

# Quantitative Weight of Evidence Model for Assessing Adverse Outcome Pathways

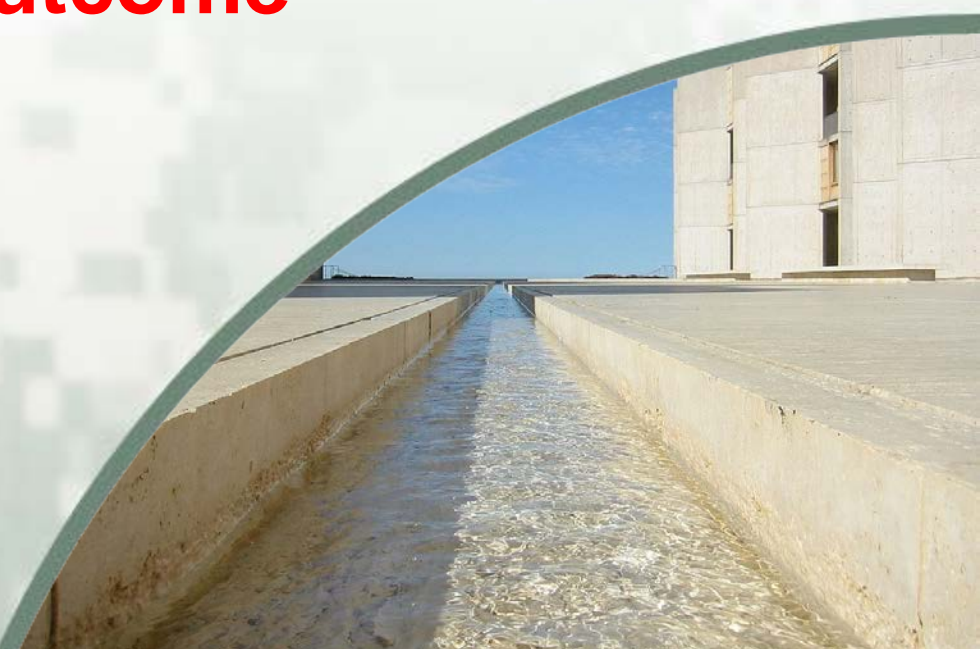
Dasha Agoulnik<sup>1,3</sup>, Olivia Massey<sup>2,3</sup>, Christy Foran<sup>3</sup>, Dan Villaneuve<sup>4</sup>, Gary Ankley<sup>4</sup>, Natalia Garcia-Reyero Vinas<sup>3</sup>, Ed Perkins<sup>3</sup>, and Igor Linkov<sup>3</sup>

<sup>1</sup>University of Massachusetts Amherst,  
<sup>2</sup>Massachusetts Institute of Technology,  
<sup>3</sup>US Army Engineer Research and  
Development Center, <sup>4</sup>US Environmental  
Protection Agency

[Igor.Linkov@usace.army.mil](mailto:Igor.Linkov@usace.army.mil)



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# Summary

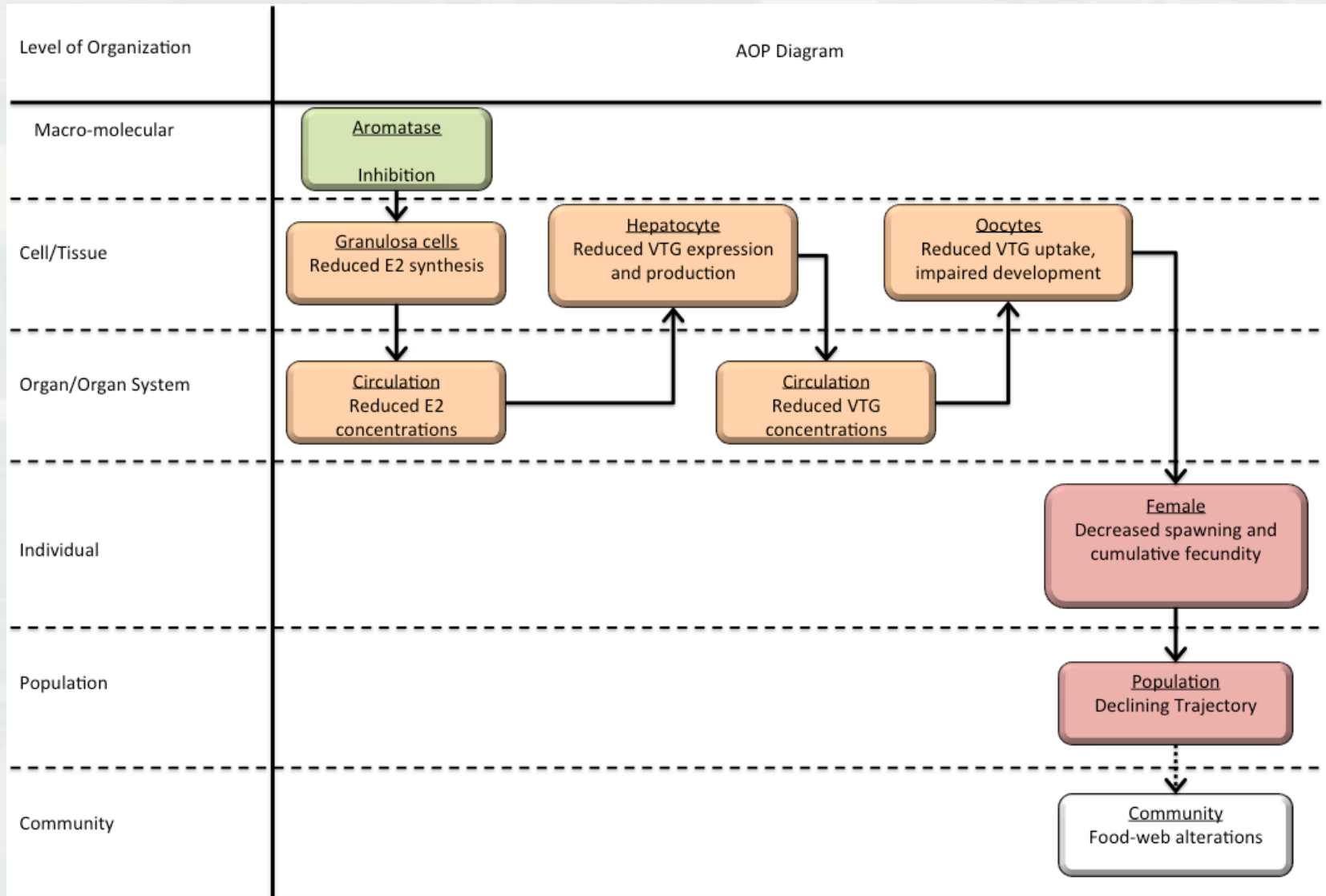
- Majority of WoE frameworks have been qualitative in nature.
- We propose a quantitative WoE framework that utilizes MCDA methodology for integrating evidence underpinning KER within an AOP.
- We developed a prototype model that was parameterized by input from a few mode of action studies and judgments of individual experts.



# Comparing WoE Approaches



# Case Study: Aromatase inhibition leading to fish reproductive dysfunction



# Logic Model Output

Key Event (upstream)	Key Event (downstream)	Weight-of-evidence for link
Aromatase inhibition	Ovarian (granulosa cell) E2 synthesis (reduction)	Strong
Ovarian (granulosa cell) E2 synthesis (reduction)	Plasma 17 $\beta$ -estradiol concentrations (reduction)	Strong
Plasma 17 $\beta$ -estradiol concentrations (reduction)	Transcription and translation of vitellogenin (reduction)	Strong
Transcription and translation of vitellogenin (reduction)	Plasma vitellogenin concentrations (reduction)	Strong
Plasma vitellogenin concentrations (reduction)	Vitellogenin uptake, impaired oocyte development (reduction)	Moderate
Vitellogenin uptake, impaired oocyte development (reduction)	Spawning and cumulative fecundity (reduction)	Moderate
Spawning and cumulative fecundity (reduction)	Population trajectory (declining)	Moderate

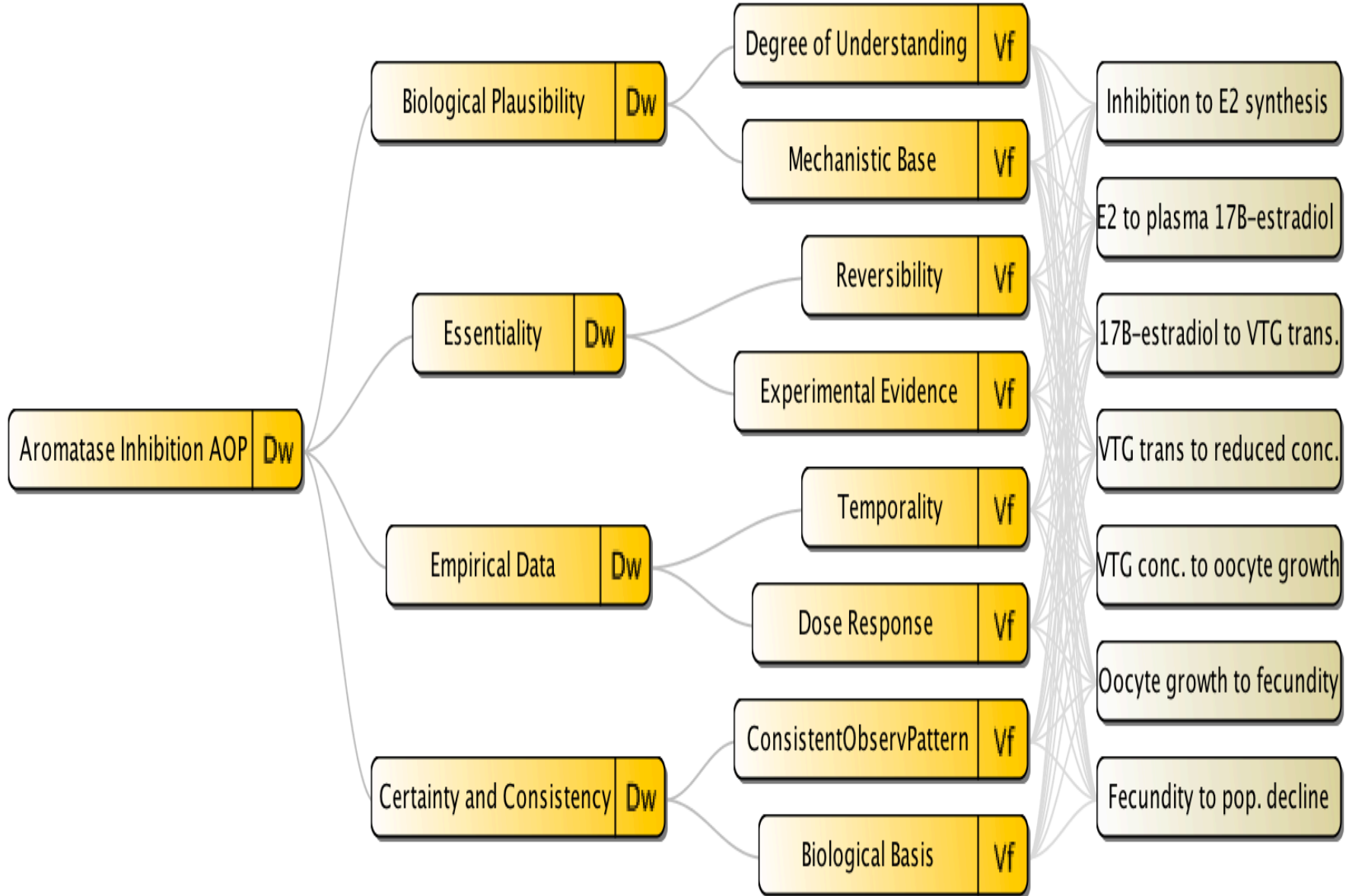


# Steps of MCDA Methodology

1. Define KERs as alternatives for which relative confidence is unknown and needs to be prioritized
2. Map out the criteria and metrics based on BH considerations and KERs as a value tree
3. Assign weights for importance of each of the criteria and metrics associated with each criterion.
4. Score each KER based on each metric (In this case, Strong=3, Moderate=2, Weak=1)
5. Integrate scores and weights for each KER to assess overall confidence level
6. Conduct sensitivity analysis



# MCDA Model



# Criteria Weights and Key Event Scores

Direct weighting - Empirical Data

Set weights directly

Criterion	Weight	
Temporality	0.500	0.500
Dose Response	0.500	0.500

Direct weighting - Aromatase Inhibition AOP

Set weights directly

Criterion	Weight	
Biological Plausibility	0.400	0.400
Essentiality	0.300	0.300
Empirical Data	0.200	0.200
Certainty and Consistency	0.100	0.100

Scores/Performances - Degree of Understanding

Scores Value function

Set scores directly

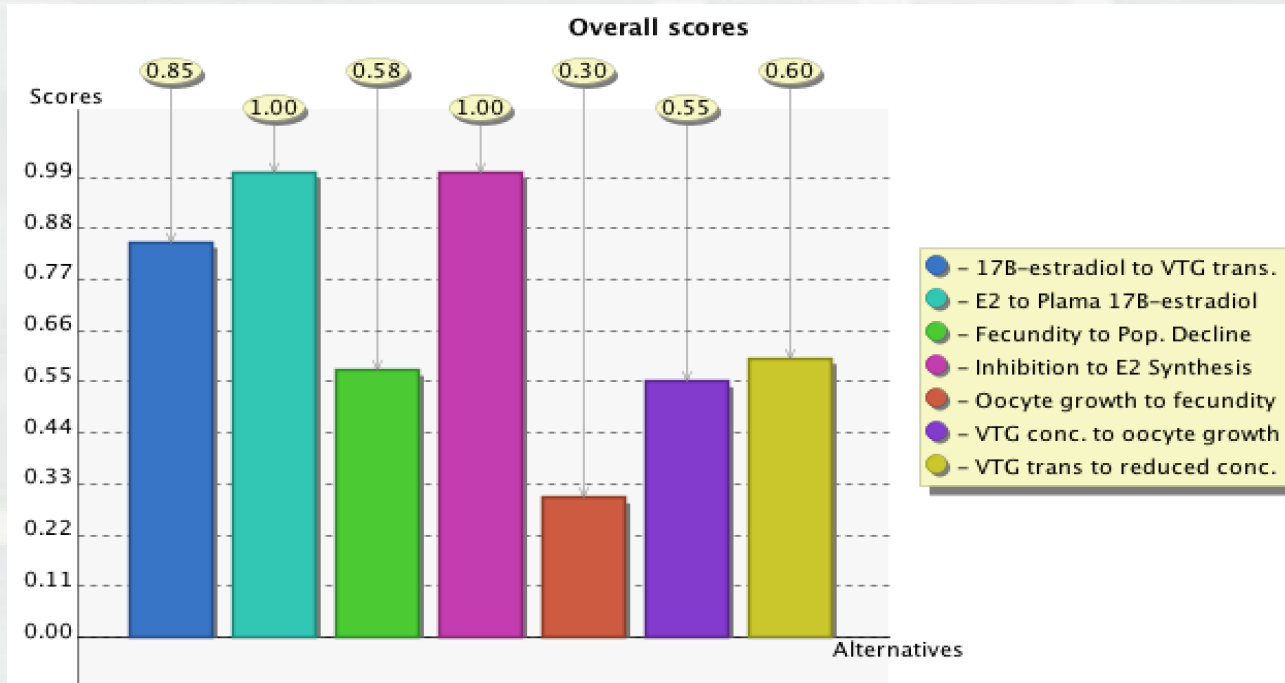
Alternatives performances and scores

Alternative	Current	Score
Inhibition to E2 Synthesis	3.000	1.000
E2 to Plasma 17B-estradiol	3.000	1.000
17B-estradiol to VTG trans.	3.000	1.000
VTG trans to reduced conc.	3.000	1.000
VTG conc. to oocyte growth	3.000	1.000
Oocyte growth to fecundity	2.000	0.000
Fecundity to Pop. Decline	3.000	1.000

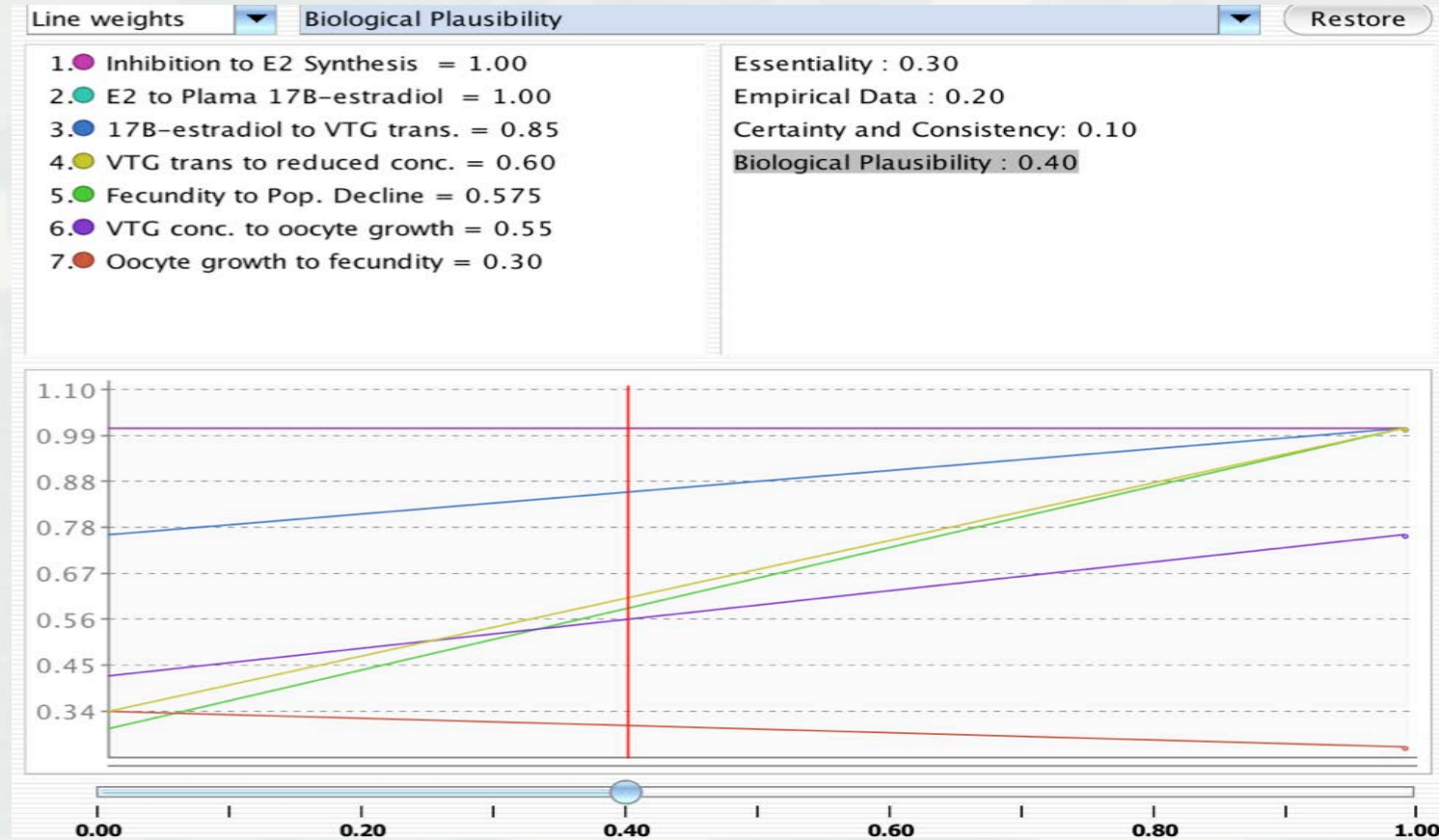




# Confidence Assessment Scores



# Sensitivity Analysis



# Conclusions

- Quantitative WoE frameworks provide an objective and transparent mean to assess AOPs
- MCDA strengthens WoE logic by adding visual effect of a mapped decision structure as well as quantitative weighing of LOE
- Restricts expert inputs to weighing evidence
- Allows for incorporation of inputs from multiple experts



# References

- Linkov, I., Welle, P., Loney, D., Tkachuk, A., Canis, L., Kim, J., Bridges, T. (2011). The use of Multi-Criteria Decision Analysis Methods to Support Weight of Evidence Evaluation in Risk Assessment. *Risk Analysis* 31:1211-1225.
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