that EPA’s registration of GHB’s precluded Monsanto from warning Plaintiff
19 of the risk of Non-Hodgkin’s lymphoma.

20 Respectfully submitted,

21
22 Dated: May 24, 2018

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23
24 By: /s/ Curtis G. Hoke

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mmiller@millerfirmllc.com

tlitzenburg@millerfirmllc.com
choke@millerfirmllc.com

Attorneys for Plaintiff

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