

## Monsanto Manuscript Clearance Form Global Regulatory

**NOTE:** this form needs to be completed and submitted for review at least 4 weeks prior to manuscript submission and a minimum of 2 weeks prior to abstract/presentation submission

Questions regarding completion of this form can be directed to Jeanna Graf [REDACTED] or Kevin Glenn [REDACTED]

Date: 2/29/2012

Please indicate type of publication: ☒ Manuscript ☐ Conference/meeting presentation ☐ Abstract

Has this information been publicly disclosed previously? ☐ No ☒ Yes If yes, where & when? This manuscript reviews glyphosate genotoxicity publications since the Williams et al. (2000) review

(If the information has previously been published, no need to include Patent Scientist & Patent Atty review)

Title: Review of Genotoxicity of Glyphosate and Glyphosate Based Formulations

Author(s): David Saltmiras, Larry Kier (consultant)

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Is this related to a Monsanto collaboration? ☐ No ☒ Yes If yes, Other

Meeting Date & Location at which Manuscript will be presented:

Journal Submitted To: Reg. Toxicol. Pharmacol.


**Lead Author's Comments:** (Please provide information on purpose of presentation/publication and any relevant patents or manuscript disclosures.) This manuscript provide a comprehensive quality check on the large number of genotoxicity publications on glyphosate since the Williams et al. (2000) glyphosate toxicology review manuscript. This work falls under the scope of the EU Glyphosate Task Force and will be a valuable resource in future product defense against claims that glyphosate is mutagenic or genotoxic.

# Title: Review of Genotoxicity of Glyphosate and Glyphosate Based Formulations

**Author Material Transfer Agreement Statement:** (Please refer questions to a patent attorney.) I have reviewed the material transfer and data disclosure requirements of the proposed journal and have discussed any such requirements with my direct supervisor. I ensure that when I submit this manuscript, the journal either does not or will not in this instance require Monsanto to provide restricted plasmids or other materials referenced in our manuscript, or that I will obtain legal approval for any such materials prior to submission of this paper.

I have reviewed that the appropriate individuals that have contributed substantial, direct, intellectual work to this manuscript/presentation are included as authors, and other significant contributors have been appropriately acknowledged.

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<b>Center Lead *</b> Nordine Cheikh, C3NA, (J. Graf)		
<b>Regulatory Team Lead (Crop/Chem)</b> Chemistry; Susan Martino-Catt C3NA (L. Billadeau)		
<b>Crop/ChemTeam Lead</b> Not Applicable		
<b>Regulatory Law</b> Not Applicable		
<b>Regulatory Law (Add'l Reviewer)</b> Brandon Neuschafer, E1NH (J. Wardlow)		
<b>Regulatory Scientific Affairs</b> Eric Sachs, C3NA, (J. Graf)		
<b>Biotechnology</b> Not Applicable		
<b>Patent Scientist</b> Not Applicable		
<b>Patent Scientist (Add'l Reviewer)</b> Not Applicable		
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Forward MCF to other TC representatives for review		x								
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Manuscript fits with crop team's overall strategic positioning and publication strategy			x	x		x	x			x
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