



EPA Activities on New Approach Methods: Update from the Office of Pesticide Programs

MONIQUE PERRON, SC.D.

ENVIRONMENTAL PROTECTION AGENCY

OFFICE OF PESTICIDE PROGRAMS, HEALTH EFFECTS DIVISION

SENIOR SCIENCE ADVISOR

PERRON.MONIQUE@EPA.GOV

Acute Toxicity

- Eye irritation (Clippinger et al., 2021)
 - Provides evaluation of human relevance, strengths, and uncertainties of *in vivo* and *in vitro* studies
 - Comparison of corneas across species
 - Proposes adverse outcome pathway for eye irritation
 - Concludes many *in vitro/ex vivo* methods are equivalent or scientifically superior to *in vivo* rabbit study
- Acute Oral (Hamm et al., 2021)
 - Pilot program to evaluate mathematical tool (GHS Mixtures Equation) as alternative to acute testing in animals
 - Worked with NICEATM to conduct retrospective analyses, which demonstrated the utility of the GHS Mixtures Equation to predict oral toxicity, particularly for formulations with lower toxicity

CUTANEOUS AND OCULAR TOXICOLOGY
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Open access


Human-relevant approaches to assess eye corrosion/irritation potential of agrochemical formulations

Amy J. Clippinger^a, Hans A. Raabe^b, David G. Allen^c, Neepa Y. Choksi^c, Anna J. van der Zalm^a, Nicole C. Kleinstreuer^d, João Barroso^e, and Anna B. Lowit^f

^a PETA Science Consortium International e.V., Stuttgart, Germany ^b Institute for In Vitro Sciences, Inc., Gaithersburg, MD, USA


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Performance of the GHS Mixtures Equation for Predicting Acute Oral Toxicity

Jon Hamm^{a,*}, David Allen^a, Patricia Ceger^a, Tara Flint^b, Anna Lowit^b, Lindsay O'Dell^b, Jenny Tao^b, Nicole Kleinstreuer^c

^a ILS, P.O. Box 13501, Research Triangle Park, NC, 27709, USA
^b Office of Pesticide Programs, U.S. Environmental Protection Agency, 1200 Pennsylvania Ave, NW, Washington, DC, 20460, USA
^c National Toxicology Program Interagency Center for the Evaluation of Alternative Toxicological Methods, National Institute of Environmental Health Sciences, P.O. Box 12233, Research Triangle Park, NC, 27709, USA



Dermal Absorption “Triple Pack”

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Research Article

Retrospective Analysis of Dermal Absorption Triple Pack Data

David G. Allen¹, John Rooney¹, Nicole Kleinstreuer², Anna Lowit³, Monique Perron³

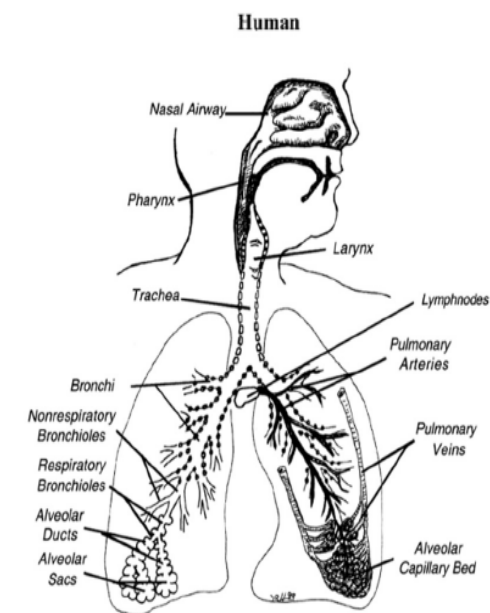
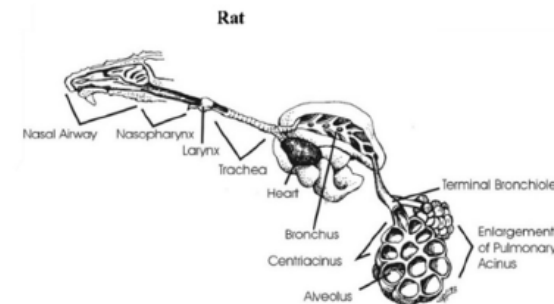
¹Integrated Laboratory Systems LLC, Research Triangle Park, NC, USA; ²National Toxicology Program Interagency Center for the Evaluation of Alternative Toxicological Methods, National Institute of Environmental Health Sciences, Research Triangle Park, NC, USA;


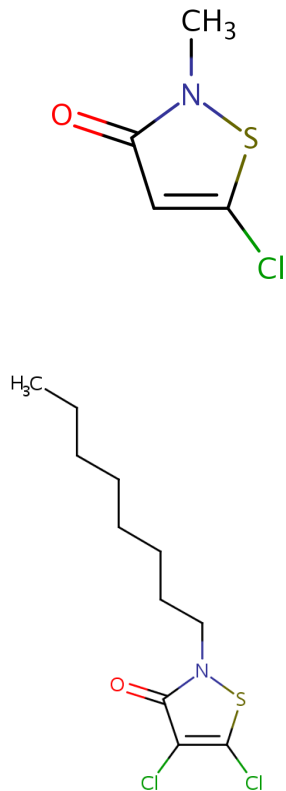
³Office of Pesticide Programs, U.S. Environmental Protection Agency, Washington, DC, USA

- Retrospective analysis of human *in vitro*, rat *in vitro*, and rat *in vivo* studies using similar protocols (e.g., same test material, doses)
- Industry partners provided >30 triple pack studies
- Demonstrates *in vitro* studies alone provide similar or more protective estimates of dermal absorption, with limited exceptions
- Incorporating shift in use of *in vitro* studies into updated OECD Guidance Notes on Dermal Absorption

Inhalation Risk Assessment

- Refined approach using *in vitro* point of departure and human dosimetry modeling incorporated into chlorothalonil risk assessment for Registration Review
- <https://www.regulations.gov/document/EPA-HQ-OPP-2011-0840-0080>
- Evaluating available *in vitro* data for other contact irritants
- Investigating potential to apply similar approach for other pesticides
- Additional projects/discussions to evaluate and compare across *in vitro* assays

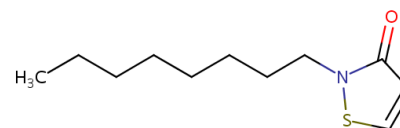
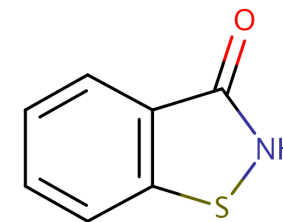
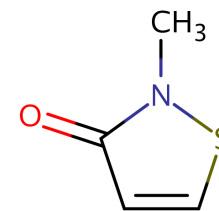
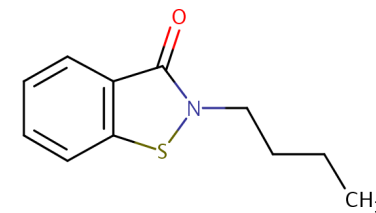




HAZARD
CHARACTERIZATION OF
ISOTHIAZOLINONES IN
SUPPORT OF FIFRA
REGISTRATION REVIEW
April 6, 2020

US Environmental Protection
Agency, Office of Pesticide
Programs, Antimicrobials Division

In Collaboration with the National
Toxicology Program's Interagency
Coordinating Committee for the
Evaluation of Alternative
Toxicological Methods



Isothiazolinone biocides are used as material preservatives to prevent the growth of microbial organisms and are used in industrial processes and consumer products

<https://www.federalregister.gov/documents/2020/05/14/2020-10376/pesticide-registration-review-draft-human-health-and-ecological-risk-assessments-for-several>

Rethinking Carcinogenicity Assessment for Agrochemicals Project (ReCAAP)

- Collaborative project to develop a waiver framework for pesticides
 - Project led by PETA-ISC
 - Retrospective & prospective case studies were developed as part of the weight of evidence (WOE) development
 - Framework provides structure to organize existing critical information, including NAM data, that can be considered by a regulatory authority when making a waiver decision
 - Review by 2020 Scientific Advisory Board (SAB)
 - Manuscript published in June 2022



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Rethinking chronic toxicity and carcinogenicity assessment for agrochemicals project (ReCAAP): A reporting framework to support a weight of evidence safety assessment without long-term rodent bioassays

Gina M. Hilton ^a ✉, Catherine Adcock ^b, Gregory Akerman ^c, James Baldassari ^d, Michael Battalora ^d, Warren Casey ^e, Amy J. Clippinger ^a, Rhian Cope ^f, Amber Goetz ^g, A. Wallace Hayes ^h, Sabitha Papineni ⁱ, Richard C. Pepper ^j, Deborah Ramsingh ^b, Brandy Williamson Riffle ^k, Mitscheli Sanches da Rocha ^k, Natalia Ryan ^g, Edward Scollon ^l, Nicolo Visconti ^m ... Anna Lowit ^c

Developmental Neurotoxicity (DNT)

- International effort to develop DNT NAM battery to assess critical processes of neurodevelopment
- Convened FIFRA Scientific Advisory Panel (SAP) soliciting comment on currently available battery of assays
 - Report published in December 2020: <https://www.epa.gov/sap/use-new-approach-methodologies-nams-derive-extrapolation-factors-and-evaluate-developmental>
- Provide chemical nominations to NTP for additional testing
- OECD DNT NAM Expert Group drafting guidance document on integration of DNT NAMs into regulatory decision-making
- Release of data evaluation records (DERs) and memos to aid in NAM evaluation
 - Docket on regulations.gov: EPA-HQ-OPP-2016-0093



Integration of toxicodynamic and toxicokinetic
new approach methods into a weight-of-evidence
analysis for pesticide developmental neurotoxicity
assessment: A case-study with DL- and L-
glufosinate ☆

Sarah Dobreniecki ^a, Elizabeth Mendez ^a, Anna Lowit ^a, Theresa M. Freudenrich ^b, Kathleen Wallace ^b, Amy
Carpenter ^c, Barbara A. Wetmore ^b, Anna Kreutz ^c, Evgenia Korol-Bexell ^c, Katie Paul Friedman ^b, Timothy J. Shafer ^b
✉

Use of DNT assays in WOE

- DNT study exists for DL-glufosinate; however, L-glufosinate isomers lack *in vivo* DNT study
- Data from *in vitro* DNT assays and toxicokinetics of DL- and L-glufosinate evaluated
- Incorporated data into a WOE assessment to support waiving *in vivo* DNT study for L-glufosinate

Eco Retrospective Analyses

- Fish acute retrospective
 - OPP uses cold freshwater fish, warm freshwater fish, and saltwater fish to assess acute risks (200 fish or more used)
 - Is there a consistently more sensitive fish across all compounds and can we reduce data sets to two or even one fish study?
 - Manuscript in preparation
- Avian reproductive retrospective
 - Currently 2 species are required: Mallard Duck and Bobwhite Quail
 - How could a protective risk assessment be done with only one species?
 - Currently in data curation and exploration stage

QSAR Evaluation for Acute Lethality

- QSAR for Fish LC₅₀
 - Data submitted to EPA by registrants
 - Much larger database than main current tool (ECOSAR) and with pesticide-specific data
 - Working on peer-reviewed publication and web-based GUI
- Collaborative Acute Toxicity Modeling Suite (CATMoS)
 - 35 participants/groups from around the globe representing academia, industry, and government contributed to the development
 - Evaluate potential replacement of rat acute oral toxicity study with LD₅₀ predictions
 - Manuscript in preparation

Honeybees

- Five tests (adult acute contact, adult acute oral, larval acute, adult chronic, and larval chronic toxicity) currently required
 - Examining EFED data holdings to determine whether all 5 tests are needed to do risk assessment
 - Manuscript published in PLOS One, April 7, 2022: “A retrospective analysis of honey bee (*Apis mellifera*) pesticide toxicity data”
- SeqAPASS (Sequence Alignment to Predict Across Species Sensitivity) model being evaluated to extrapolate honeybee data to non-*Apis* bees
- Colony models (e.g., VarroaPOP)



Updated OPP Webpage

Main page:

<https://www.epa.gov/pesticide-science-and-assessing-pesticide-risks/strategic-vision-adopting-new-approach-methodologies>

Metrics: <https://www.epa.gov/pesticide-science-and-assessing-pesticide-risks/adopting-21st-century-science-methodologies-metrics>



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Adopting 21st-Century Science Methodologies —Metrics

On Sept. 10, 2019, EPA Administrator Andrew Wheeler issued a [directive](#) to prioritize efforts to reduce animal testing, which included the goals of reducing mammal study requests and funding 30 percent by 2025 and eliminating them by 2035. The administrator's directive specifically charged the Agency to establish baselines, measurements and reporting mechanisms to track its progress.

Additionally, the U.S. Government Accountability Office (GAO) released a [report](#) to Congress in 2019 recommending that Federal agencies develop metrics to assess the progress made toward reducing, refining and replacing animal use in testing. The activities and policies EPA has implemented over the past several years demonstrate significant impacts in reducing the number of animals used in testing and saving resources for the Agency and stakeholders.

Details on these reduction and replacement metrics are described on their respective pages. EPA's Pesticide Program reports its progress in the [Annual Reports on PRIA Implementation](#), and began to release specific metrics in FY2015.

On this page:

- [Hazard and Science Policy Council \(HASPOC\) metrics](#)
- [Chemistry and Acute Toxicology Science Advisory Council \(CATSAC\) metrics](#)
- [Acute Dermal Retrospective Waiver Requests](#)



Evaluate regulatory flexibility for accommodating



Develop baselines and metrics for assessing



Establish scientific confidence and demonstrate



Develop NAMs that fill critical information gaps



Engage and communicate with stakeholders

EPA NAM Workplan

- Original EPA NAMs Work Plan released in June 2020, which laid out the Agency's objectives and strategies
- Committed to regularly reviewing the work plan and acknowledge the work plan will evolve as EPA's knowledge and experience grows, and as outside experts offer their perspectives and contributions
- Work plan recently updated in December 2021
 - Main objectives and strategies were left unmodified

Thank You!
