



Research centre
for toxic compounds
in the environment

Ecotoxicology

New topics and future issues

Ludek Blaha + ecotox colleagues

cecoen



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OP Research and
Development for Innovation



Take home messages from this presentation

- Traditional (eco)toxicity testing (based on simple standardized bioassays) and related chemical risk assessment is likely to change in this century...
- ... towards the use of mechanistic data and knowledge (omics) – through – for example - Adverse Outcome Pathways, (AOPs) and mathematical models
 - The paradigm shift is strongly promoted by influential players – OECD, US EPA, European Commission (example shown – OECD AOPWiki)
 - Also in line with minimizing use of animals and implementation of „3R“ policies (examples shown)
 - Toxicological predictions = computational (AI) models are becoming more and more advanced

Current approaches

(black box of apical endpoints)

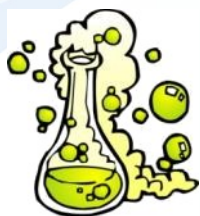
VS

Future

(mechanistic understanding & AOPs)

Hazard assessment

Traditionally – Evaluation of adverse effects using the whole organism models



Chemical



Organism



Adverse Effects

Death
Altered Reproduction
Inhibition of Growth

Tumorigenicity
Skin irritation

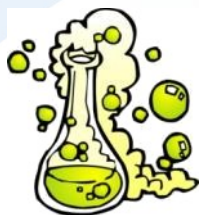
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**REGULATORY FOCUS
(APICAL ENDPOINTS)**

Hazard assessment

Traditionally – Evaluation of adverse effects using the whole organism models



Chemical



Organism



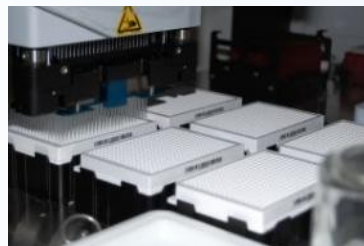
Adverse Effects
Death
Inhibition of Growth
Altered Reproduction
Tumor
Skin irritation
...



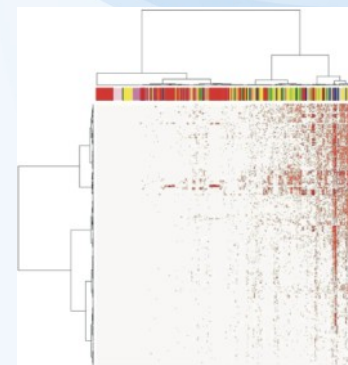
New – Ex vivo / in vitro / In chemico / In silico Methods



10^4 Chemicals



HTS
High-Throughput-Screening



**Chemical-biological interactions,
Mechanistic Toxicological Data**

Key task/question:

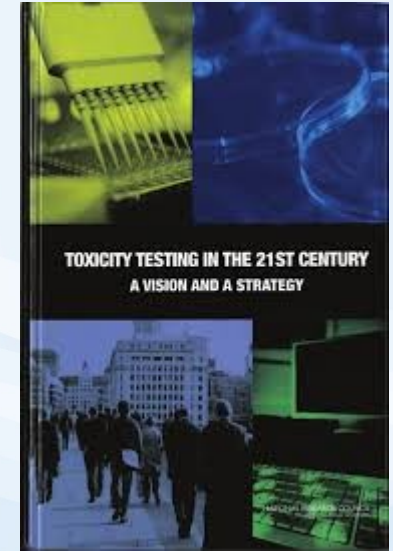
How to link MECHANISTIC INFORMATION with APICAL ENDPOINTS ?

MoA and omics are supported by strategic documents

Toxicity Testing in the 21st Century: A Vision and a Strategy

US National Academies of Sciences

<http://www.nap.edu/catalog/11970.html>



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Computational Toxicology Research

You are here: [EPA Home](#) » [Research & Development](#) » [CompTox](#) » [ToxCast™](#)

Key Links

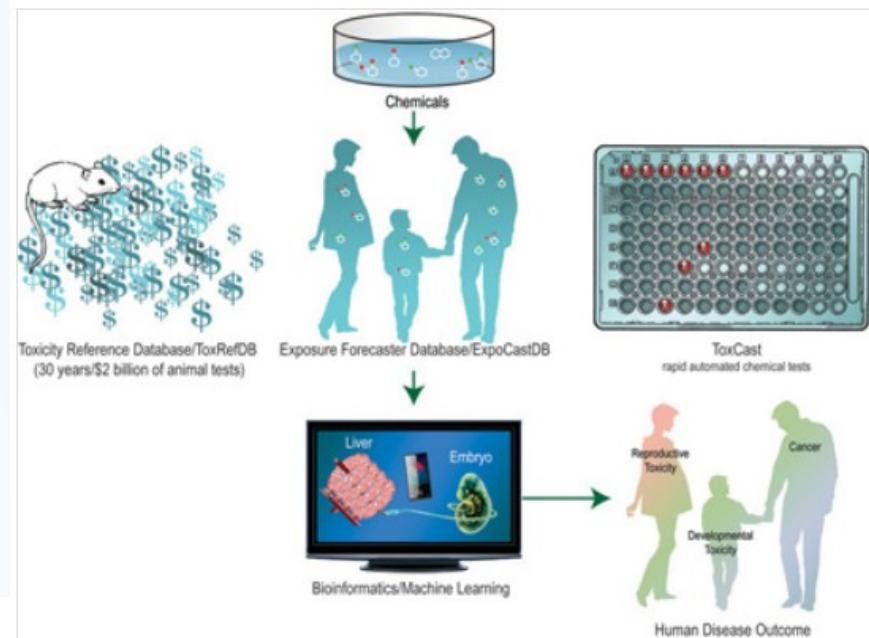
[CompTox Home](#)
[Basic Information](#)
[Organization](#)

[Research Projects](#)
[Chemical Databases](#)
[CompTox Events](#)

R
S
C

ToxCast™

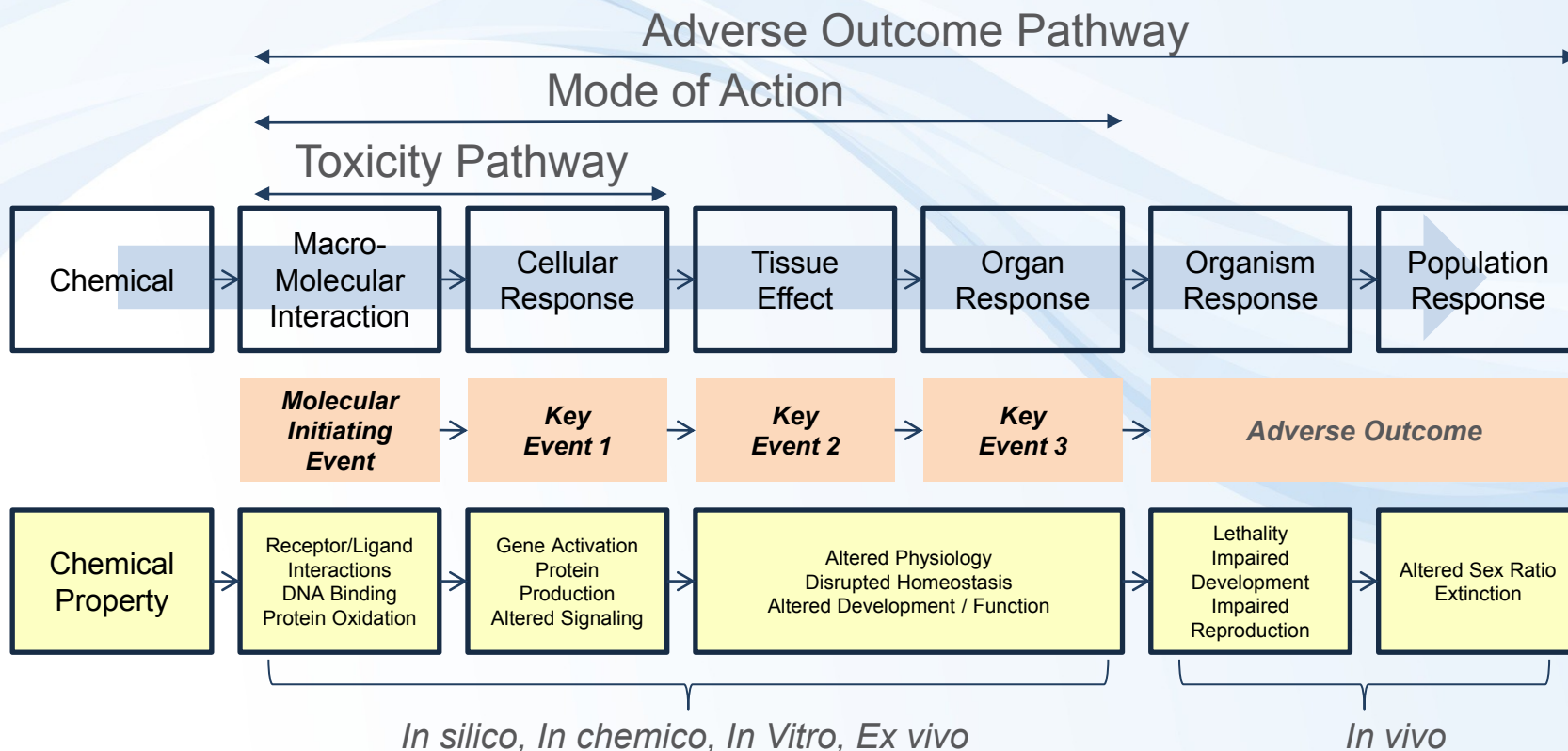
Screening Chemicals to Predict Toxicity Faster and Better



How ToxCast Fits Into CompTox Research




Adverse Outcome Pathways




The **EXISTING KNOWLEDGE** is used **to link the** two anchor points: **Molecular Initiating Event (MIE)** and **Adverse Outcome (AO)** **via a series** of intermediate steps: **Key Events**

AOP = Global strategy with support from OECD, EU, USA



OECD.org

Data Publications More sites News Job vacancies



BETTER POLICIES FOR BETTER LIVES

> A to Z

OECD Home About Countries Topics Français

[OECD Home](#) > [Chemical safety and biosafety](#) > [Testing of chemicals](#) > Adverse Outcome Pathways, Molecular Screening and Toxicogenomics

- > Testing of chemicals
- > Assessment of chemicals
- > Risk management of chemicals
- > Chemical accident prevention, preparedness and response
- > Pollutant release and transfer register
- > Safety of manufactured nanomaterials
- > Agricultural pesticides and biocides
- > Biosafety - BioTrack

Adverse Outcome Pathways, Molecular Screening and Toxicogenomics

WHAT'S NEW

SURVEY ON ADVERSE OUTCOME PATHWAYS (AOPS) TO IDENTIFY DEVELOPMENT PRIORITIES

The OECD has launched a survey to explore the utility of AOPs for regulatory assessment of chemicals and to identify development priorities. The objective is to collect feedback on how the AOP concept and/or existing AOPs are already being used for regulatory purposes, to understand where they fall short regarding their utility, and to identify what directions and priorities future AOP development work should embrace to increase their impact on regulatory toxicology and chemical risk assessment.

The survey is mainly for chemical safety regulators who are experiencing a transition in their work towards an increased use of 'alternative' methods and AOPs. However, stakeholders that come from the regulated community and environmental NGOs are also welcome to participate.

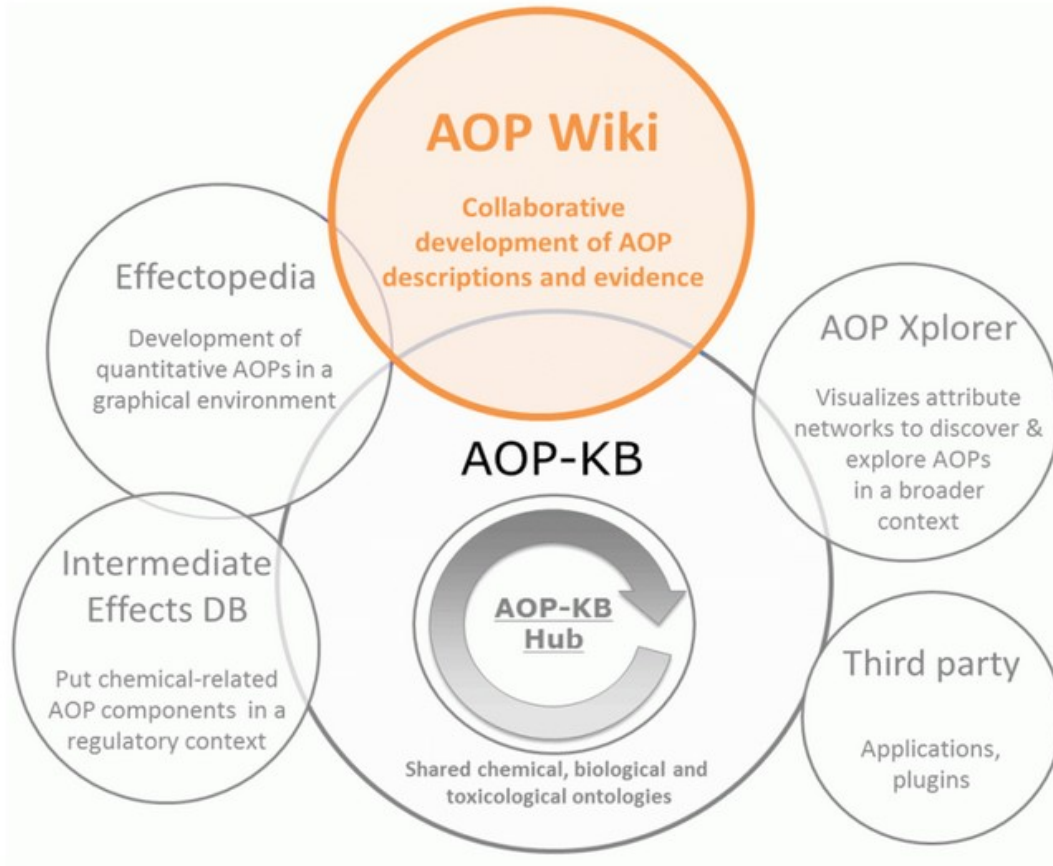
> **The survey is now closed. Thank you for your submissions.**

<http://www.oecd.org/chemicalsafety/testing/projects-adverse-outcome-pathways.htm>



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Please click on any of the AOP-KB elements you want to use.
Please note that the AOP-KB is work in progress and more elements will become available over time.

<http://aopkb.org/>

Key documents

OECD Guidance document and a template for developing and assessing adverse outcome pathways (Series No. 184, Series on Testing and Assessment)

Handbook for AOP developers

AOP Wiki

- https://aopkb.org/aopwiki/index.php/Main_Page
- Wiki-based platform for development of AOPs
- Only members of an OECD AOP development project can create / edit AOPs



What AOPs are now in AOP Wiki (autum 2024) – total 501



OECD Endorsed (WNT and TFHA)	35	1x ecotoxicology: Aromatase inhibition leading to reproductive dysfunction (in fish)
Status: EAGMST Under review	20	
Status: Under Development	83	
(No status: working versions)	363	



<https://aopwiki.org/aops>

- OECD Extended Advisory Group on Molecular Screening and Toxicogenomics (EAG MST)
- The Working Group of the National Coordinators of the Test Guidelines Programme (WNT)

No title search results matched your request

AOP Fulltext Search Results

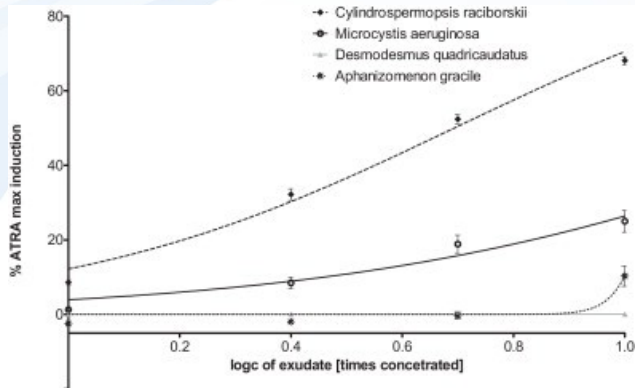
Filtered AOP Count: 5

?

ID	Title	Point of Contact	License	MIE	AO	OECD Status	OECD Project
			Clear			Clear	
201	Juvenile hormone receptor agonism leading to male offspring induction associated population decline	Knut Erik Tollefsen	BY-SA	<ul style="list-style-type: none"> Activation, Juvenile hormone receptor 	<ul style="list-style-type: none"> Increased, Male offspring Decline, Population Alteration, Food-web structures 		
113	Glutamate-gated chloride channel activation leading to acute mortality	Helen Poynton	BY-SA	<ul style="list-style-type: none"> Activation, Glutamate-gated chloride channel 	<ul style="list-style-type: none"> Increased Mortality Decreased, population 1 		1.29
4	Ecdysone receptor agonism leading to incomplete ecdysis associated mortality	Knut Erik Tollefsen	BY-SA	<ul style="list-style-type: none"> Increase, Ecdysone receptor agonism 	<ul style="list-style-type: none"> Increase, Mortality 		
505	Reactive Oxygen Species (ROS) formation leads to cancer via inflammation pathway	John Frisch	BY-SA	<ul style="list-style-type: none"> Increased, Reactive oxygen species 	<ul style="list-style-type: none"> Increase, Cancer 		
360	Chitin synthase 1 inhibition leading to mortality	Simon Schmid	BY-SA	<ul style="list-style-type: none"> Inhibition, Chitin synthase 1 	<ul style="list-style-type: none"> Increase, Mortality 	WPHA/WNT Endorsed	1.94

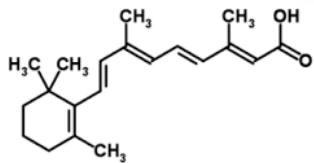
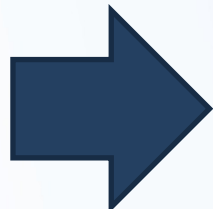
AOP Example from RECETOX:

Modulation of RAR/RXR → developmental toxicity in fish → feeding → population growth

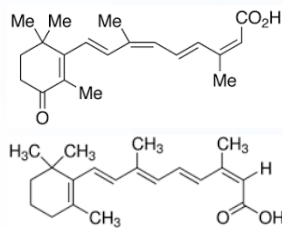


Activation of RAR/RXR

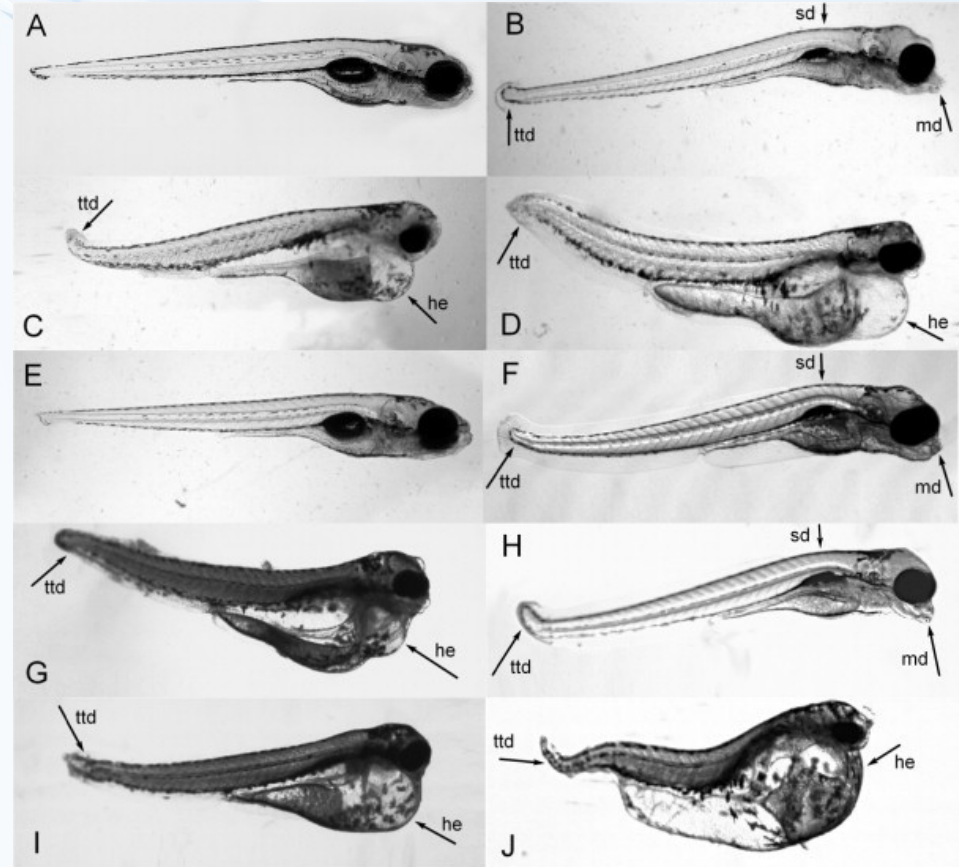
in P19/A15 cells by atRA and cyanobacterial metabolites



atRA



other RAs in cyanos



ZF exposed to ATRA and cyanobacterial (120 hpf) - Control (A), exudates of *C. raciborskii* 3.3 (B) and 10 (C), *M. aeruginosa* 10 (D) and *D. quadricaudatus* 17 (E). ATRA 4 µg/L (13.3 nM) (F), 12 µg/L (40 nM) ((G) and (H)), 36 µg/L (I) and 108 µg/L (J).

Related Projects & Studies & Databases

- **TOXNET** - <http://toxnet.nlm.nih.gov/>
 - searching databases on toxicology, hazardous chemicals, environmental health, and toxic releases
- **Tox21** - <http://www.epa.gov/ncct/Tox21/>
 - 10,000 chemicals
 - 14 concentrations, 4 logs, 3 replicates
 - 1536 well plates, 2-8 uL volumes
 - 50+ assays
- **ToxCast** - <http://www.epa.gov/ncct/toxcast/>
 - App. 2000 chemicals
 - 700+ assay, 300 signaling pathways
 - DATA AVAILABLE iCSS Dashboard
 - <http://actor.epa.gov/dashboard>
 - <http://ww.epa.gov/ncct/toxcast/data.html>



Related Projects & Studies & Databases

- **ToxRefDB (Toxicity Reference Database)**
 - *in vivo* toxicological data
 - <http://actor.epa.gov/toxrefdb/faces/Home.jsp>
- **ExpoCast**
 - information on human exposures
 - <http://www.epa.gov/ncct/expocast/>
- **Human Toxome Project**
 - information on human exposures
 - <http://www.ewg.org/sites/humantoxome/>
- **Agriculture Health Study**
 - Occupational Exposure to Pesticides – a cohort study
 - <http://aghealth.nih.gov/>

Summary

- **Toxicology is about doses**

- The goal is LD(LC)50 or NOAEL/NOEC



- **Legislation defines**

... what assays and how to do them

- About 30 assays
- The most widely used standard - OECD Guidelines for Testing of Chemicals



- **Replacing „black box“ in traditional testing**

- Synthesis of mechanistic and omics data
- Adverse Outcome Pathways
- Strategically supported by OECD, EU, USA



Environmental exposures → Transgenerational effects (Epigenomics)

Maternal predator-exposure has lifelong consequences for offspring learning in threespined sticklebacks



Daniel P. Roche, Katie E. McGhee* and Alison M. Bell

School of Integrative Biology, University of Illinois, Urbana, IL 61801, USA

*Author for correspondence (kemcghee@illinois.edu).

Stress

→ multigeneration effects



Epigenetics

→ DNA methylations

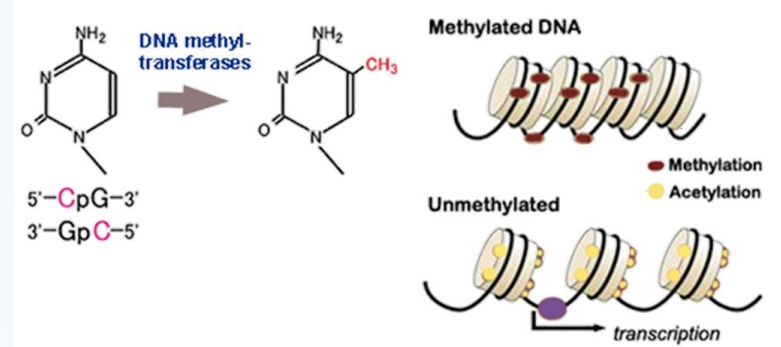


Table 1. Behaviours (mean \pm s.e.) of the offspring from the maternal treatments.

	offspring of predator-exposed mothers (s)	offspring of unexposed mothers (s)
initial exploratory behaviour (day 1: 09.00):		
latency to first begin moving	49 \pm 30	56 \pm 20
latency to enter either chamber for the first time	330 \pm 70	326 \pm 78
learning the colour association:		
day 1 (09.00): latency to find food reward	426 \pm 65	427 \pm 61
day 3 (09.00): latency to find food reward	533 \pm 48	304 \pm 74
day 5 (09.00): latency to find food reward	337 \pm 61	158 \pm 68

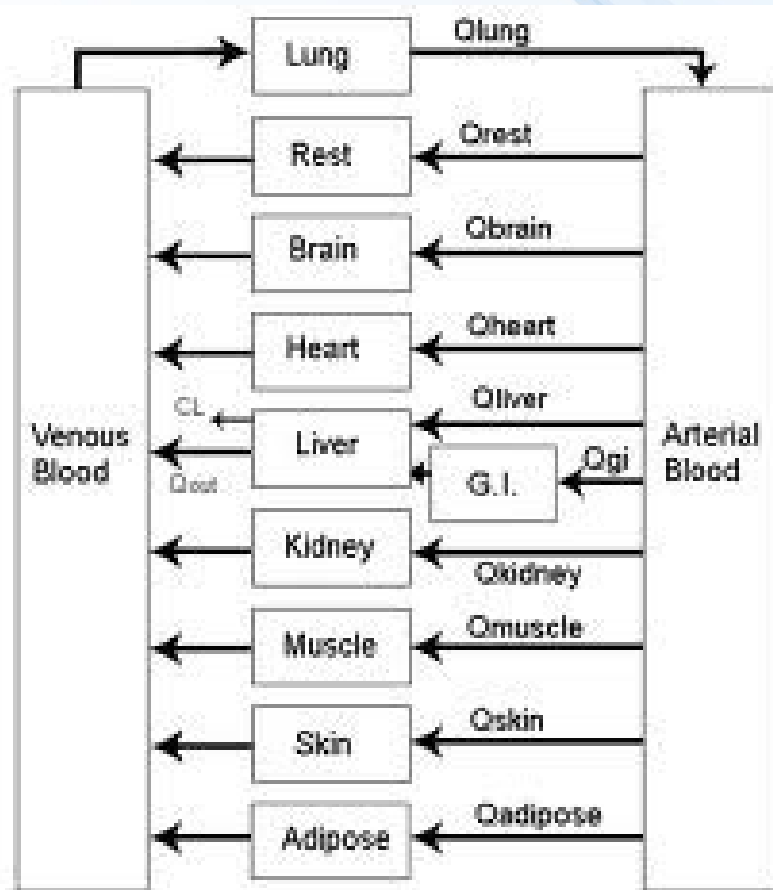
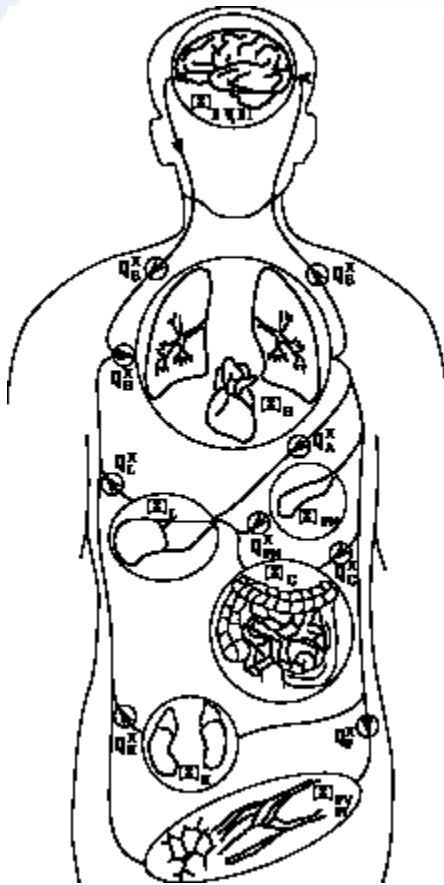
2x difference

COMPUTATIONAL (ECO)TOXICOLOGY

PBPK models

PBPK (PBTK)

Physiologically based pharmacokinetic (toxicokinetic) models



Fragmentation of a complex system to „boxes“

→ All Processes described by arrows (mathematical equations)



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Example – computational toxicology for EDCs

Li et al. *BMC Systems Biology* 2011, 5:63
<http://www.biomedcentral.com/1752-0509/5/63>



RESEARCH ARTICLE

Open Access

A computational model of the hypothalamic - pituitary - gonadal axis in female fathead minnows (*Pimephales promelas*) exposed to 17α -ethynylestradiol and 17β -trenbolone

Zhenhong Li¹, Kevin J Kroll², Kathleen M Jensen³, Daniel L Villeneuve³, Gerald T Ankley³, Jayne V Brian⁴, María S Sepúlveda⁵, Edward F Orlando⁶, James M Lazorchak⁷, Mitchell Kostich⁷, Brandon Armstrong⁸, Nancy D Denslow² and Karen H Watanabe^{1*}

Li (2011) BMC Systems Biology

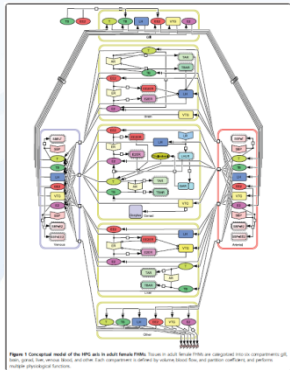
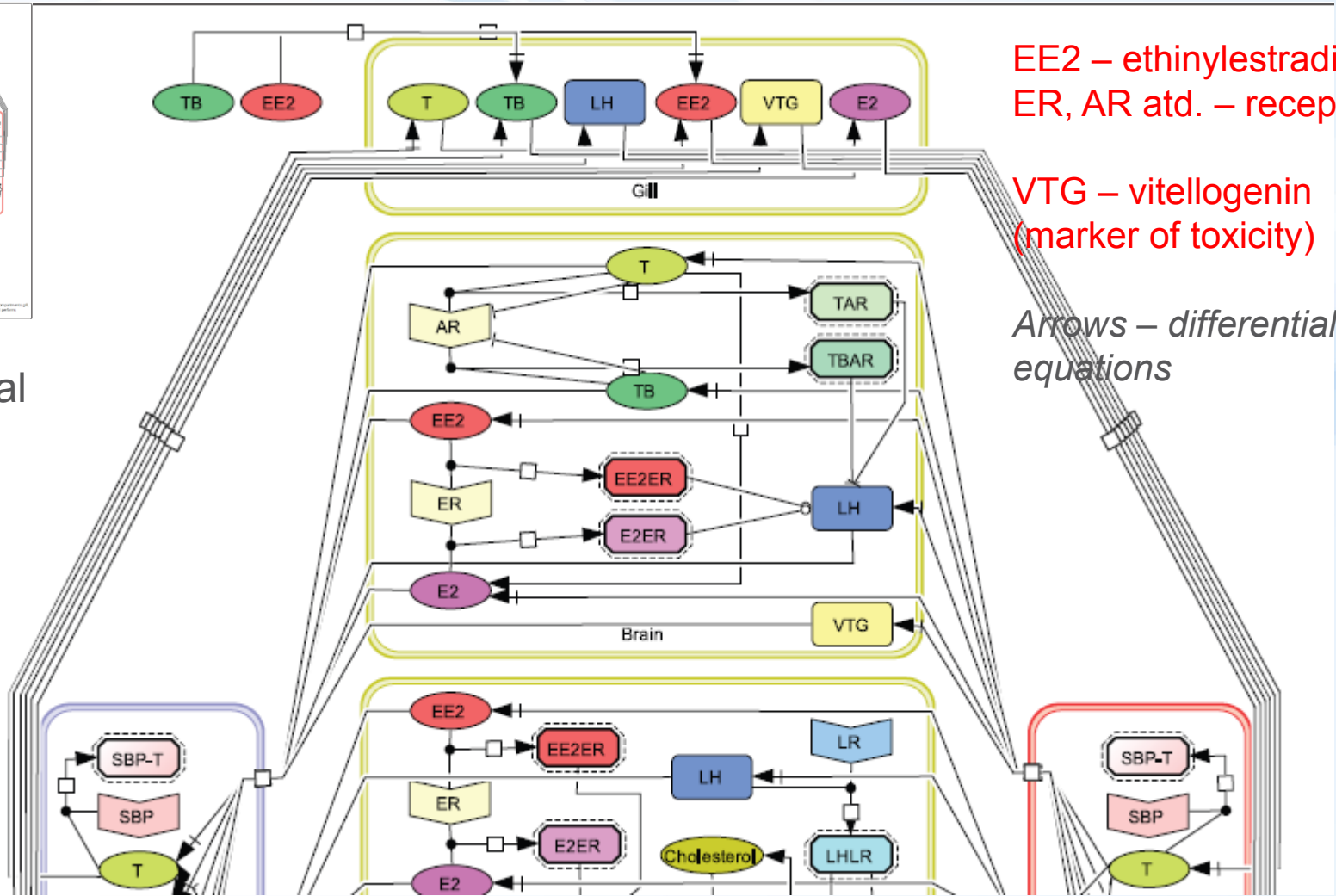


Figure 1 Conceptual model of the HPG axis in adult female F0 fish. The model is a hierarchical flow of information from the hypothalamus through the pituitary gland to the gonads (ovary and testis). The model is represented as a complex network of nodes and arrows.

Conceptual model



EE2 – ethinylestradiol
ER, AR atd. – receptors

VTG – vitellogenin
(marker of toxicity)

Arrows – differential equations

Li (2011) BMC Systems Biology

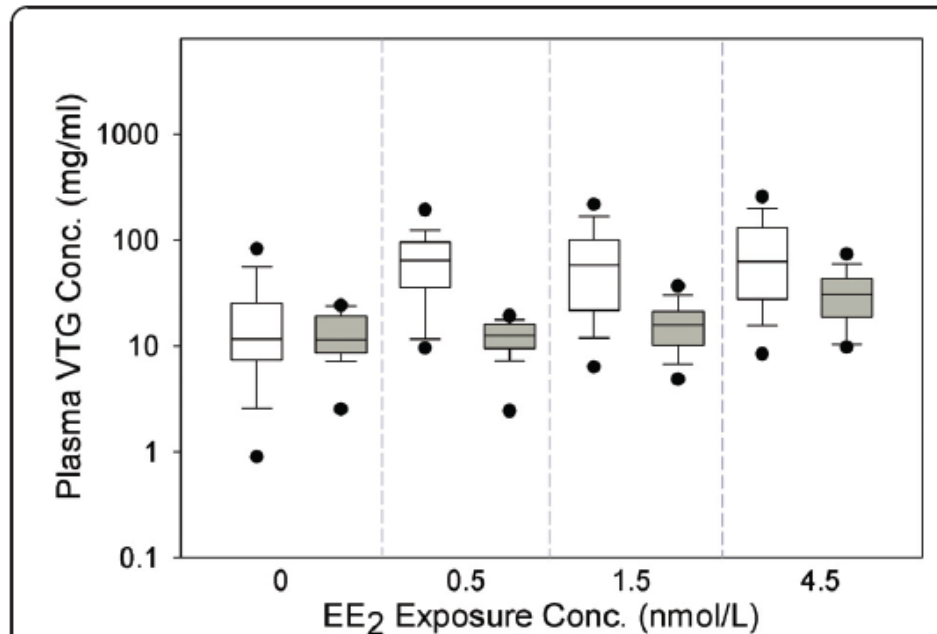


Figure 6 Comparison of model predictions with measured data in female FHM_s exposed to EE₂. $n = 28$ at each sampling time.

White boxes represent model predictions, and grey boxes represent measured data [42]. The x-axis represents EE₂ concentrations in ng/L. The solid line within the box marks the median; the boundary of the box farthest from zero indicates the 75th percentile; the boundary of the box closest to zero indicates the 25th percentile; the whisker (error bar) farthest from zero marks the 90th percentile; whisker (error bar) closest to zero marks the 10th percentile; the circle farthest from zero marks the 95th percentile; and the circle closest to zero marks the 5th percentile.

Results:

MODELLED (white)
Vs
MEASURED (grey)

...good comparable




Update – quantitative mechanistic/computational toxicology

 OPEN ACCESS  PEER-REVIEWED

RESEARCH ARTICLE

A Computational Model of the Rainbow Trout Hypothalamus-Pituitary-Ovary-Liver Axis

Kendall Gillies, Stephen M. Krone, James J. Nagler, Irvin R. Schultz 

Published: April 20, 2016 • <https://doi.org/10.1371/journal.pcbi.1004874>

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Abstract

Reproduction in fishes and other vertebrates represents the timely coordination of many endocrine factors that culminate in the production of mature, viable gametes. In recent years

Subject Areas



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Update – quantitative mechanistic/computational toxicology

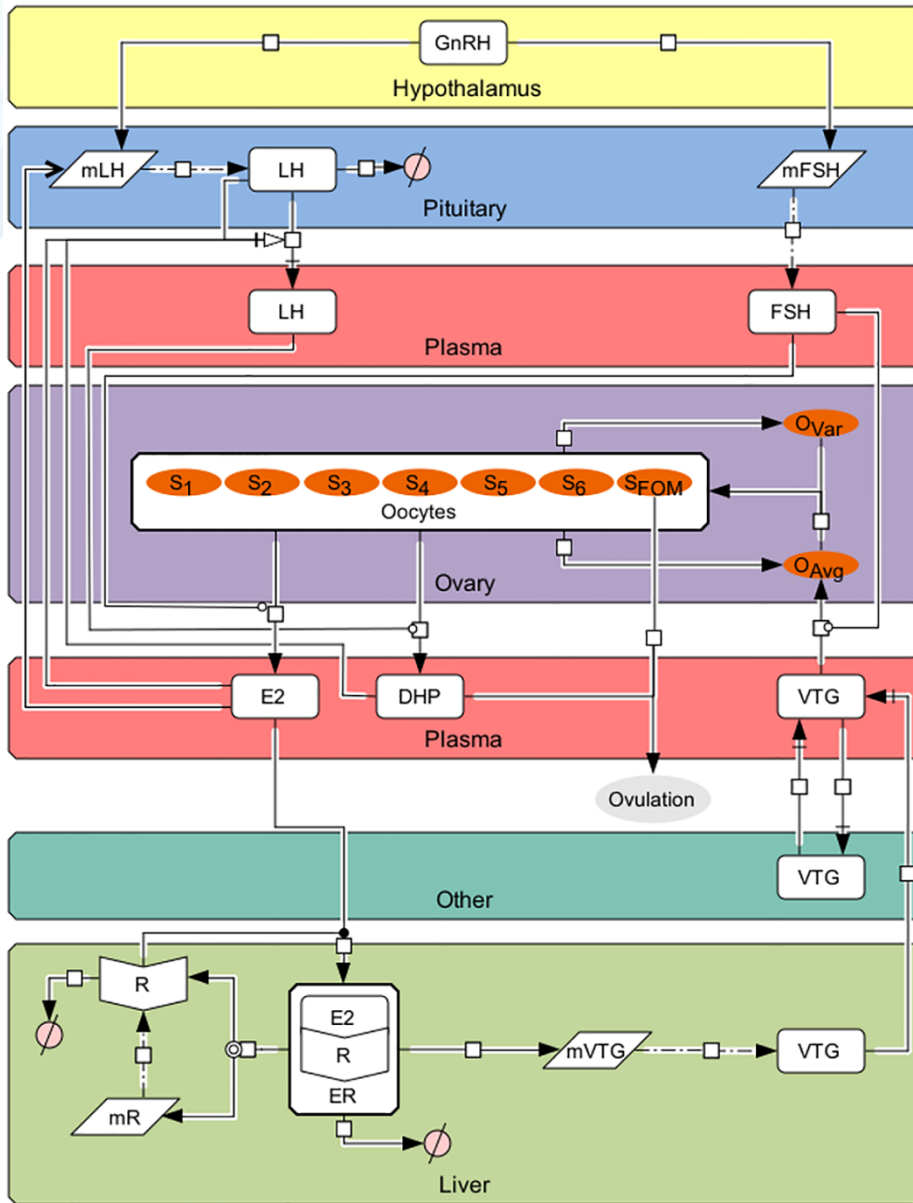
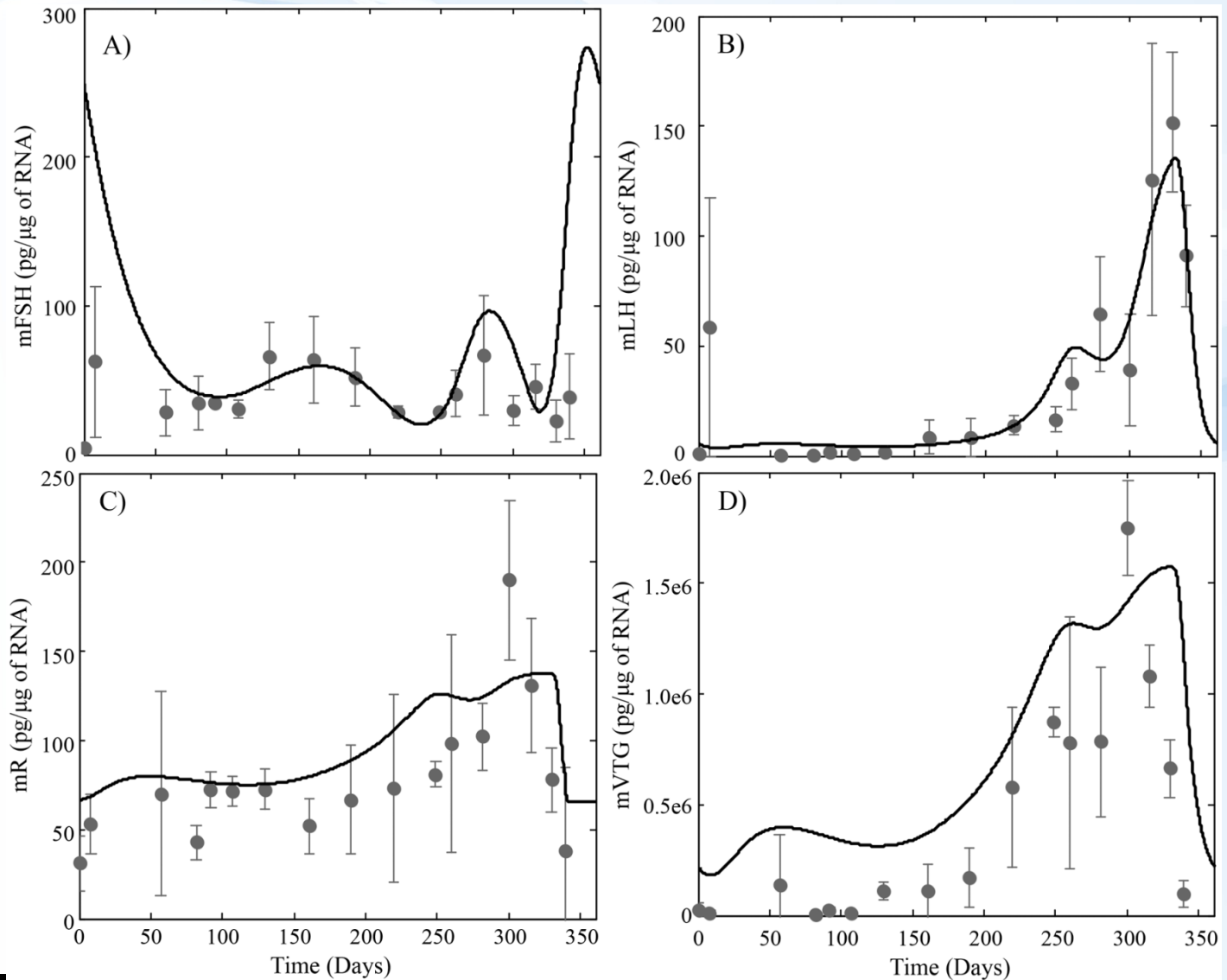


Fig 1. The HPOL signaling network in rainbow trout as formulated in our model.

Arrows and symbols on graph follow CellDesigner vs. 4.4 notation (www.celldesigner.org). GnRH is secreted from the hypothalamus into the pituitary stimulating the production of mFSH and mLH, which then leads to formation of FSH and LH, respectively. FSH, which is being continuously secreted from the pituitary, travels to the ovaries to stimulate production of E2. E2 then travels to the liver to bind with E2 receptors (R; translated from mR) to form ER. ER then stimulates the production of mVTG, which produces VTG_L. Secreted VTG then travels from the liver to the ovaries via the plasma (VTG_P) where it is absorbed by follicles in stages 3 through 6 (the proportion of follicles in these stages are denoted by S_j, j = 3, 4, 5, and 6) during vitellogenesis, the rate of which is affected by FSH_P, to promote oocyte growth (O_{Avg}). Oocyte growth then progresses the oocytes through the stages using a Weibull distribution created from O_{Avg} together with O_{Var}. In the later stages LH_P stimulates the oocytes to produce DHP. Finally, oocytes undergo final maturation (S_{FOM}) and combined with DHP, determine when the fish ovulates.

Update – quantitative mechanistic/computational toxicology

Fig 3. HPOL **model predictions** for (A) pituitary levels of FSH $_{\beta}$ subunit mRNA, (B) pituitary levels of LH $_{\beta}$ subunit mRNA, (C) Hepatic levels of E2 receptor mRNA and (D) Hepatic levels of VTG mRNA
Observed data (dark grey circles; mean TG mRn = 3)



Closing remarks

- Ecotoxicology is exciting **science!**
- **Interface**: science and society
- Many **opportunities**
- Sometimes **hard work**
10% inspiration and 90% „perspiration“



- Be **creative**: move frontiers
- **Keep the purpose** in mind
- **Be critical**: do not accept perceptions as facts
- **Do not hesitate to speak up** ..