

A Unified Aquatic Life Framework for Addressing the Affected Percentages of Individuals, Species, and Time

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Two Approaches for Addressing Time Variability

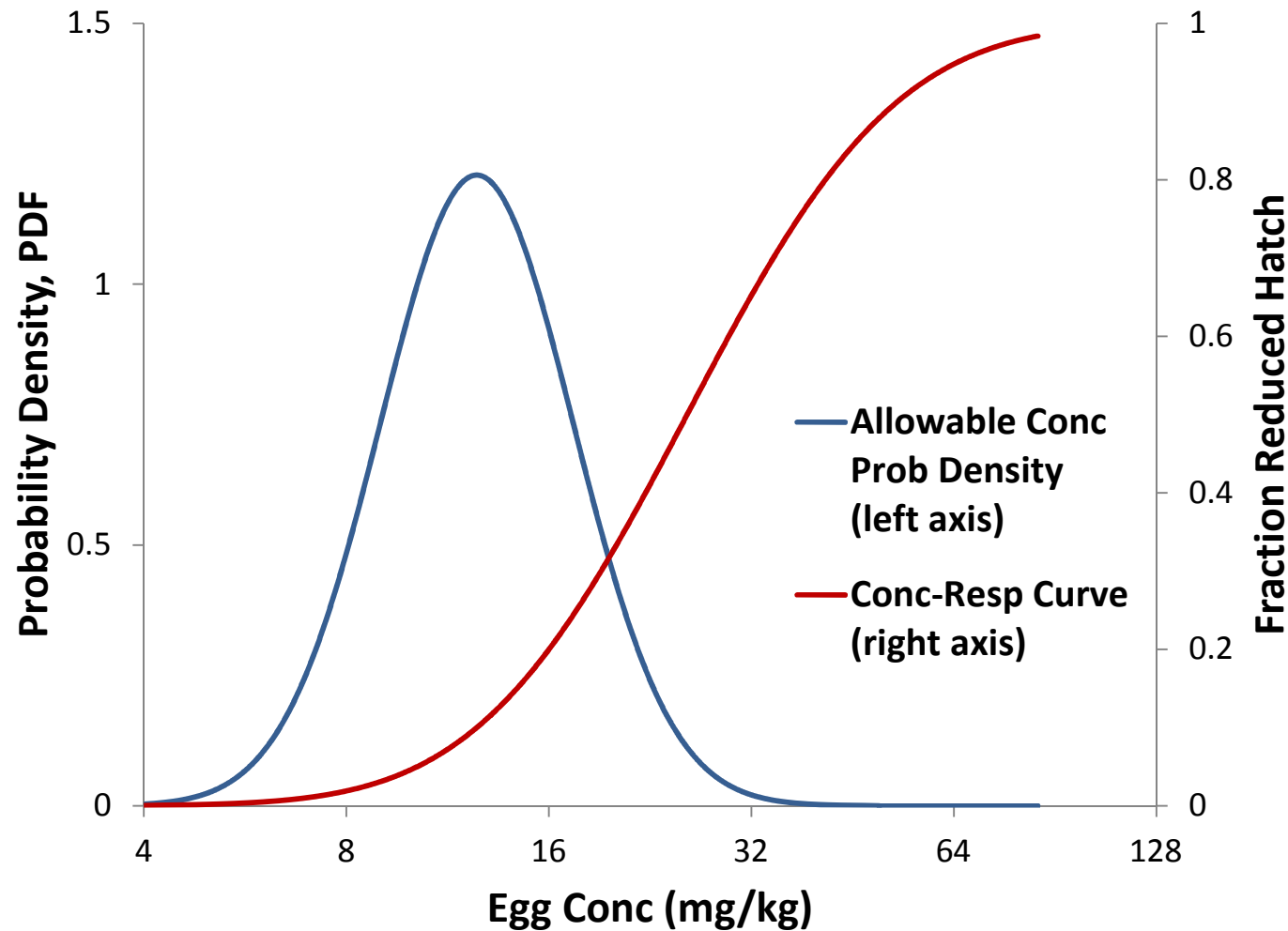
- Simple approach uses distribution of exposure concentrations.
- Complex approach uses a long time series of concentrations.

An Actual Application Applying the Simple Distribution of Concentrations

- State of Utah adopted a selenium criterion for the Great Salt Lake, Gilbert Bay.
 - Applies to the Se concentration in bird eggs.
 - Set at 12.5 mg/kg, the State's estimate of the EC10 for mallard duck, the most sensitive known species.
 - Applies as the geometric mean concentration.
- Question: What is the aggregate level of effect if the water body geometric mean rises to the EC10 and the variability of concentrations (expressed as a CV or log standard deviation) remains as present?

Combining Ambient Concentration Distribution with a Species Concentration-Response Curve:

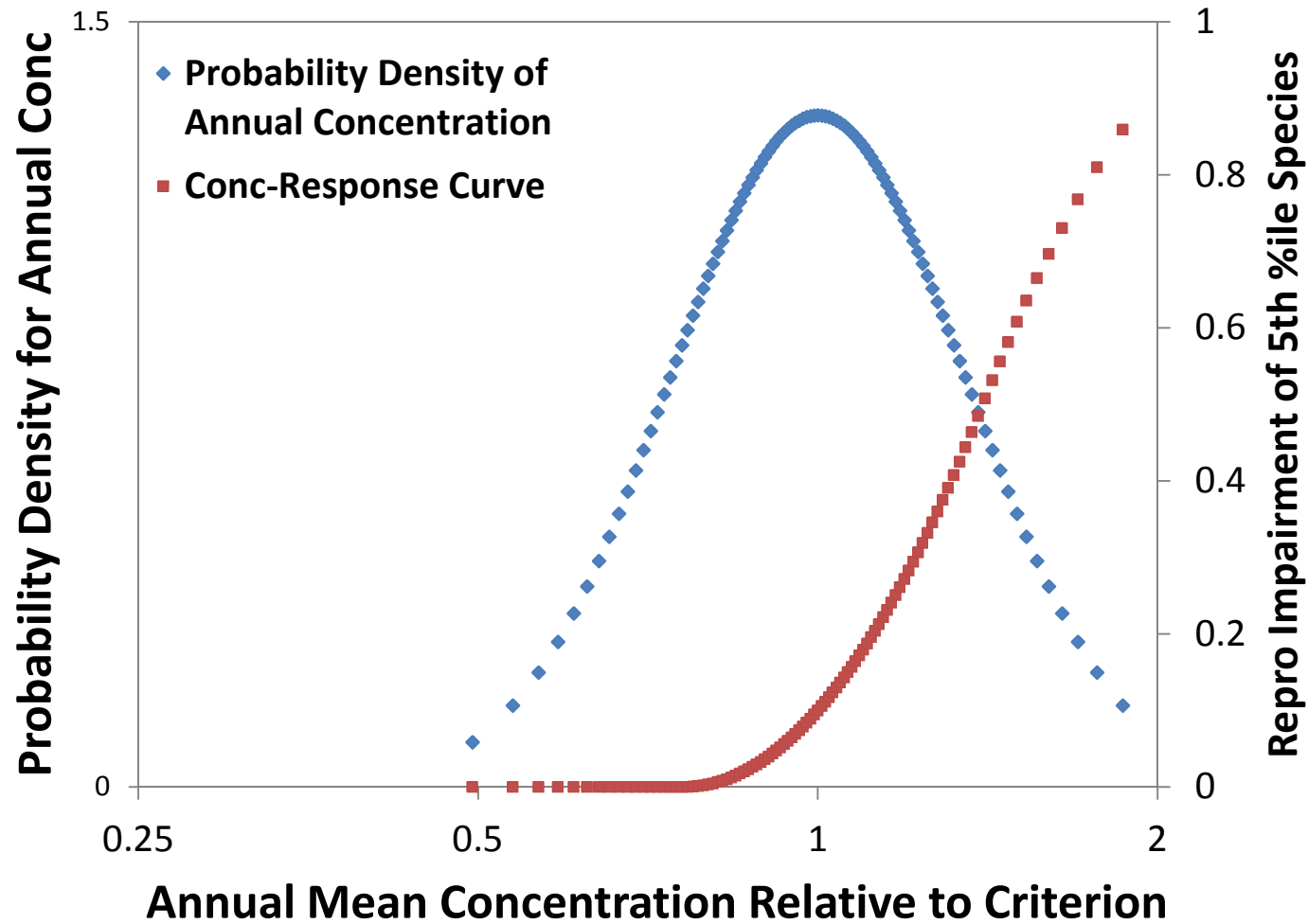
$$\text{Aggregate Effect} = \sum \text{probability}_i \times \text{Effect}_i$$



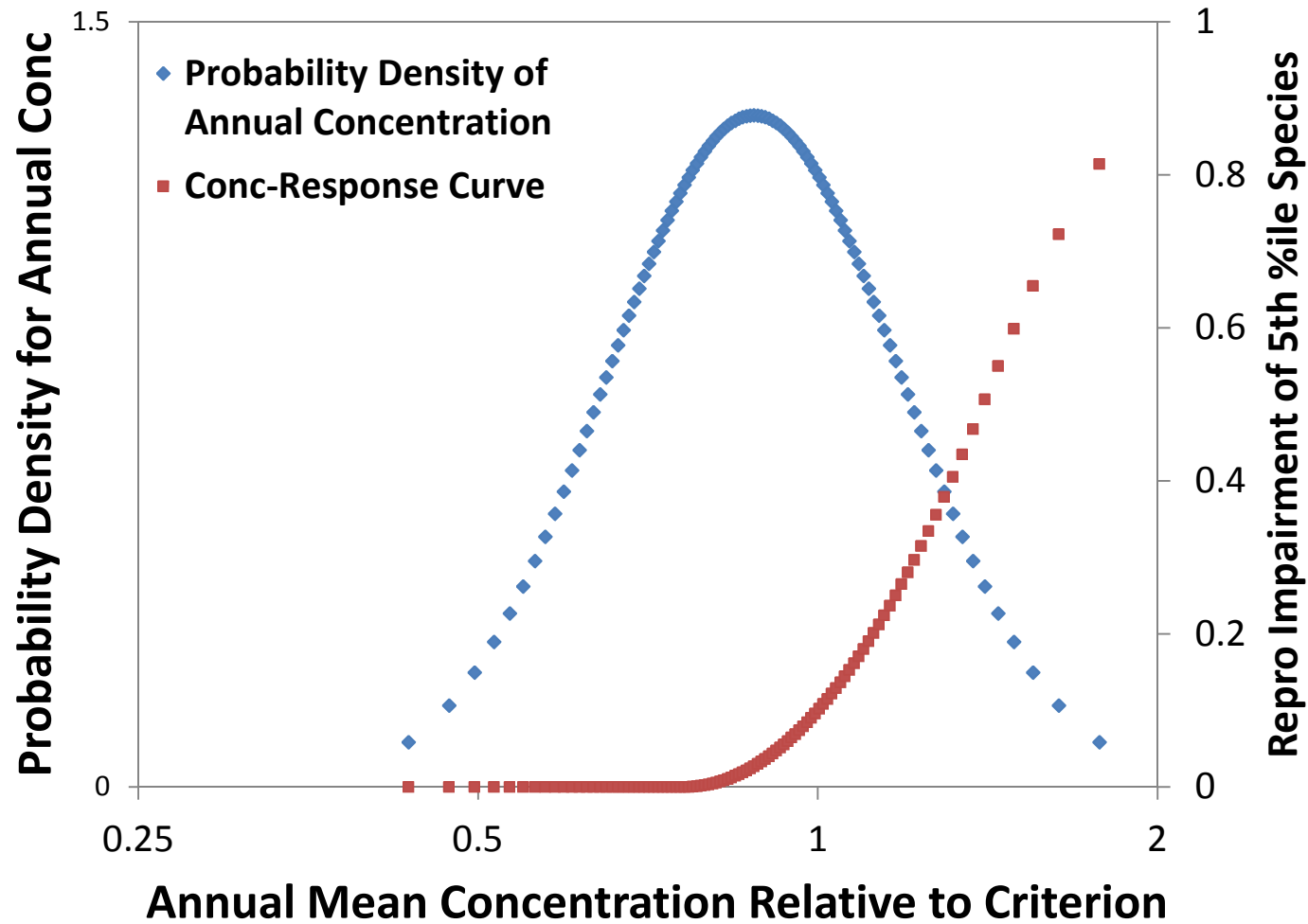
Hypothetical Illustration: Selecting a Return Interval for Exceeding a Criterion

- Possible application: let's say a state wants to allow the annual reproductive season mean Se fish tissue concentration to exceed the criterion only once in "X" number of years.
- We ask: for various values of the return frequency, X, what is the level of aggregate effect on the hypothetical 5th percentile species having EC10=Criterion?

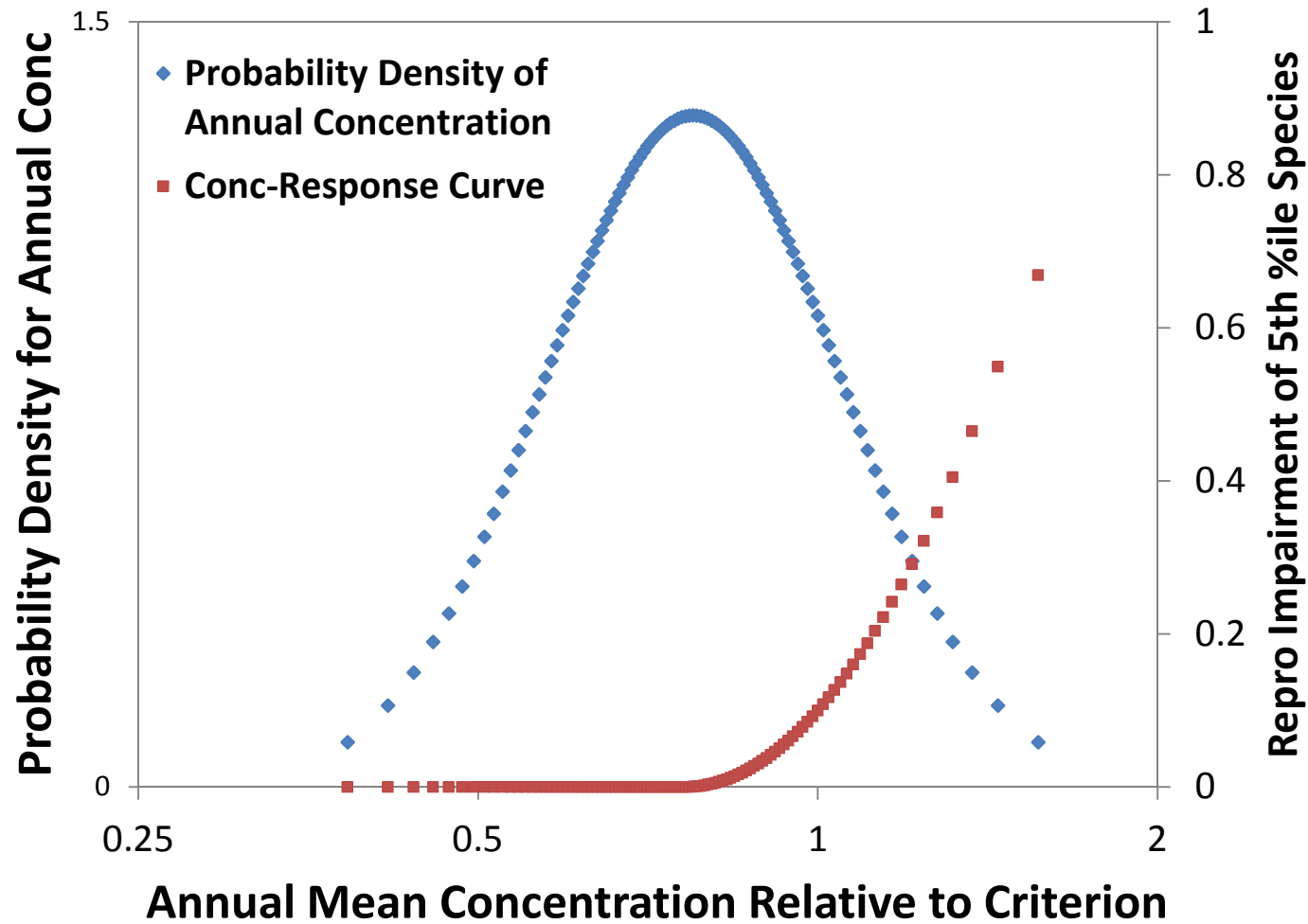
Trial 1: Once in 2 Years



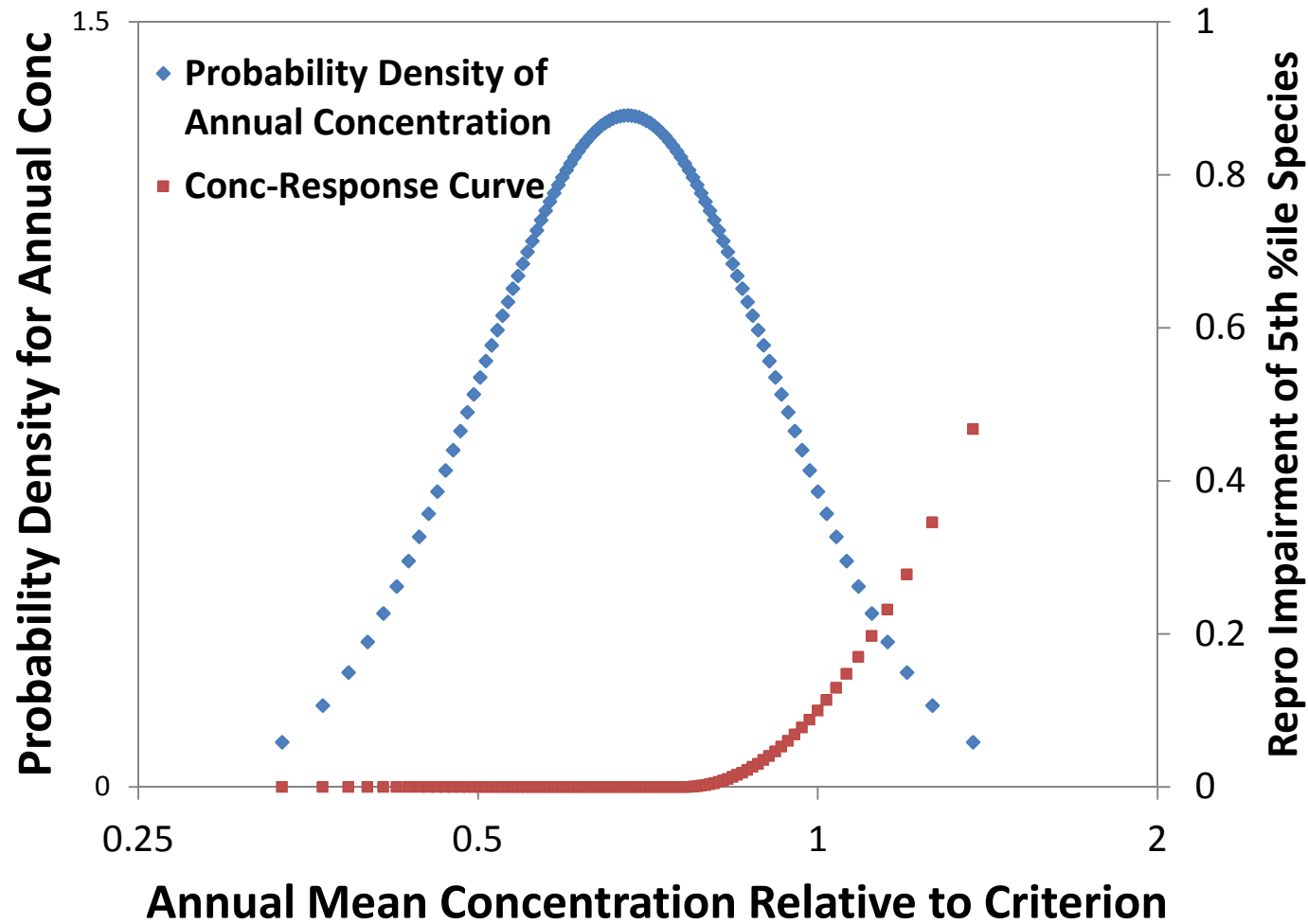
Trial 2: Once in 3 Years



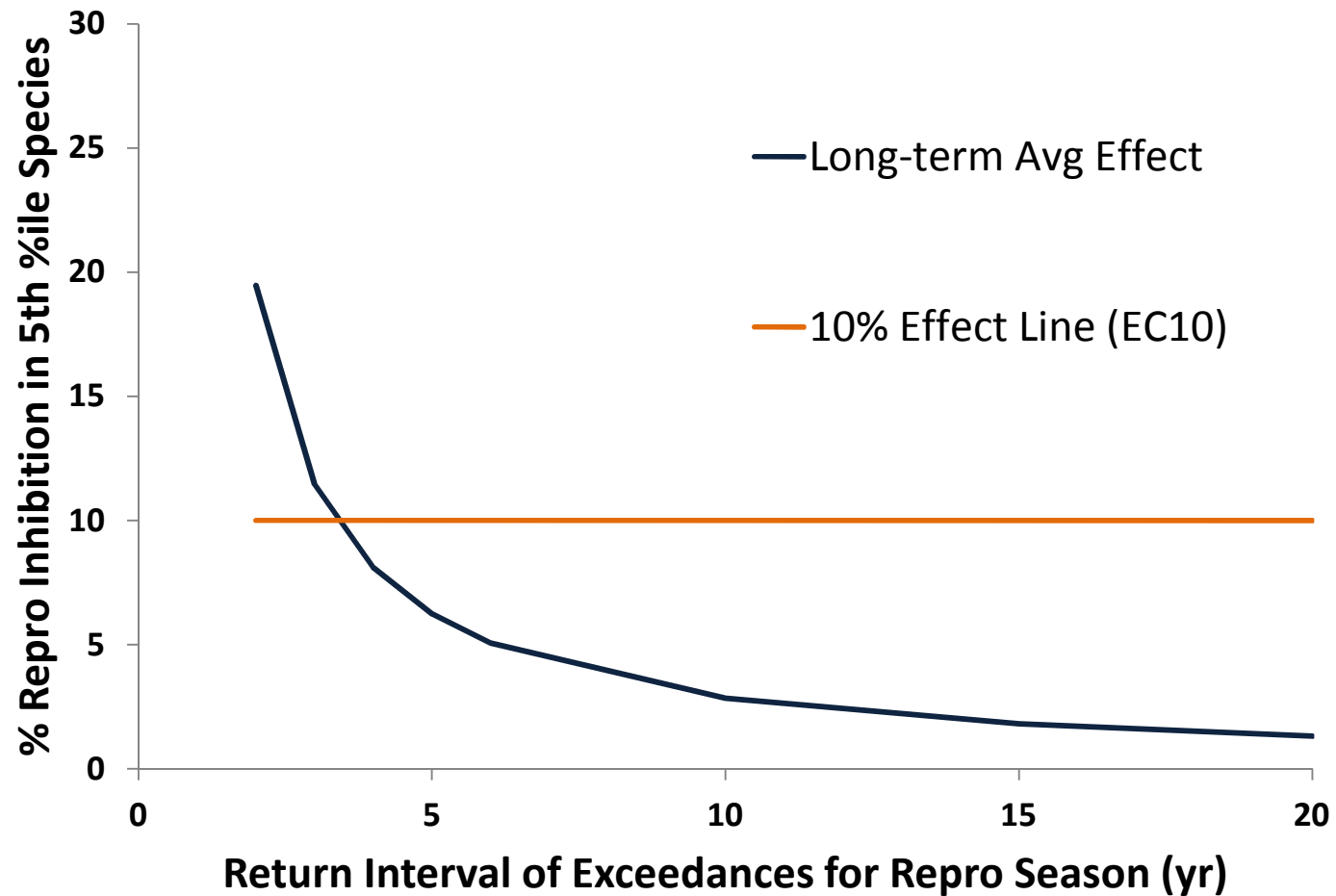
Trial 3: Once in 5 Years



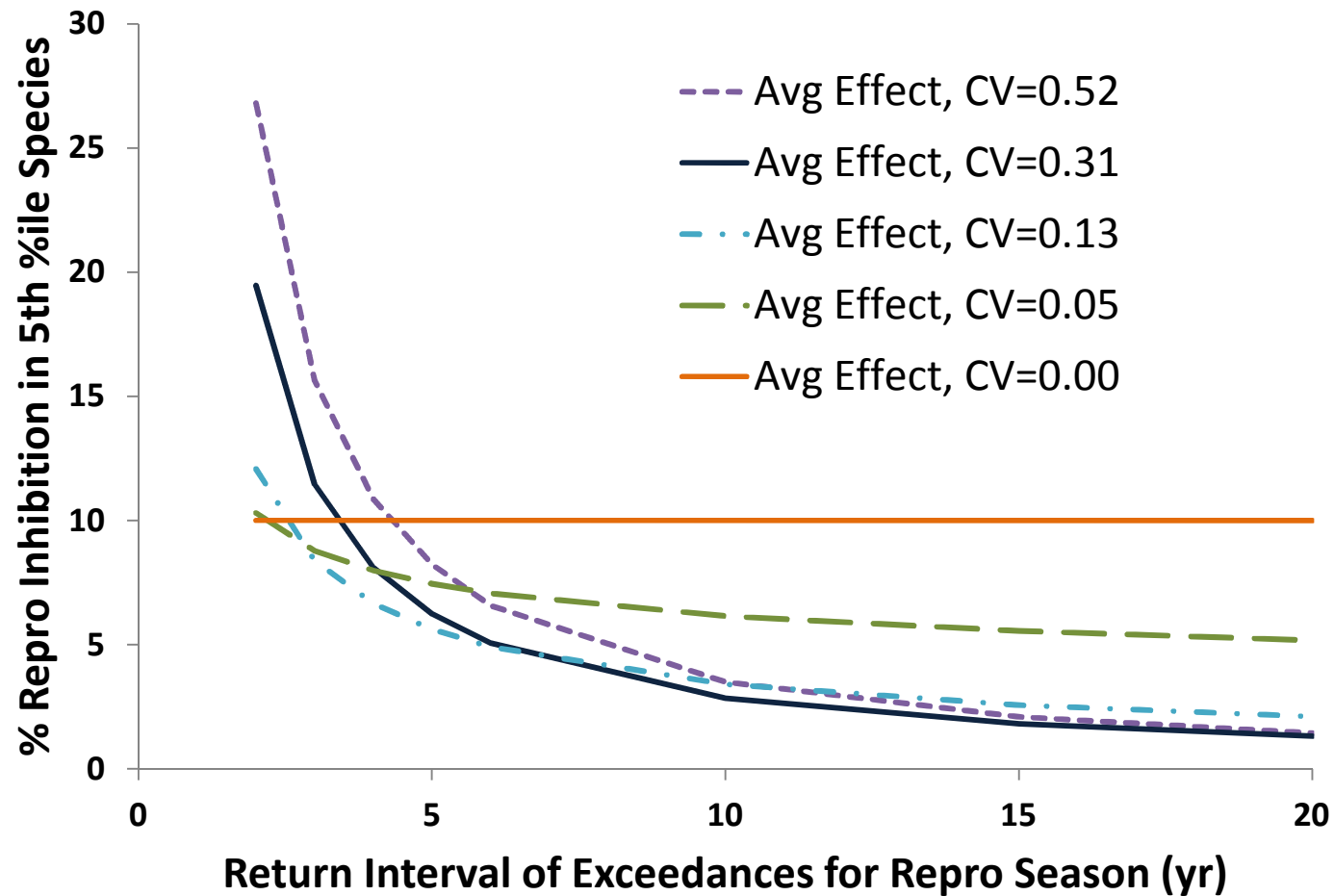
Trial 4: Once in 10 Years



Results for Various Exceedance Frequencies



Influence of Annual Concentration Variability



Complex Approach

- The next set of slides addresses the complex approach, which uses a time series of exposures rather than the statistical distribution of exposures.

Higher Tier Assessment: Combined Application of Kinetic Toxicity Model and Population Model

- For Each Species, Apply Two Models -

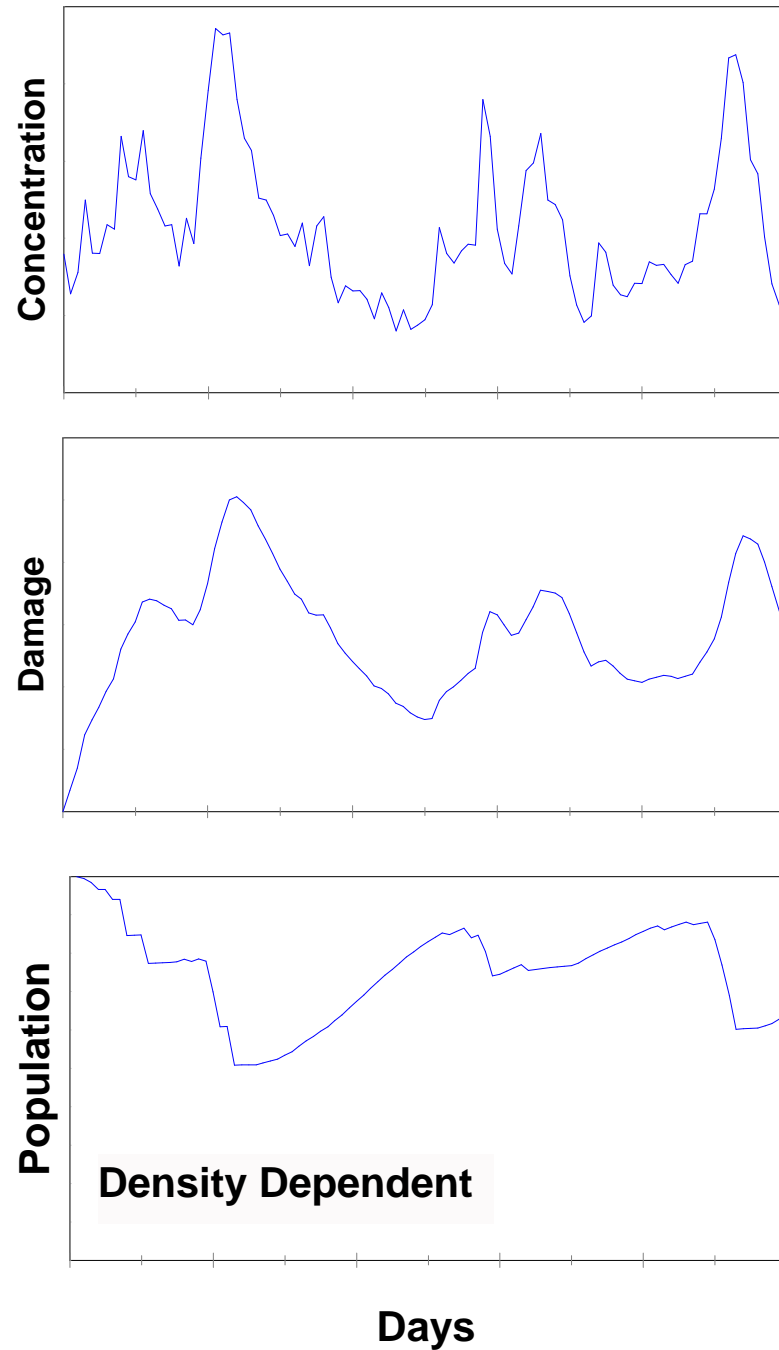
- Kinetic toxicity model to translate from lab test exposures to continuously variable concentrations.
- Life-stage structured population model.

Higher Tier Assessments: Combined Application of Kinetic Toxicity Model and Population Model

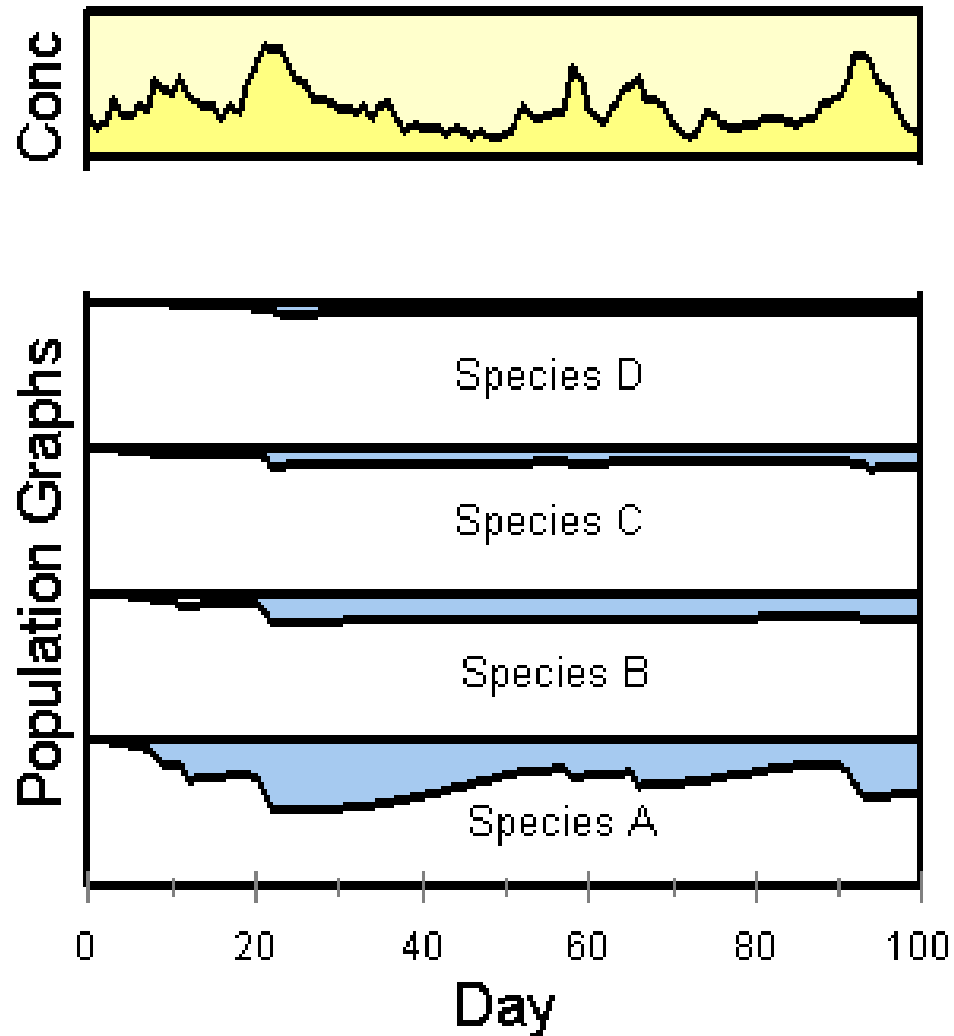
- For Each Species, Apply Two Models -

- Kinetic toxicity model to translate from lab test exposures to continuously variable concentrations.
- Life-stage structured population model.
- The combination allows discerning:
 - How sensitivity differs among individuals and between life stages.
 - How reductions in survival and reproduction differ in their effect on populations.
 - How population effects differ between species that recover rapidly and species that recover slowly.

- Toxicant concentration, short example
- Accumulation of stress in individuals of one species
- Population response in one species



Generating an Assemblage Toxicity Index for Tested Representative Species



Kinetic Toxicity Model

- Needed to predict toxicity of continuously variable concentrations
- Provides input (such as death rates) to population model.

Minimum Data Needed to Calibrate Toxicity Model for Each Animal Species

- Required

- Acute LC50
- Chronic survival EC50 or EC20
- Chronic EC50/EC20 ratio (conc-response slope)

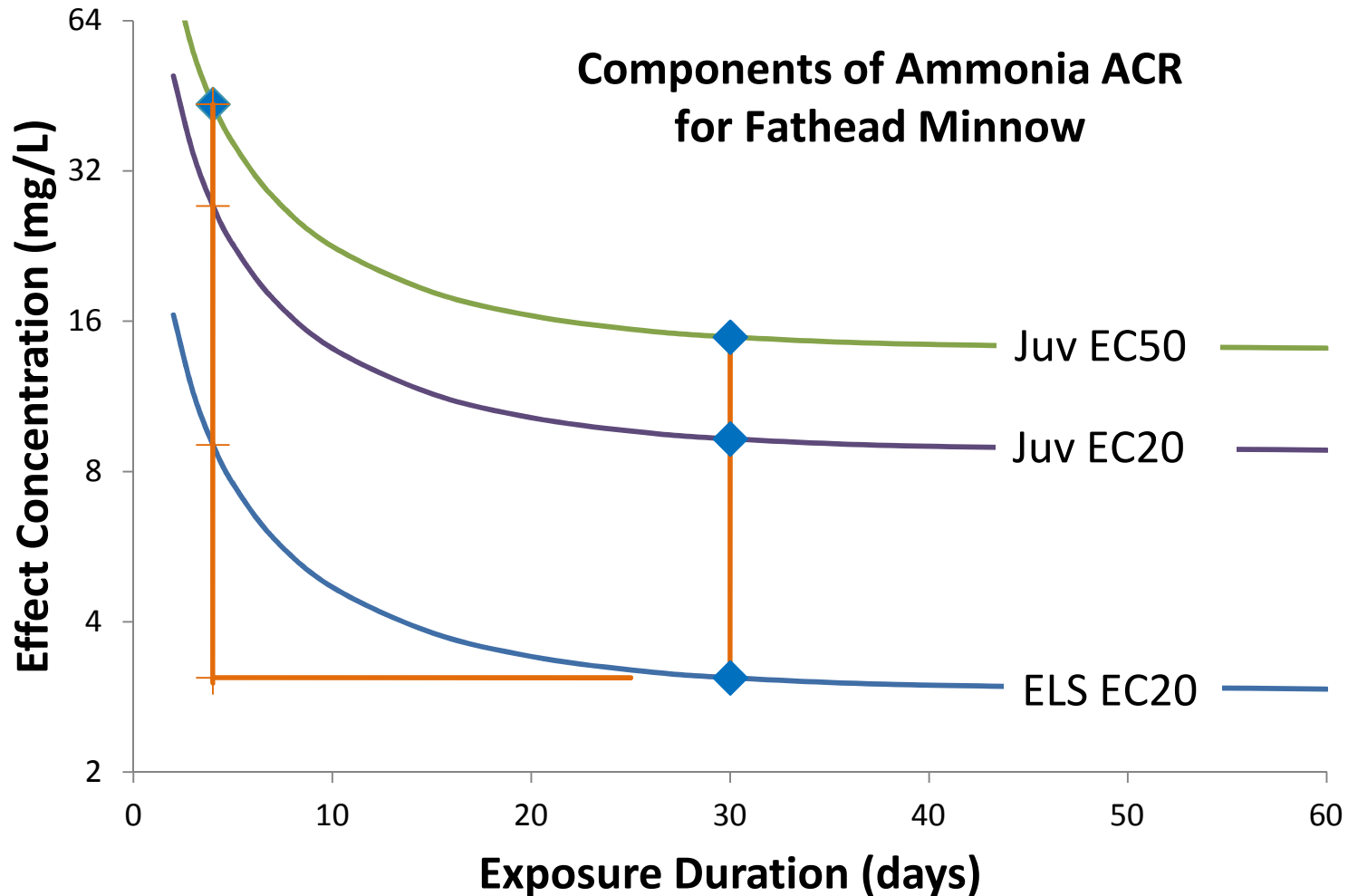
- Desirable

- Chronic ECx differences between early life stages and juvenile-adult stages

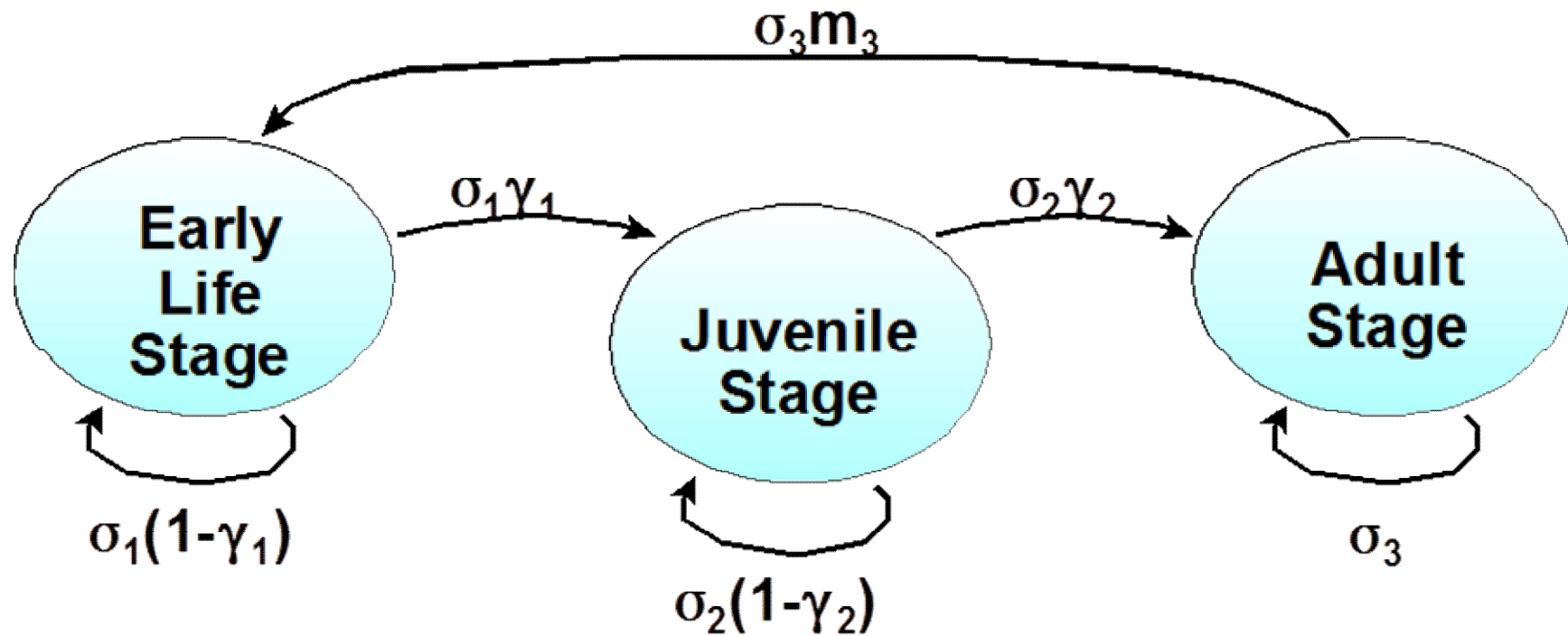
- Optional

- Chronic ECx differences between lethal and sublethal effects (growth-reproduction)

Simple Use of the Kinetic Model to Help Understand Acute-Chronic Ratios



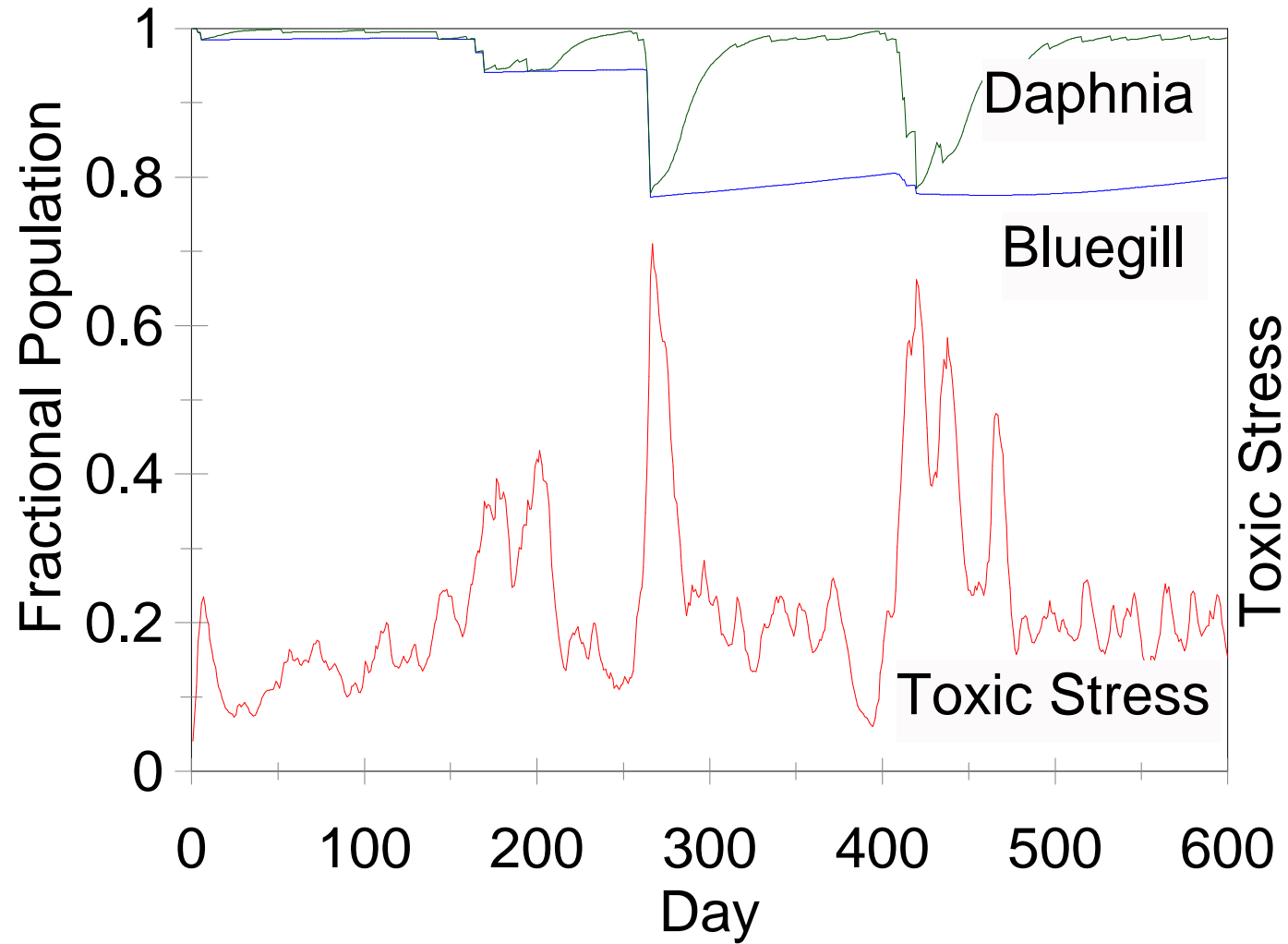
Stage-Structured Population Model



Population Model Input Parameters

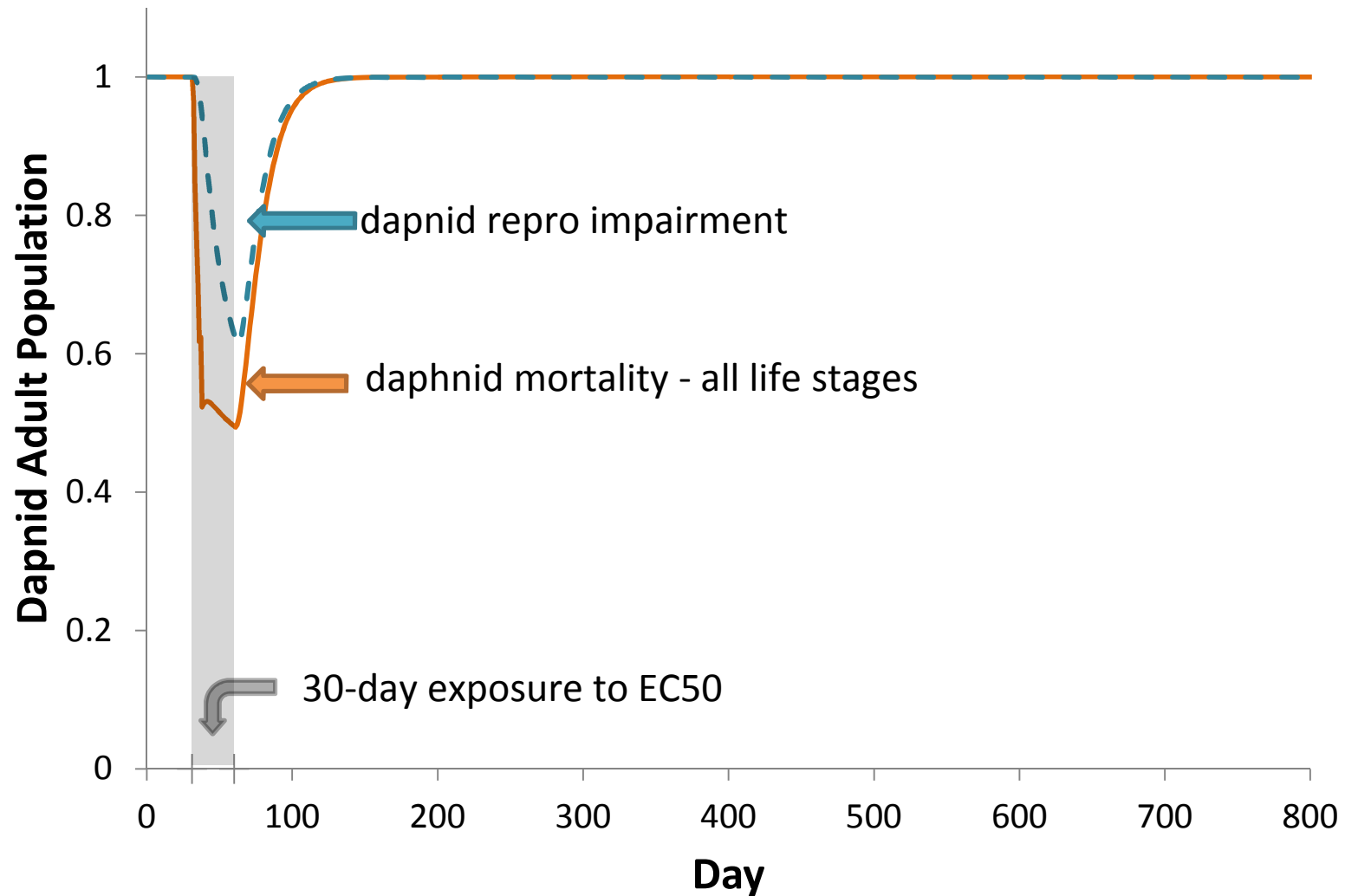
- Decide how many life stages you want to divide the species lifespan into.
- For each life stage, specify its:
 - Duration
 - Background survival rate
 - Reproductive rate – for adult stage(s) only.

Modeling Effects on Populations



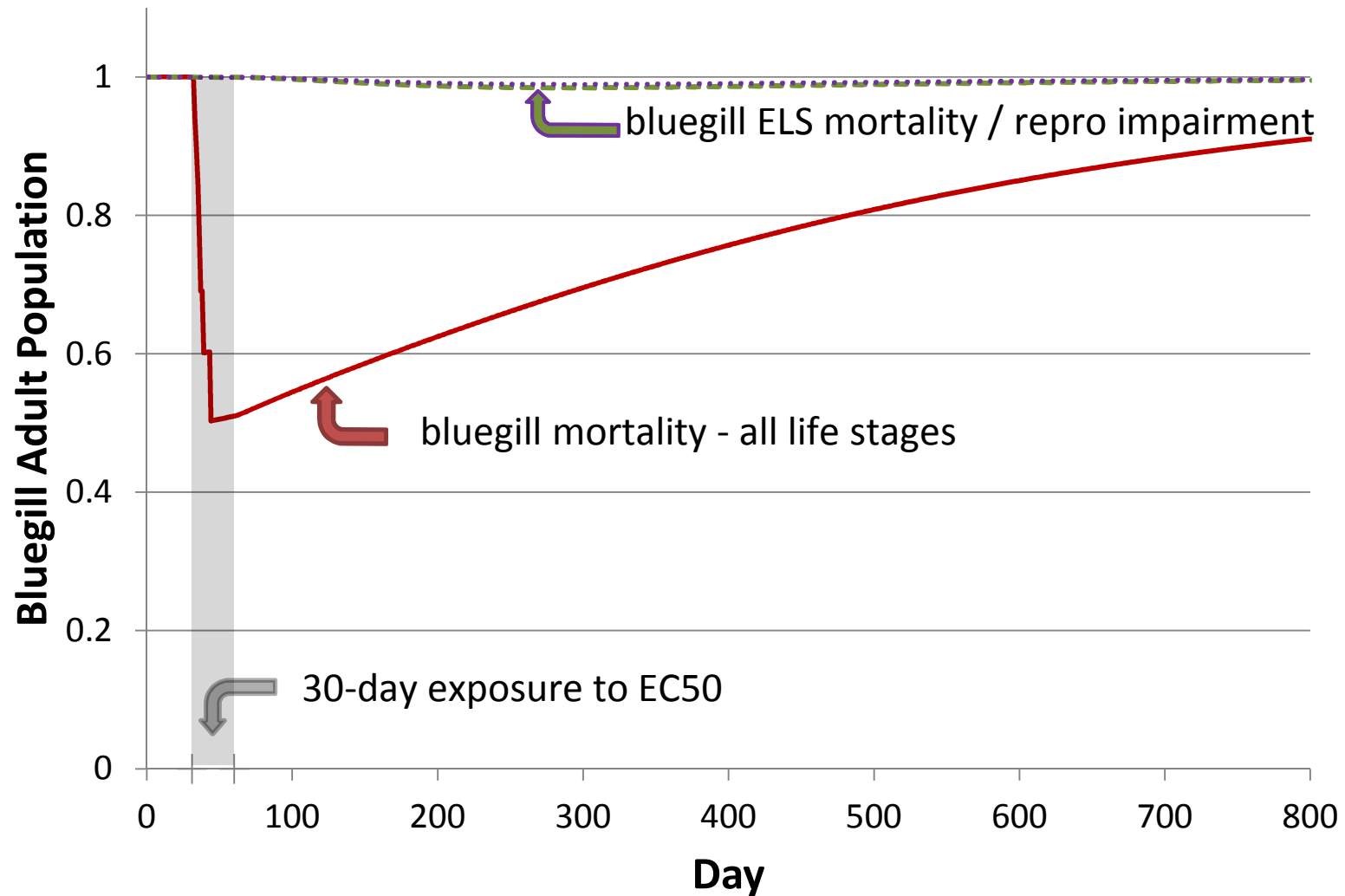
Mortality v. Repto Effects

Daphnia response to 30-day Pulse Exposure at EC50

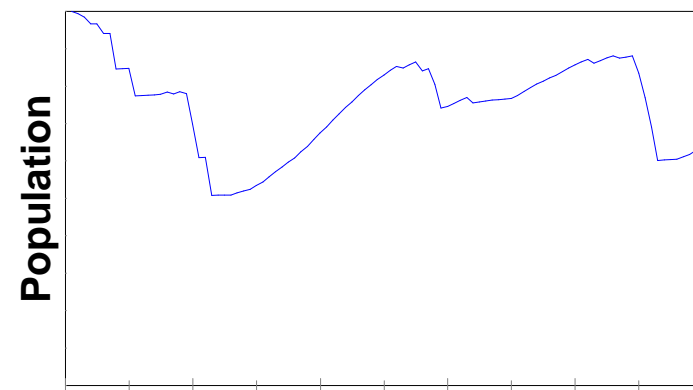
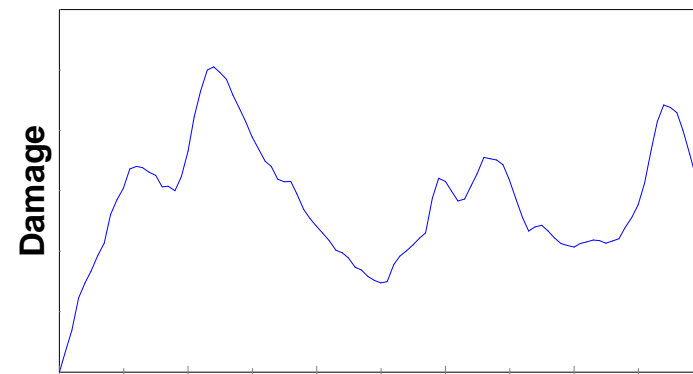
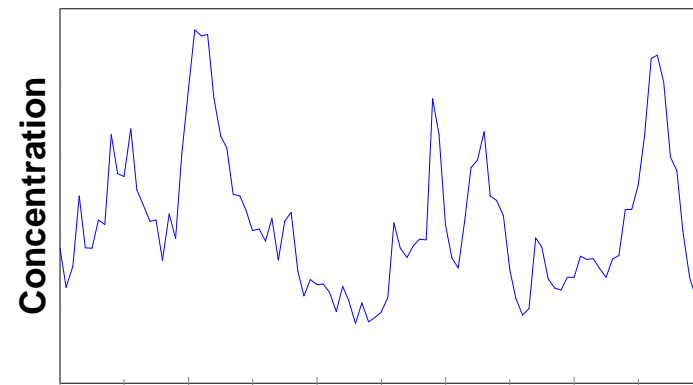
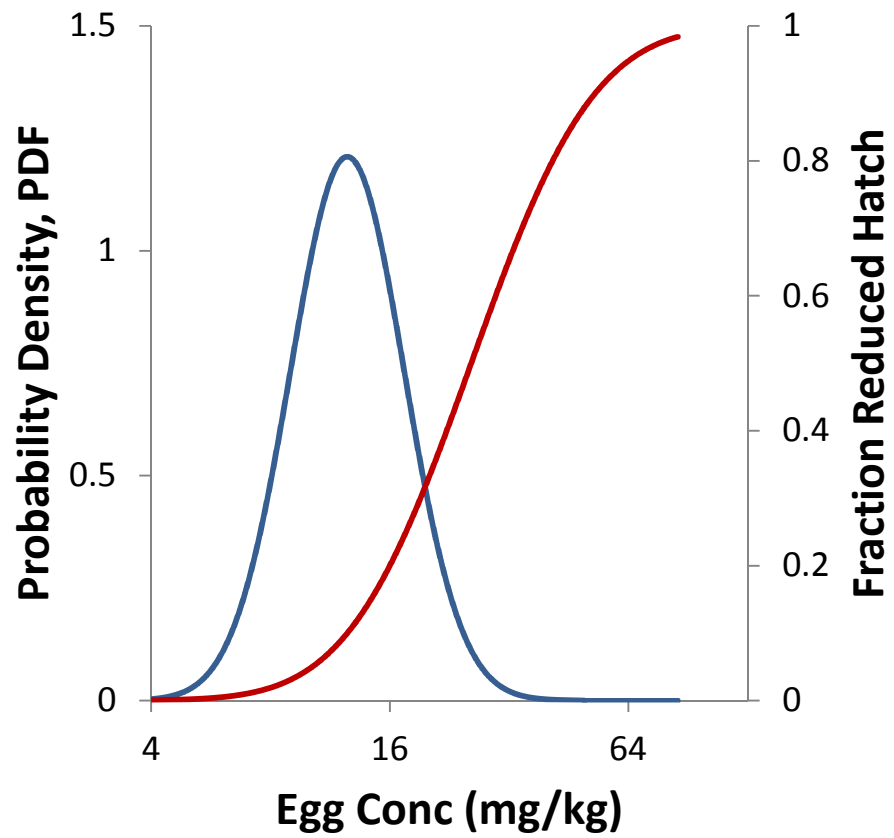


Mortality v. Repro Effects

Bluegill response to 30-day Pulse Exposure at EC50



Comparing the Simple and Complex Approaches:
Coupled Concentration Distribution & Response Curve
vs.
Coupled Toxicity Model & Population Model



Days

Comparing the Simple and Complex Approaches:
Coupled Concentration Distribution & Response Curve
vs.
Coupled Toxicity Model & Population Model

- Simple approach:
 - Bypasses kinetics of toxicity.
 - Bypasses sequencing of events.
 - Cannot discern life-stage sensitivity differences.
 - Cannot discern chronic lethal from sublethal effects.
 - Omits persistence of loss concepts (recovery time): cannot discern short-lived from long-lived species.