

<https://gitlab.com/swejk12/esa-workshop>



# Generalized Linear Mixed Effects Modeling Deep Dive:

*Selecting Conditional Distributions, Checking Assumptions, and Drawing Inference for Ecological Data*

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## Introductions



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Associate Statistician



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Associate Statistician

## Who is WEST?



# Where is WEST?

<https://gitlab.com/swejk12/esa-workshop>

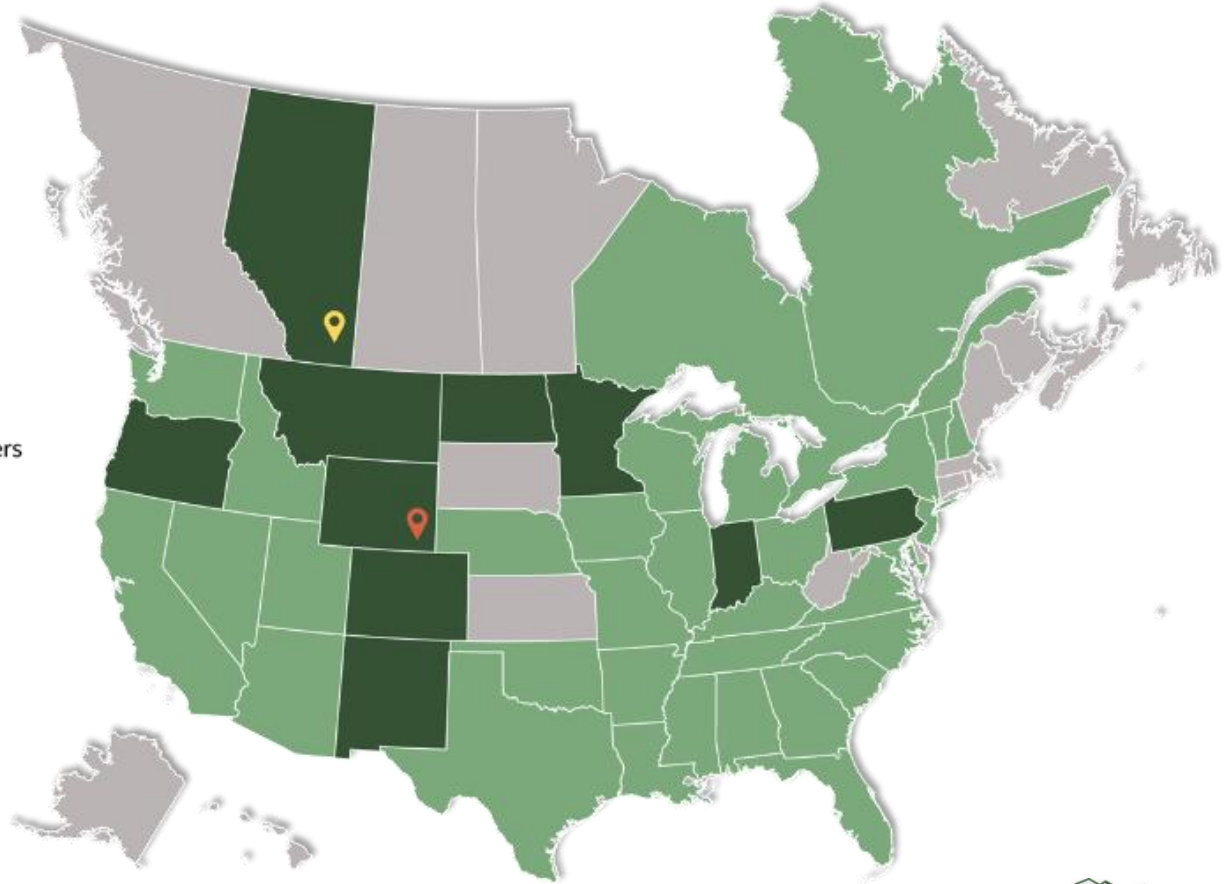
**WEST Office Locations**

- 📍 WEST Headquarters
- 📍 WEST, ULC Headquarters
- Branch Offices
- Field Offices

The map shows the following distribution:

- Field Offices (light green):** Most states are covered by field offices.
- Branch Offices (dark green):** Includes California, Nevada, Arizona, New Mexico, Texas, Oklahoma, Kansas, Nebraska, Minnesota, Wisconsin, Illinois, Indiana, Michigan, Ohio, Pennsylvania, Maryland, Delaware, Virginia, North Carolina, South Carolina, Georgia, Florida, Alabama, Mississippi, Louisiana, Arkansas, Missouri, Iowa, Kentucky, Tennessee, West Virginia, Maryland, Delaware, Virginia, North Carolina, South Carolina, Georgia, Florida, Alabama, Mississippi, Louisiana, Arkansas, Missouri, Iowa, Kentucky, Tennessee, West Virginia.
- WEST Headquarters (yellow pin):** Located in Washington state.
- WEST, ULC Headquarters (orange pin):** Located in Colorado.

**WEST**





## Our Expertise

- Aerial Surveys
- Bat and Avian Surveys
- Bird and Bat Conservation Plans
- Big Game and Mammal Surveys
- Collision Risk Modeling
- Ecosystems Study Design
- Statistical Analyses
- Estimation of Biological Size
- Resource Selection Studies
- Eagle Conservation Plans
- Evaluation of Mitigation Measures Effectiveness
- Expert Testimony
- Fatality Monitoring Studies
- Habitat Conservation Plans
- Impact Assessments and more...



## Outline

# Objectives

### Build

- Use graphical tools to determine *candidate* conditional distributions
- Specify random and fixed effects appropriately based on study design

### Assess

- Use DHARMA package to assess residuals
- Understand application of REML and ML
- Understand uses of AIC/BIC

### Draw Conclusions

- Use graphical tools to display results from a model
- Interpret results from a model to make biological conclusions.
- Understand variance structure and what it says about your system

# Generalized Linear Mixed Effects Modeling Basics

Although selection procedures are helpful exploratory tools, the model-building process should **utilize theory and common sense**

- Agresti 2002

There is no consensus in the statistics community about what constitutes correct practice, and there likely never will be. On the other hand, **there is broad consensus on what constitutes poor practice...**

- Tredennick et al. 2021

Whereas GLMMs themselves are uncontroversial, describing how to use them to analyze data necessarily touches on controversial statistical issues. **We acknowledge the difficulty while remaining agnostic.**

.....different methods are appropriate for different problems .... **how one analyzes data depends strongly on one's philosophical approach.**

- Bolker et al. 2008

# What is a Generalized Linear Mixed Model?

Mixed model  
(incorporates fixed and random effects)



Generalized linear model  
(non-normal data, link functions)

## Why Random Effects?

VARIATION

STRUCTURE

**temporal and spatial, account for replication, etc**



# Biological Conclusions and GLMM

Draw biological conclusions from estimates and confidence intervals to explain or understand relationships in ecological systems.

## Estimate Parameters

- Covariate effects and interactions
- Treatment effects
- Quantify variation
- Useful in exploration, inference, and prediction

## Exploration/Ecological Relationships

- Describe patterns and develop hypothesis about the natural world
- Trade-offs: Including variables of interest vs. spurious relationships
- Type-I errors (false discoveries)
- Make use of biological intuition

## Inference

- Relationships/associations between response and covariates
- Evaluate strength of evidence for patterns
- Null-hypothesis testing
- Fewer models: lower risk of Type-I errors

## Prediction

- Overlaps with exploration and inference
  - Model that best explains process should improve prediction, right?
- Focused on predicting the mean

# What Makes it a "Mixed" Model?

“nuisance variable”, if it  
contributes a lot to variation,  
“unmeasured variation that  
needs to be accounted for”

	Fixed Effects	Random Effects
Use:	Test <u>statistical significance or relationship</u>	<i>Don't want to test statistical significance</i>  Test another effect across all levels
Models:	Mean structure	Variance structure <i>nesting of sites within islands, include island.</i>
Levels of a variable:	Fully represented in data  Ex) <i>All measurements have an associated habitat or island of covariate that we can assess a relationship</i>	Treated as from a population effects  <i>Think block design or repeat measures. Samples from a greater population</i>
Numbers of levels:	Finite	Many
Basis for inference:	Levels of the variable present in the data	Larger population

Hierarchy and correlation of your variables

# GLMM Components

Conditional Distribution

Link Function

Random Effect Structure

Fixed Effects Structure

## Simple Linear Regression

- $y = \beta_0 + \beta_1 x + \varepsilon$ 
  - $y$  is the dependent variable
  - $\beta_0$  and  $\beta_1$  are regression coefficients define mean of y
  - $x$  is the independent linear predictor
  - $\varepsilon$  is the error term
- Fixed effects:  $\beta_0, \beta_1$ 
  - Define the **mean** of  $y$
- Random effect:  $\varepsilon$ , where  $\varepsilon \sim N(0, \sigma^2)$  doesn't effect mean structure at all, but does impact variance
  - Defines the **variance** of  $y$

## Breaking Down Random Effects

transformed mean, E is the expected mean, conditioning on u, random

Simple Linear Regression:  $y = \beta_0 + \beta_1 x + \varepsilon$

GLMM:  $g[E(y_{ijk} | \mathbf{u}_{ijk})] = \beta_0 + \beta_1 w_j + b_j + a_i + t_i w_j + c_{ijk}$

- Random year effect:  $b_j \sim N(0, \sigma_b^2)$  normal with mean 0, and year to year variation
- Random site effects:  $\begin{pmatrix} a_i \\ t_i \end{pmatrix} \sim \text{MVN} \begin{pmatrix} 0 & \sigma_a^2 & \sigma_{at} \\ 0 & \sigma_{at} & \sigma_t^2 \end{pmatrix}$  correlated site effects, t is random site slope giving trendline of each site
- Random site-by-year interaction effect:  $c_{ijk} \sim N(0, \sigma_c^2)$  same site within a year, variation accounted for
- $\sigma_b^2, \sigma_a^2, \sigma_t^2$ , and  $\sigma_c^2$  are variance components



## GLMM Form

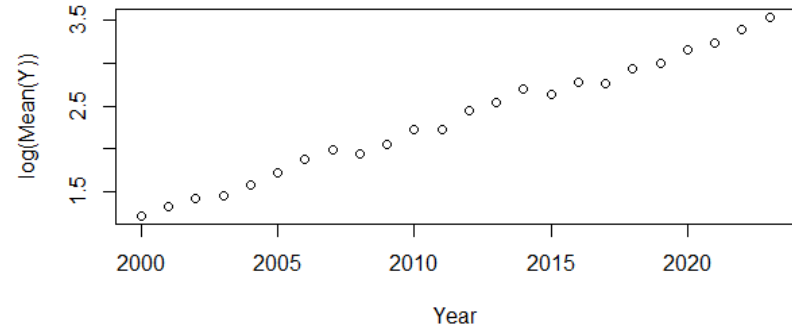
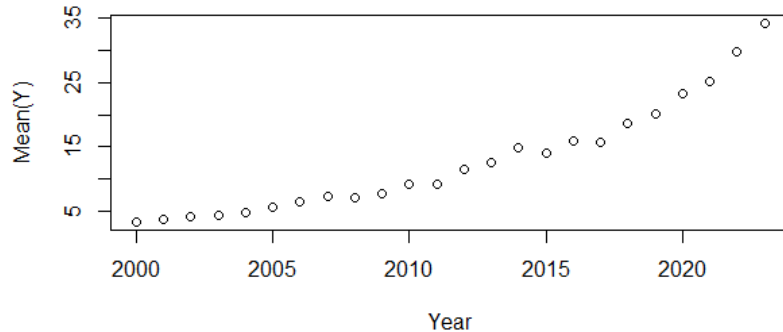
$$g[E(\mathbf{y}|\mathbf{u})] = \boldsymbol{\eta} = \boldsymbol{\beta}\mathbf{X} + \mathbf{Z}\mathbf{u}$$

- $g(.)$  is the link function (e.g. log, logit) modeling the mean of  $\mathbf{y}$ ,
- $E$  is the expectation for the conditional distribution of  $\mathbf{y}$
- $\boldsymbol{\eta}$  is the linear predictor point of link is to get linear relationships for this model
- Fixed effects and random effects

If cant get linear, its time for GAM

# Link Function

- Transforms the expected value of the response variable so that relationships with predictors are **linear**
- Common link functions:
  - Gaussian: identity (LMM)
  - Counts, non-negative continuous data: log
  - Proportions: logit, probit, log-log, clog-log



# GLMM Components

Conditional Distribution

Based on data structure  
Exponential family

Random Effect Structure

Link Function

Transforms the expected value  
of the response variable so that  
relationships are linear

Fixed Effects Structure

# Conditional Distributions of the Response

- Also called “error distributions”
- Exponential family
  - Poisson
  - Negative binomial (two forms)
  - Gamma
  - Tweedie “amazing”
  - Binomial
  - Beta
- glmmTMB package also includes mixture and truncated distributions
  - Zero inflation (excess zeros)
  - Zero truncation (no zeros)

Distribution

Data Type

Assumptions

Application

# Conditional Distributions of the Response

Binomial	Beta	Beta-Binomial
Binary, presence-absence, proportions	<u>Proportion</u> : not based on Bernoulli trials	Overdispersed or correlated binomial data: presence/absence
Independent trials with a binary outcome	Two scale parameters	Binomial probability is a Beta random variable
Family = binomial Link = logit, probit, log-log, clog-log	Family = beta Link = logit, probit, log-log, clog-log	Family = betabinomial Link = logit, probit, log-log, clog-log



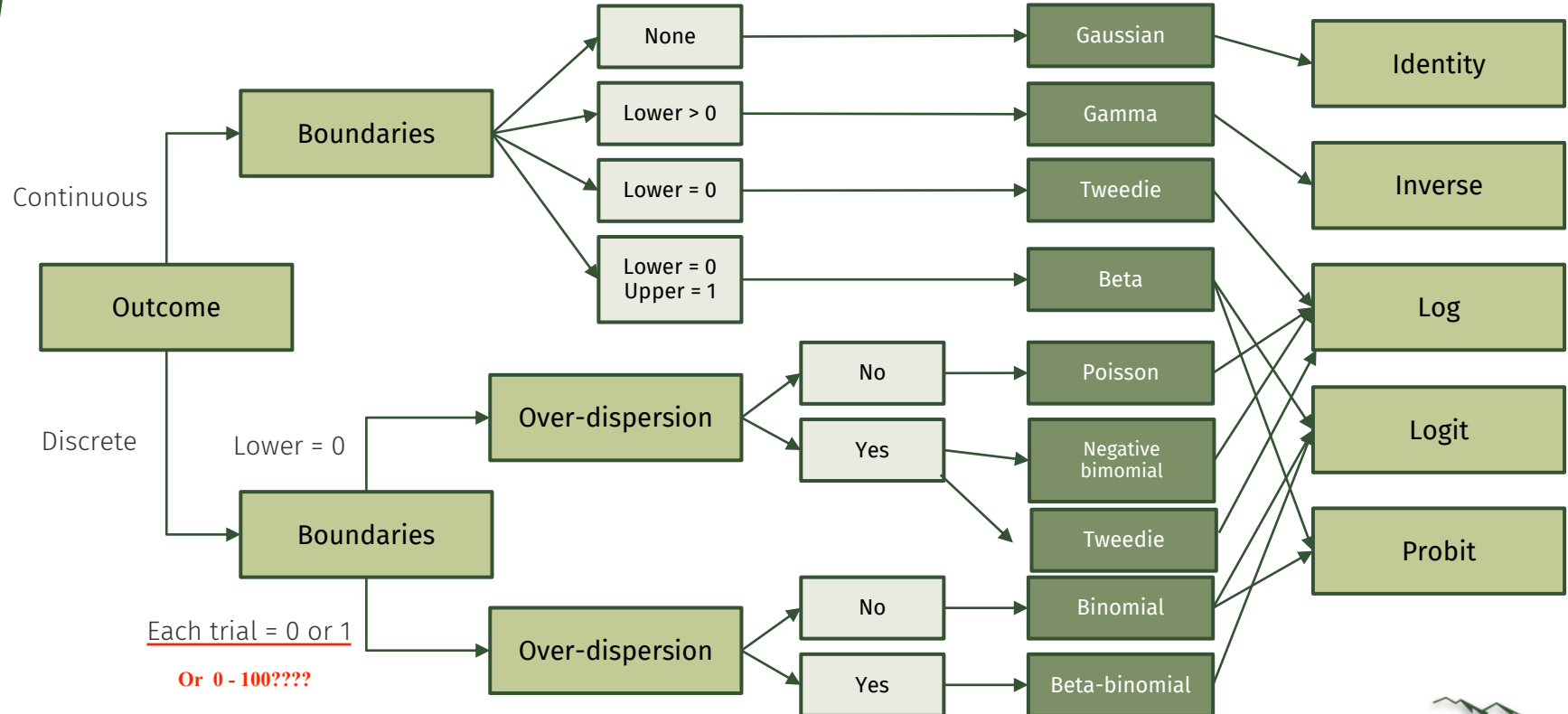
# Conditional Distributions of the Response

Poisson	Negative Binomial
Counts: abundance	Counts with overdispersion: abundance
Variance is equal to the mean	Relationship between variance and mean: <i>nbinom1</i> : linear <i>nbinom2</i> : quadratic
Family = poisson Link = log, identity	Family = nbinom1, nbinom2 Link = log, identity

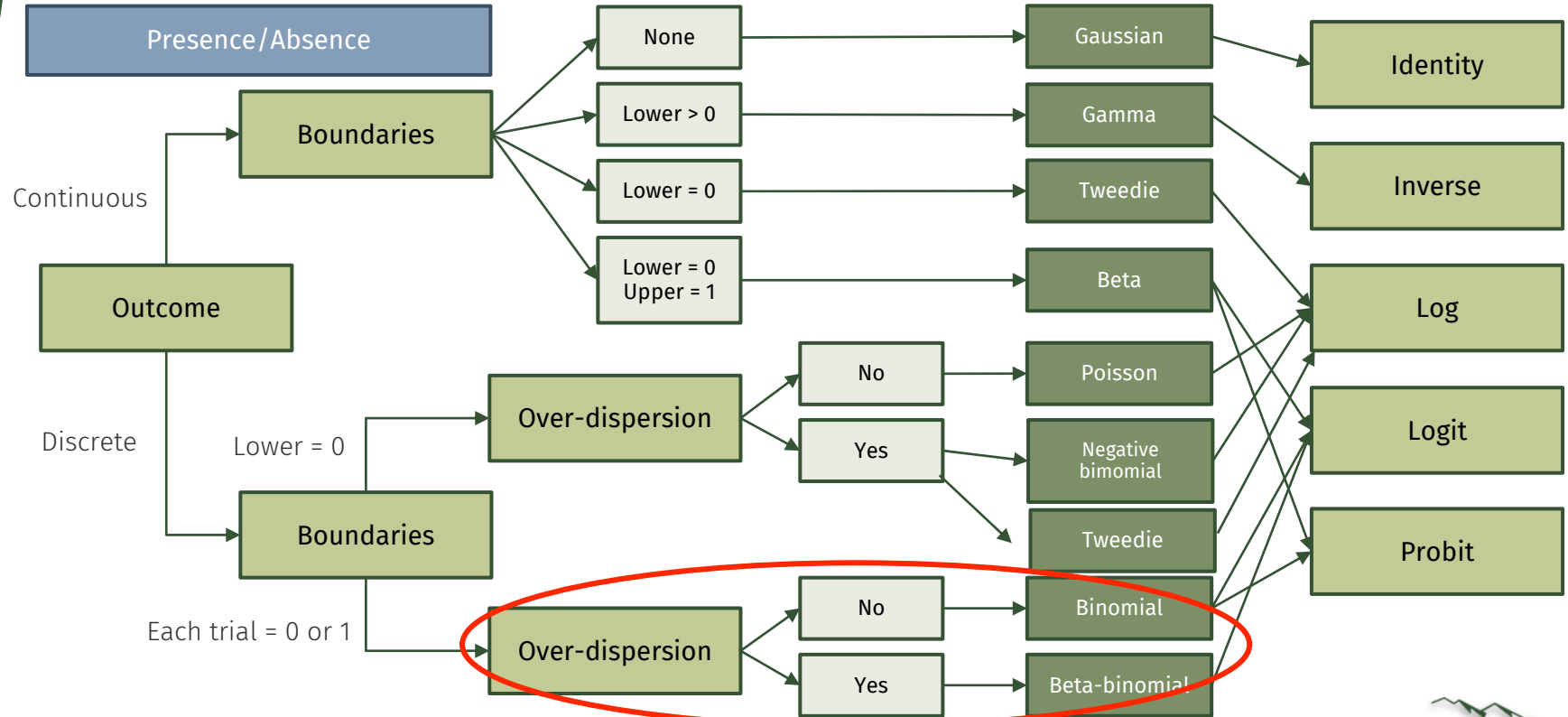
# Conditional Distributions of the Response

Gaussian	Gamma	<u>Tweedie</u>
Continuous metric: temperature, time	Positive continuous data: time	Continuous or count, <u>may contain zeros</u> : abundance, weight, ...
Linear mixed model Variance is not a function of the mean	Mean is linear in scale parameter, variance is quadratic in scale parameter	Useful for continuous data with zeros
Family = gaussian Link = identity	Family = gamma Link = inverse	Family = tweedie Link = log

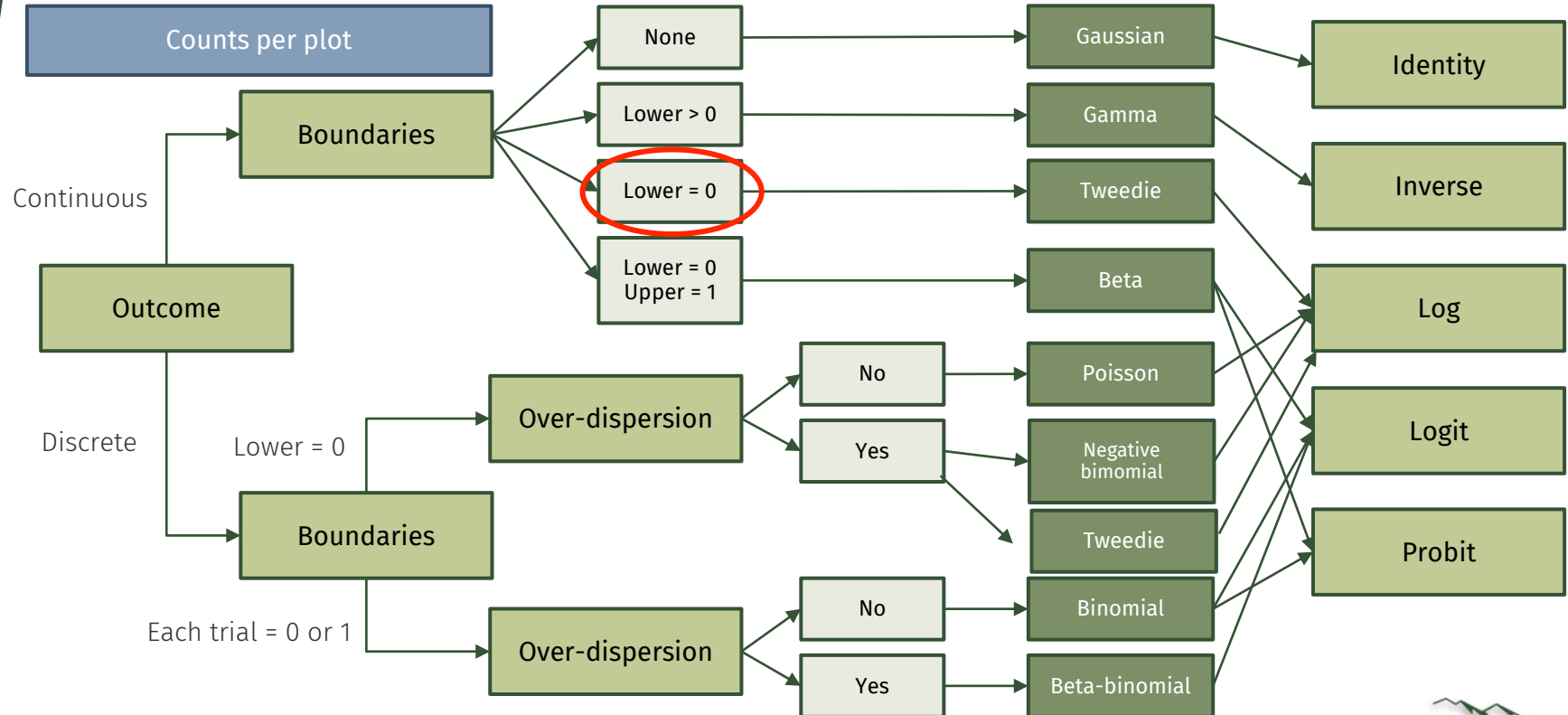
# Conditional Distributions and Link Functions



# Conditional Distributions and Link Functions

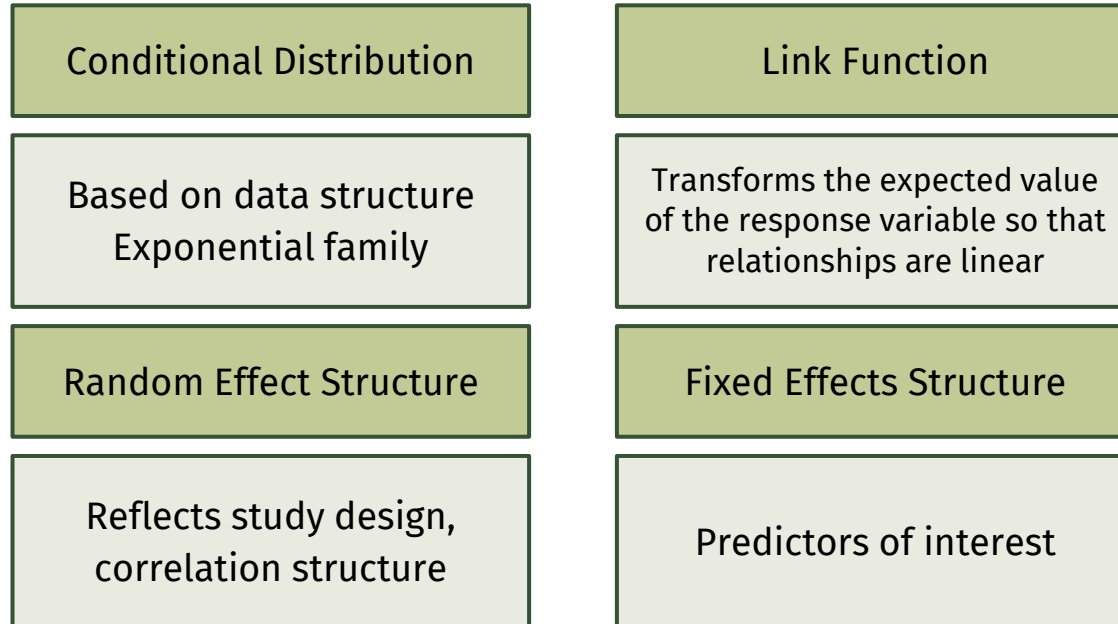


# Conditional Distributions and Link Functions





# GLMM Components



# Specifying Random Effects in R

Random intercept

$(1|\text{Site})$

Correlated random intercept (site)  
and slope (year)

$(1+W\text{Year}|\text{Site})$

Nested random effects

$(1|\text{Site}/\text{Island})$

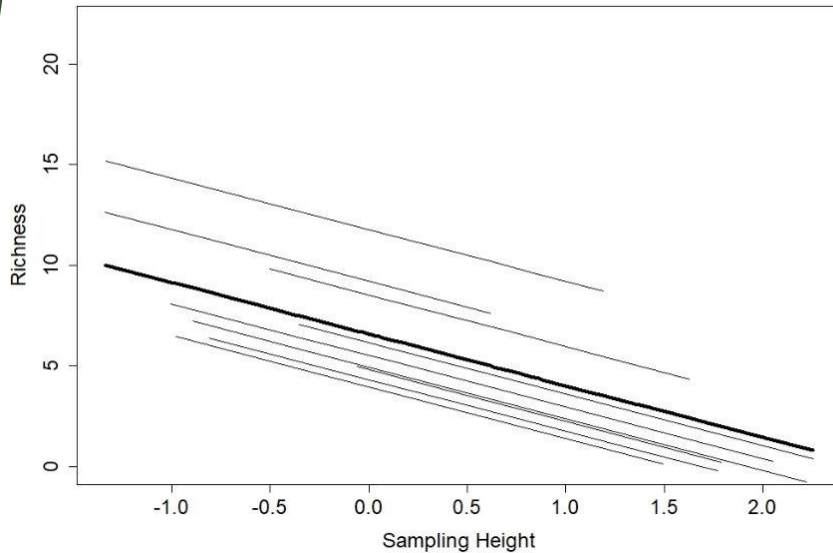
Uncorrelated random intercept and  
Year slope

$(1+W\text{Year}||\text{Site})$  or  
 $(1|\text{Site}) + (-1+W\text{Year}|\text{Site})$

## Random Intercept

Richness ~ Sampling Height +  
(1|Beach)

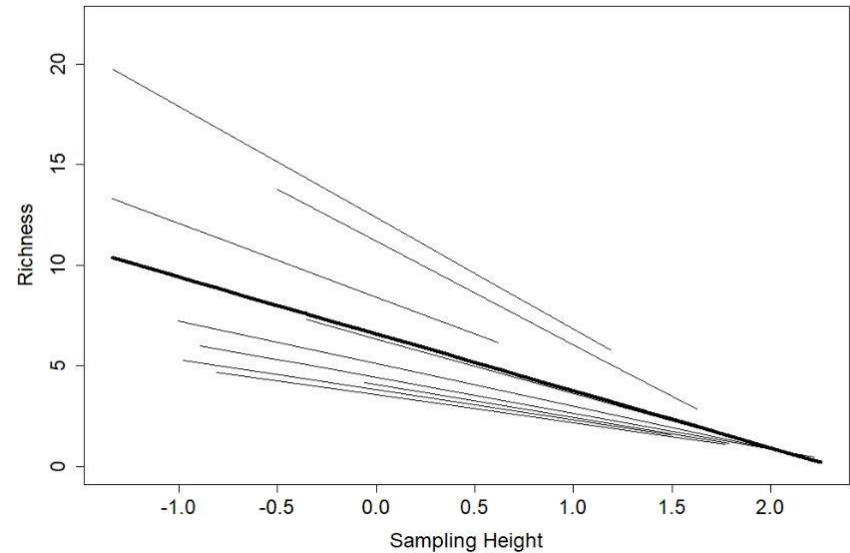
intercept varies slope does not



## Random Slope and Intercept

Richness ~ Sampling Height +  
(1 + Sampling Height|Beach)

random slope by height and beach



## Build and Assess

Conditional Distribution and Link

Determine family and link  
Check for linearity & homogenous  
variance  
Identify outliers

Check Model Assumptions

Specify and Fit the Full Model

Finalize Model

## Exercise 1:

Pick the *candidate* conditional distributions

Explore relationships and consider fixed and random effects for model fitting

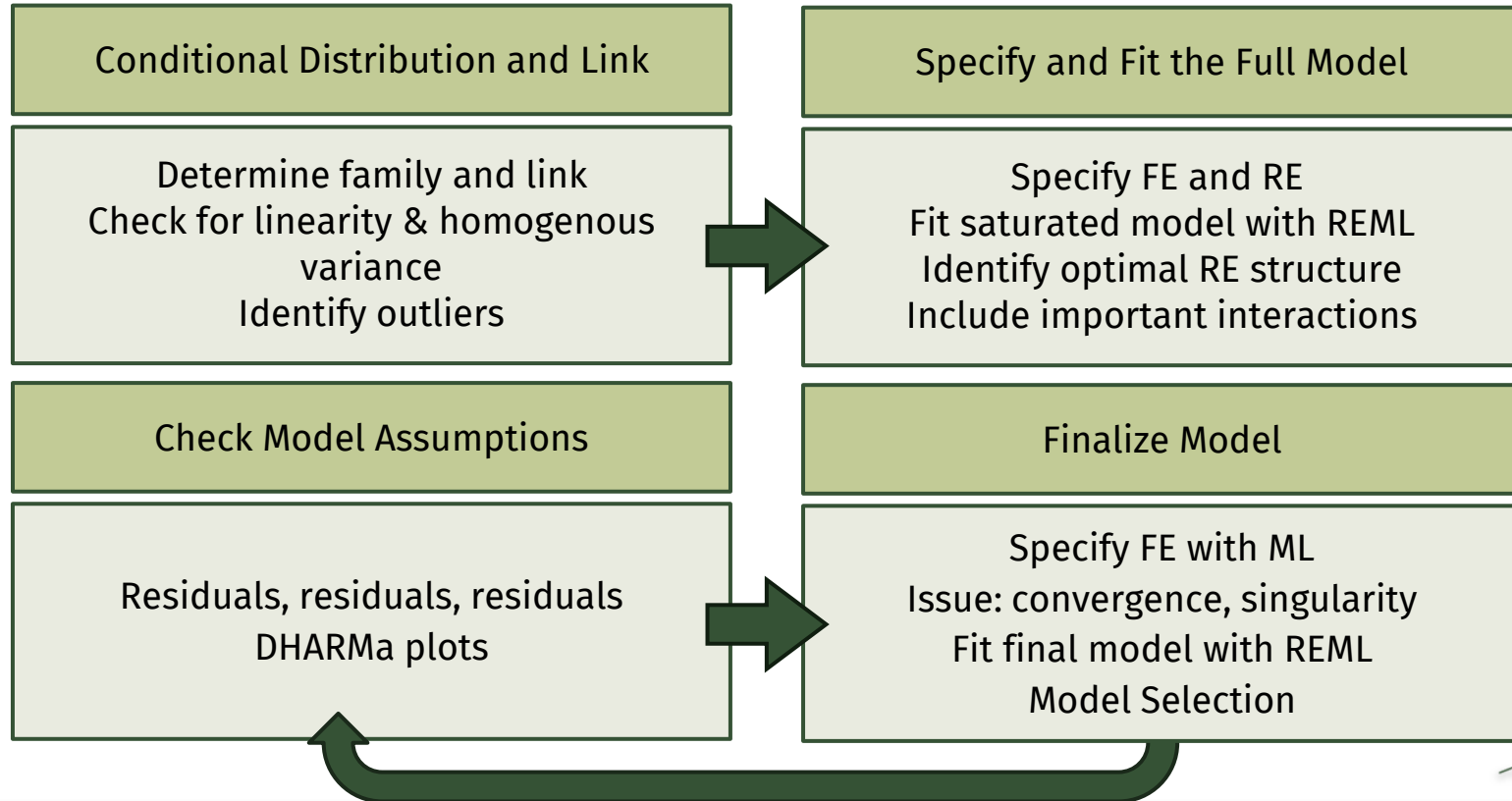
Objectives:

Use graphical tools to determine *candidate* conditional distributions

Examine data and variance structures to think about possible random effect structure



## Build and Assess



## Residual Diagnostics - "Goodness of fit" - DHARMa

### Kolmogorov-Smirnov Test

Test of uniformity of residuals  
Does the assumed distribution match the data?

### Overdispersion

More variation than expected based on distribution → **misspecified model**  
Underestimate FE SEs  
Inflated type-I errors

### Underdispersion

Less variation than expected based on distribution  
Low power to detect relationship

Visual detection or test (e.g. Pearson chi-squared)

### Zero inflation

Common cause of overdispersion  
Compare ZI family to alternative

### Heteroscedasticity

Patterns in the residuals  
Check quantile deviations plot  
Some variation not accounted for in structure  
Missing predictors?

### Residual correlation

Spatial or temporal

# Check Model Assumptions

Distribution and Link

Check linear relationship with link-transformed means

RE are Independent of Residuals

Pearson's correlation tests

Independence

Partial autocorrelation plots  
of residuals

Normality of Random Effects

Quantile-quantile plots and histograms of  
random effects

## Exercise 2:

Fit a GLMM that we expect to have variation across space and time.

### Objectives:

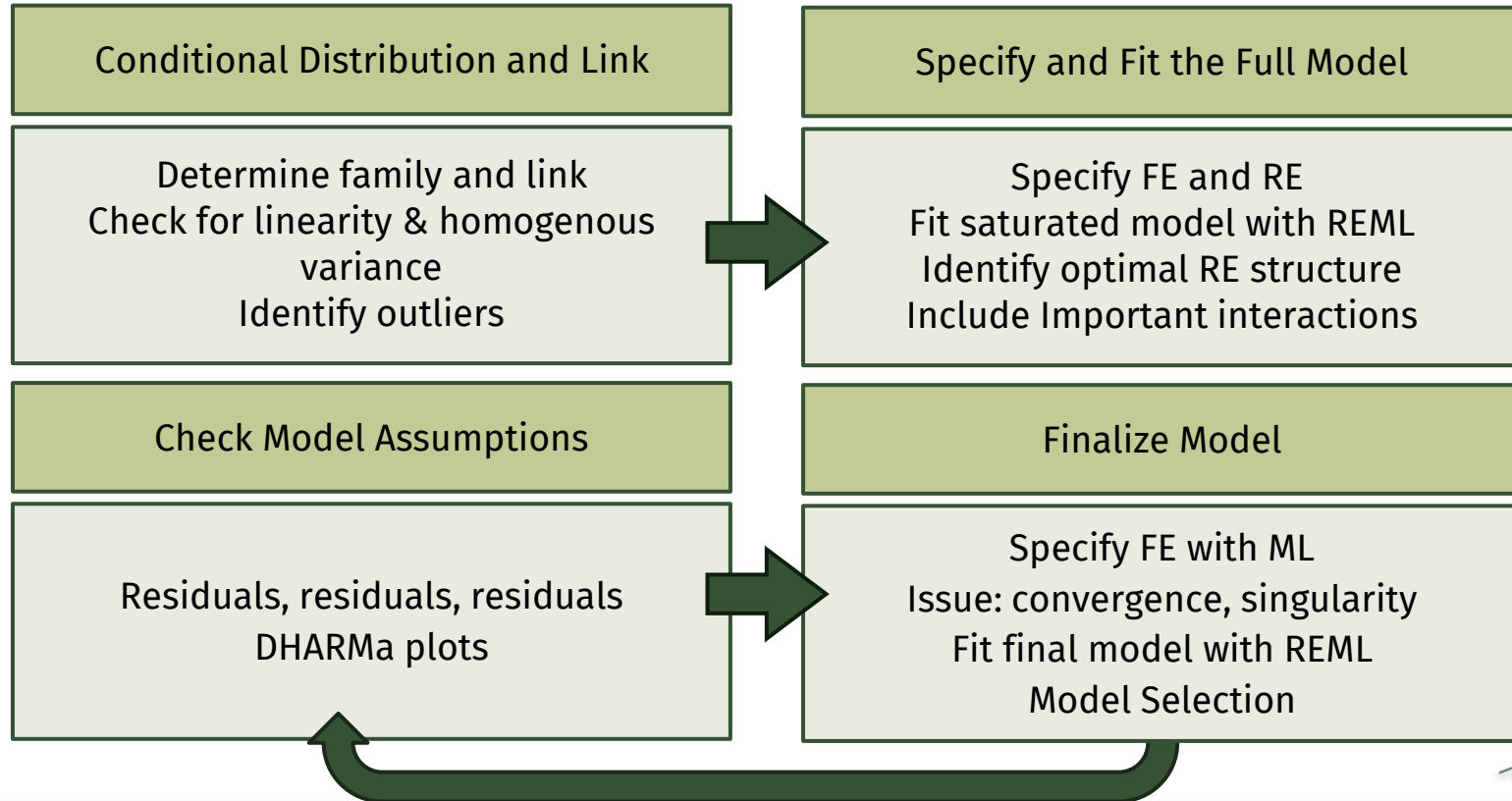
Identify the conditional distribution.

Specify random and fixed effects.

Compare a random intercept model to a random slope and intercept model.

Assess residuals using DHARMA.

## Build and Assess



## Exercise 3:

Incorporate design variables into random effects structure

Objectives:

Specify design variable in our model.

Assess residuals using DHARMa.

## Specifying Random Effects in R (reminder)

Random intercept

$(1|\text{Site})$

Nested random effects

$(1|\text{Site}/\text{Island})$

Correlated random intercept and  
Year slope

$(1+W\text{Year}|\text{Site})$

Uncorrelated random intercept and  
Year slope

$(1+W\text{Year}||\text{Site})$  or  
 $(1|\text{Site}) + (-1+W\text{Year}|\text{Site})$

# Importance of Variance Components

- Decomposing the error into variance components tells us something about the response variable
  - High variation from year-to-year  $\Rightarrow$  Not a useful metric to monitor over time, requires long time periods to overcome noise in data and detect signal
  - High variation from site to site  $\Rightarrow$  Adequate sample size of sites needed
  - High variation among trend lines at a site  $\Rightarrow$  Possible subpopulations responding differently to stressors
  - High site-by-year interaction variance  $\Rightarrow$  need replication within a site and year (e.g., quarterly surveys), possible crew variability, may need to focus on a specific time period each year



## AIC and BIC

- Both are used in model selection to assess model fit and complexity
- Fit based on likelihood
- Complexity is based on the number of parameters
- BIC also accounts for number of observations and penalizes more complexity
- AIC/BIC use:
  - ✗ REML  $\leftrightarrow$  ML
  - ✓ REML  $\leftrightarrow$  REML
  - ✓ ML  $\leftrightarrow$  ML
- AIC is best for prediction because it might be more forgiving of spurious correlations than NHST. More covariates typically improve prediction (Tredennick et al. 2021)
- AIC can be adjusted for overdispersion (QAIC)
- Model selection with LRT could be an abuse of hypothesis testing (Bolker et al. 2008)
- Bayes factor can be used as LRT alternative. Outcome similar to recommendations from BIC

## REML and ML

- REML handles variance better than ML
- ML produces more biased variance estimates
- Balanced vs. unbalanced
- Slope and intercept may be similar across models, but variance should differ
- AIC/BIC use:
  - ✗ REML <--> ML
  - ✓ REML <--> REML
  - ✓ ML <--> ML
- REML should be used for finalizing random effects
- ML should be used to finalize fixed effects
- Final model should be presented using REML.

## Covariate and Model Selection

- Depends on the goal
- Adds interactions that are biologically meaningful
- Too many variables:
  - Remove variables that have correlation coefficient of  $\geq 0.3$
  - Biologically informed
  - Question/Objective informed
  - LASSO
- AIC/BIC/QAIC can be used for comparison of model
  - REML for Random Effects, ML for fixed effects
- Top-down approach for covariate selection
  - Fit full model first
- `stats::drop1()` function
  - Compares models by dropping covariates one at a time
  - Should use correction for multiple hypothesis tests

# References

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