

INTERNATIONAL EFFORTS
IMPLEMENTING NAMS FOR
ASSESSING DEVELOPMENTAL
NEUROTOXICITY IN CHEMICAL RISK
ASSESSMENT

Iris Mangas, Regulatory Toxicologist Pesticides Peer Review Unit



# DEVELOPMENTAL NEUROTOXICITY (DNT)

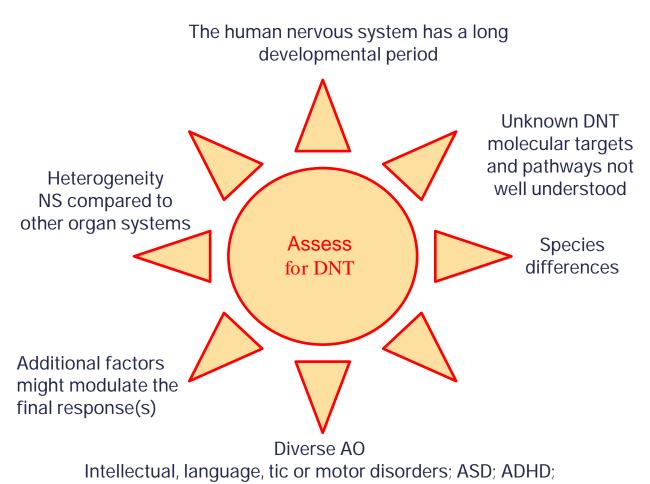
DNT chemicals: diverse set of substances that have the potential to interfere with the normal development of the nervous system, which, if perturbed without compensation, may lead to adverse effects on nervous system structures and/or functions.

Evaluating DNT as one of the endpoints to assess chemical safety is embedded in various regulations and several TGs are available for DNT in vivo screening.

However, use have been limited, potentially due to their methodological and interpretation complexity, ethical concerns and high costs.

## OECD N377., Key success factors:

- Collaborative effort
- Use the data in real case studies

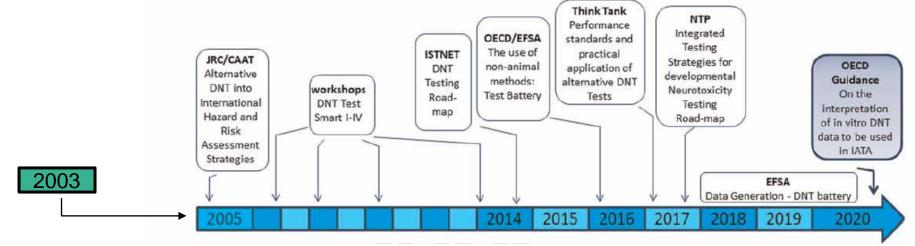


Developing and identifying an appropriate test system for DNT testing is challenging.

NDD



# BUILDING THE DNT IVB



Advances using in vitro neural models combined with technology for higher throughput analysis has fostered the development of assays.

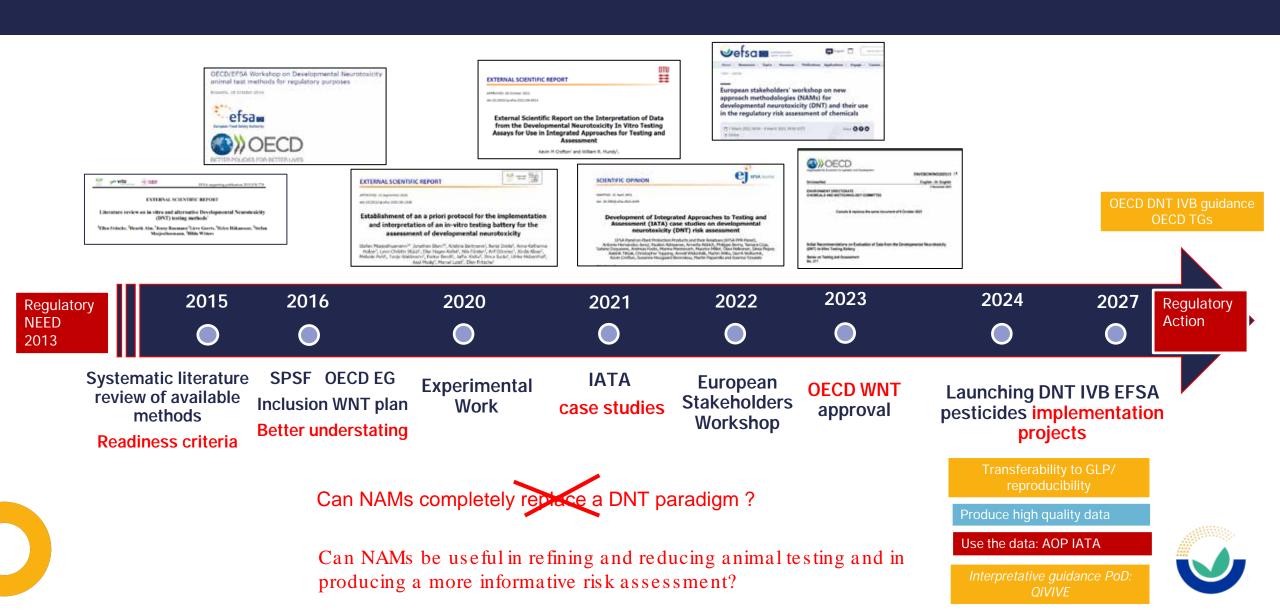


Pamela Leina, b, \*, Ellen Silbergelda, Paul Lockea, Alan M. Goldberga

Bringing together basic researchers in the fields of developmental neurobiology and mechanistic toxicology with policy analysts and decision makers



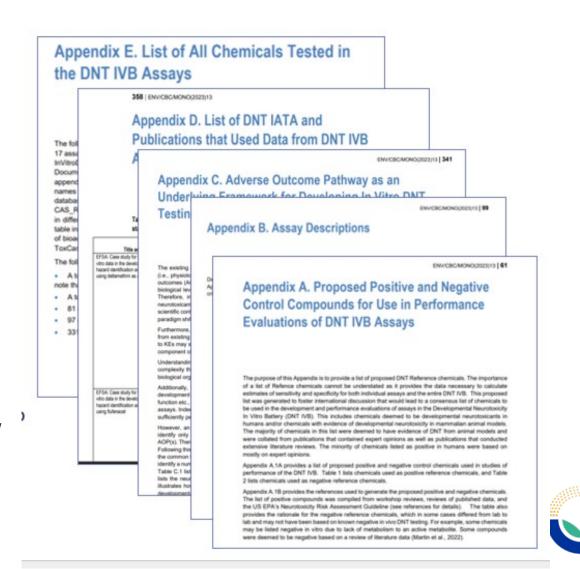
## EFSA SUPPORT TO THE INTERNATIONAL EFFORT



# COMMON NEEDS: THE OECD NO. 377



- •1. Review on the science on DNT IVB and provide GD on how to evaluate the DNT IVB and its relevance for assessing DNT
- •2. Context (AOP informed IATA) and description DNT IVB. + Appendix B
- •3. WoE considerations
- •4. Integration of evidence. Standardize use.

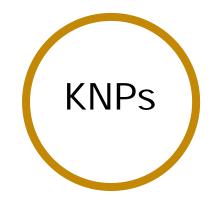


# THE OECD DNT IVB INITIAL RECOMMENDATIONS DOCUMENT NO.377

- The purpose is to provide guidance on **how to evaluate** *in vitro* data from the assays comprising the DNT IVB (e.g., hit vs non-hit, biological coverage). Understand the uncertainties in the assays and the data outputs.
- It is not intended to provide interpretative guidance on the use of results in human risk assessments. Specific criteria for such use will likely be developed by regulatory agencies who will determine acceptability based on their needs.



- A battery of assays judged ready for use by international working groups
- Tested a common set of chemicals 81
- 3. Assay descriptions expanded version of OECD GD211



- Assays are designed to recapitulate basic processes of neurodevelopment
- 2. AOPs as a Framework for In Vitro DNT. Review of existing DNT AOPs

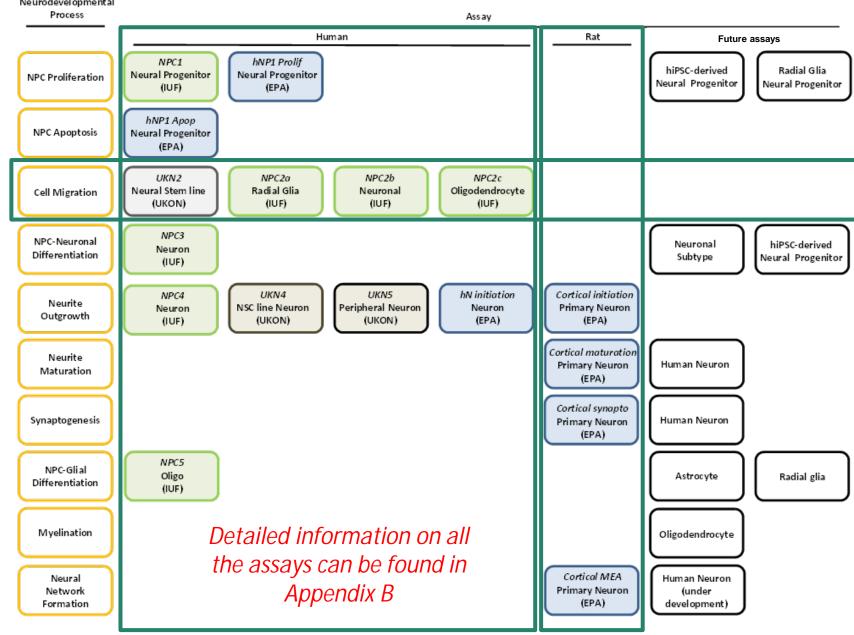


- 1. As a tool to integrate it in an WoE with all other available data.
- 2. Balancing the uncertainty in the data, with the uncertainty acceptable for the regulatory decision.
- 3. Series of proof of concept case studies **different CoU** 6

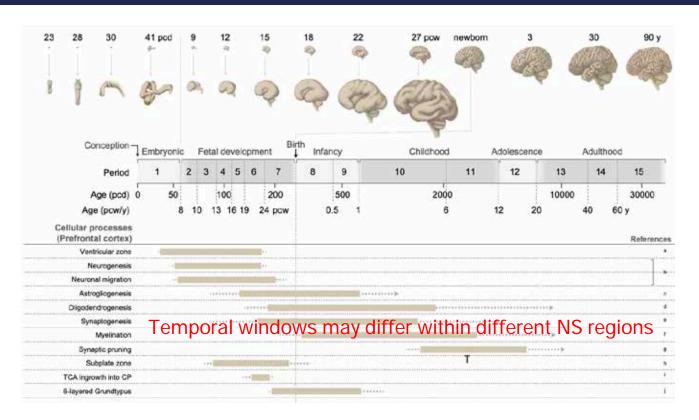


Neurodevelopmental

- Assays are not based on MiE (e.g., receptors).
- The assays in the DNT IVB instead measure neurodevelopmental processes that are critical for normal development of the nervous system and based on the following assumptions:
- 1. Changes in these cellular processes will reflect the integration of chemical disruptions in multiple up-stream molecular events.
- Chemicals affecting one or more of these neurodevelopmental processes in vitro have the potential to do so in vivo if exposure.



# WHAT DO THEY MEASURE? KNPS



Brain development is a highly complex procedure that covers time and a delicate balance of a large variety of key neurodevelopmental processes

Silbereis et al. 2016

Biological processes at cellular level necessary for NS development. Multiple cellular pathways.

DNT IVB KNDP	DEVELOPMENTAL TIMING test system	
	Embryonic	Fetal
NPC proliferation		Χ
NPC apoptosis		X
NPC migration		Х
Radial glia migration		X
Neuronal migration		Χ
Oligodendrocyte migration		Χ
Neuronal differentiation	X	X
Neurite outgrowth	X	X
Neuronal maturation	X	Χ
Synaptogenesis	X	X
Oligodendrocyte differentiation		X
Neuronal network formation	Х	Х

Overview of neurodevelopmental timing of the current DNT IVB assays

# **ELEMENTS FOR WOE: IATA**

- 1. Generic issues for all in vitro assays.
- 2. Issues specific for DNT IVB assays (unknown metabolic competence, applicability domain, nominal concentrations,
- 3. Transparency in data analyses
- 4. Individual assay interpretation
- 5. Evaluation of data from the entire battery of assays

 Treat the data from the DNT IVB like all other data used in a WoE in hazard assessments



# ACCEPTABILITY EFS A DNT IATA CASE STUDY PROJECT EU PESTICIDES REGULATORY CONTEXT OF USE AS AN EXAMPLE

Establishment of EFSA PPR Panel IATA WG

April 2021 Submission to OECD IATA case study program WPHA endorsement and publication flufenacet and deltamethrin IATA case studies

EC mandate Acetamiprid : Test an existing IATA\* MS experts endorsed the IATA conclusions during the Peer Review Meeting for deltamethrin RMS Austria EFSA PPP Panel endorsed IATA on acetamiprid

MS experts endorsed the IATA conclusions for acetamiprid

May 2019

June 2021

Sep 2022

Jan 2023

June 2023

April 2024



6 WG experts + hearings Observers: EC JRC, ECHA, MS, industry, NGO



EFSA PPP Panel endorsed DLM and Flufenacet IATAs PPR Panel agreed with the approach for DNT assessment of pesticides



Reviewed by Canada, Germany, Italy, Australia, Netherlands, OECD, BIAC.



EUTOX Risk AOP informed IATA DNT IVB for quantitative risk assessment

⊗» OECD



EFSA IATA on acetamiprid published in EFSA Journal and accepted by EC Launching outsourcing projects

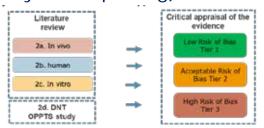


# EFSA AOP INFORMED IATA CASE STUDIES WOE WORKFLOW: EVIDENCE BASED



#### DNT HAZARD IDENTIFICATION PROBLEM FORMULATION

**Step 1.** Systematic literature review and critical appraisal to assess data quality (reliability and reporting)



In the context of the European pesticides Regulation (EU) 283/2013 and 1107/2009

**Step 2**. Data integration: AOP framework



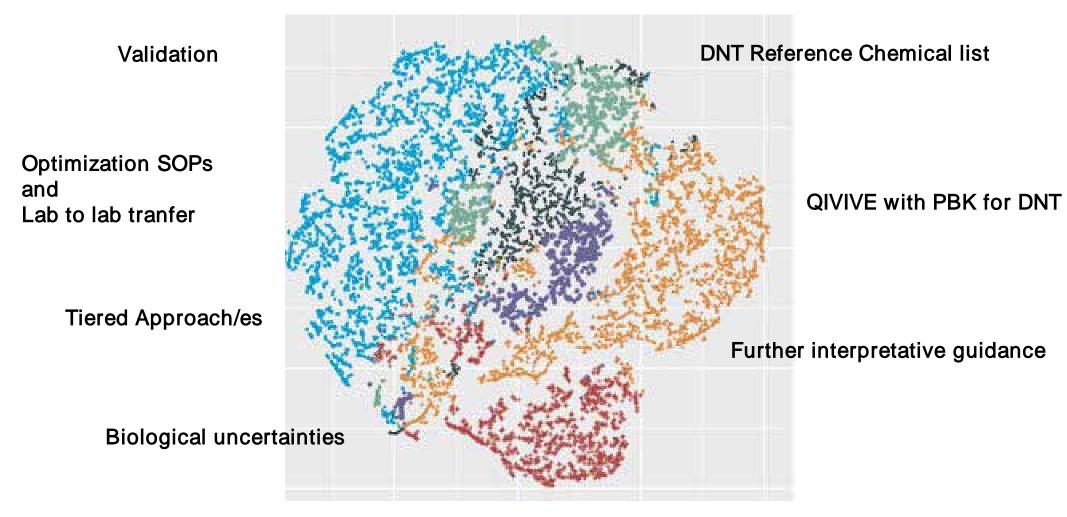
• Dose and temporal concordance considerations

Step 3. Data gap and uncertainty analysis

Step 4. AOP informed IATA conclusion and Regulatory conclusion

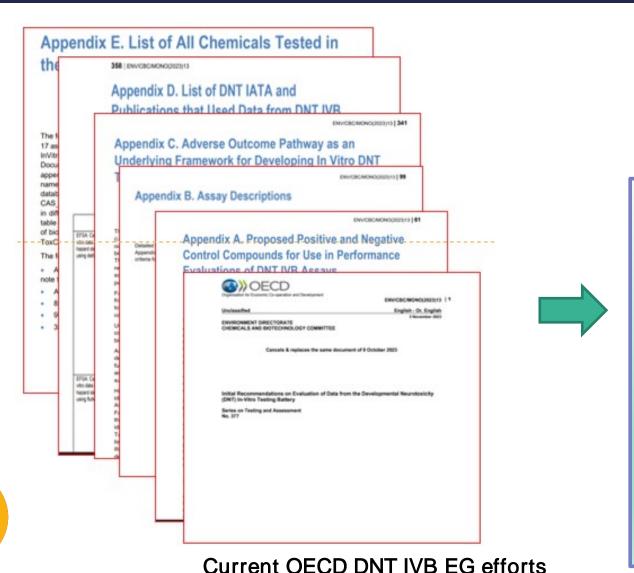


## MOVING FORWARD: REGULATORY IMPLEMENTATION





# TOWARDS AN OECD GUIDANCE: ONGOING WORK REFERENCE CHEMICAL LIST



Appendix A: OECD Guidance document provided a draft list.

Ongoing update: developing an OECD consensus list

A soon to be released EFSA report

"Recommended DNT Reference Chemical Test Set For In Vitro Assay Development"

Mundy and Crofton., 2024

Updates and prioritizes a list of 164 chemicals

# TOWARDS AN OECD GUIDANCE: ONGOING WORK COMMON AGREED ANALYTICAL PIPELINE

There are a number of on-going efforts to begin analyses of all the data in the DNT IVB that will foster understanding of the worth of the assays and the battery.



#### Research Article

#### The Impact of Biostatistics on Hazard Characterization Using In Vitro **Developmental Neurotoxicity Assays**

Hagen Eike Kellel<sup>1</sup>, Stefan Maziasthusmann<sup>1</sup>, Kristina Bartmann<sup>1</sup>, Jonathan Blum<sup>2</sup>, Arif Dönmez<sup>1</sup>, Nils Förster<sup>4</sup>, Jördiz Klozel, Axel Mozigl, Melanie Pakil, Marcel Leizel, Martin Scholzell and Ellen Fritzchell. J.

UCF - Leibnir Research Institute for Environmental Medicine, Disseldorf, Germany; In vitro Toxicology and Biomedicine, Dept imaginated by the Doeserkamp-Zoinden Foundation, University of Konstaur, Koustaur, Germany, Medical Faculty, Heinrich-Heine-University, Disseldorf, Germany, "Bioantematics Group, Rule University Bochum, Bochum, Germany, "Brandl University, London, UK.

in chemical safety assessment, benchmark concentrations (BMC) and their associated uncertainty are needed for the toxicological evaluation of in vitro data sets. A BMC estimation is derived from concentration response modelling and results from various statistical decisions, which depend on factors such as experimental design and assay endpoint features. In current data practice, the experimenter is often responsible for the data analysis and therefore relies on statistical software. often without being aware of the software default settings and how they can impact the outputs of data analysis. To provide more insight into how statistical decision-making can influence the outcomes of data analysis and interpretation, we have developed an automated plafform that includes statistical methods for BMC estimation, a novel endpoint-specific hazard classification system, and routines that flag data sets that are outside the applicability domain for an automatic data evaluation. We used case studies on a large dataset produced by a developmental neurotoxicity (DNT) in vitro battery (DNT IVB). Here we focused on the BMC and its confidence interval (CI) estimation as well as on final hazard classification. We identified five crucial statistical decisions the experimenter must make during data analysis; choice of replicate overtaging, response data normalization, regression modelling, BMC and CI estimation, and choice of benchmark response levels. The insights agrined are intended to raise more awareness among experimenters on the importance of statistical decisions and methods but also to demonstrate how important Ri-for-purpose, internationally harmonized and accepted data evaluation and analysis procedures are for objective hazard classification.

#### JOURNAL ARTICLE

#### Integrating Data From In Vitro New Approach Methodologies for Developmental Neurotoxicity @

Kelly E Carstens, Amy F Carpenter, Melissa M Martin, Joshua A Harrill, Timothy J Shafer, Katie Paul Friedman

Toxicological Sciences, Volume 187, Issue 1, May 2022, Pages 62-79, https://doi.org/10.1093/toxsci/kfac018

Published: 16 February 2022











#### Abstract

In vivo developmental neurotoxicity (DNT) testing is resource intensive and lacks information on cellular processes affected by chemicals. To address this, DNT new approach methodologies (NAMs) are being evaluated, including: the microelectrode array neuronal network formation assay; and high-content imaging to evaluate proliferation, apoptosis, neurite outgrowth, and synaptogenesis. This work addresses 3 hypotheses: (1) a broad screening battery provides a sensitive marker of DNT bioactivity; (2) selective bioactivity

Standardize and simplify analysis of data



Concentration Response BMC by Lab Integrative Analyses Resources

Developmental NeuroToxicity Data Integration and Visualization Enabling Resource (DNT-DIVER)



Research shows that a child's developing nervous system is far more vulnerable to chemical. exposures than an adult nervous system. Recent increases in the rise of neurodevelopmental disorders such as attention deficit hyperactivity disorder (ADHD), dyslexia, and autism spectrum disorder have prompted scientific interest in the potential contribution of environment toxicants to these

Traditional animal, or in vivo, studies provide important information about developmental neurotoxicity (DNT) but they are time and resource intensive. NTP has also developed more rapid screening tools that use human cell-based, or in vitro, assays, as well as alternate animal models such as zebratish and planaria to identify toxicants with potential for DNT. Multiple tests, or assays, are often required to represent the complexity of the developing nervous system, but that can make it challenging to compare and summarize results

NTP designed the Developmental NeuroToxicity Data Integration and Visualization Enabling Resource (DNT-DIVER) to analyze, compare, and visualize multiple DNT assays in an interactive webapplication.



# TOWARDS AN OECD GUIDANCE: ONGOING EFFORTS TRANSFERABILITY, TESTING AND MORE IATAS







Using only pesticides approved in EU

Test system accesibility standarization



# **Environmental Neurotoxicants**

Generate and use the data for assessing for NT and DNT Transferability DNT IVB Produce guidances and databases

Interpretative guidance qiVIVE Guidance Databases Develop regulatory action

### **Brain Health**

Develop AOPs
Molecular characterization
test systems
Refine the testing strategy
Expand the applicability
domain

Only some









Implementation/standardization of biokinetic data in the DNT IVB and PBK models is needed

There are a number of on-going efforts to transfer the DNT IVB or some of the assays to "naïve" laboratories and to develop new assays. Here EFSA plan to contribute to this international and collaborative efforts is summarized

# LESSONS AND LEARNINGS

- The DNT IVB successfully used, accepted and trusted by the stakeholders. The DNT IVB
  demonstrated to provide relevant, high-quality data on information regarding DNT potential of
  chemicals that are perturbing early cellular processes that are difficult to measure in vivo.
- The DNT-IVB data is publicly available and should be integrated in the regulatory pesticides risk assessment in EU. The available data for approved pesticides in EU is limited. To make a further regulatory action (change in data requirements) the transferability to naïve laboratories able to conduct the assays under GLP and make them commercially available is needed.
- AOP informed IATA is the EFSA current recommended framework to integrate the DNT IVB mechanistic data in pesticides RA since allows identification and quantification of uncertainties and transparent report.
- The final aim is to have an internationally agreed **interpretative guidance** and an **agreed standard workflow** facilitating the mutual use in the regulatory frameworks for quantitative risk assessment while retaining necessary flexibility depending on the context of use and advancement of science.
- This future work will continue to imply international and collaborative effort paving the way also a broader use of NAMs.

# **ACKNOWLEDGMENTS**

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- Mary Gilbert (US EPA)
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# THANK YOU VERY MUCH STAY CONNECTED

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