

Introductions



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Who is WEST?

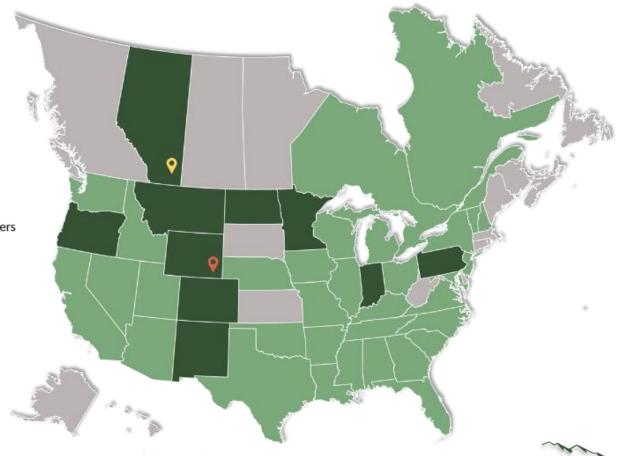




Where is WEST?

WEST Office Locations

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



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

Inference

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

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

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 statistical significance or relationship	Don't want to test statistical significance
		Test another effect across all levels
Models:	Mean structure	Variance structure nesting of sites within
Levels of a variable:	Fully represented in data	islands, include island. Treated as from a population effects
	Ex) All measurements have an associated habitat or island of covariate that we can assess a relationship	Think block design or repeat measures. Samples from a greater population
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
 - β_0 and β_1 are regression coefficients define mean of y
 - x is the independent linear predictor
 - ε is the error term
- Fixed effects: β_0 , β_1
 - Define the mean of y
- Random effect: ε , where $\varepsilon \sim N(0, \sigma^2)$ doesnt effect mean structure at all, but does impact variance
 - Defines the variance of y



Breaking Down Random Effects

transformed mean, E is the espected 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: $b_i \sim N(0, \sigma_b^2)$ normal with mean 0, and year to year variation
- Random site effects: $\binom{a_i}{t_i} \sim \text{MVN} \begin{pmatrix} 0 & \sigma_a^2 & \sigma_{at} \\ 0' & \sigma_{at} & \sigma_t^2 \end{pmatrix}$ 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
- σ_b^2 , σ_a^2 , σ_t^2 , and σ_c^2 are variance components



GLMM Form

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

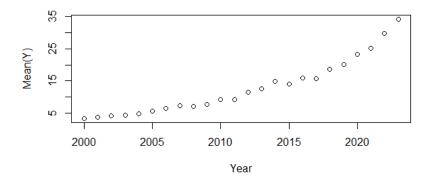
- g(.) is the link function (e.g. log, logit) modeling the mean of y,
- $oldsymbol{\cdot}$ E is the expectation for the conditional distribution of $oldsymbol{y}$
- η is the linear predictor point of link is to get linear relationships for this model
- Fixed effects and random effects

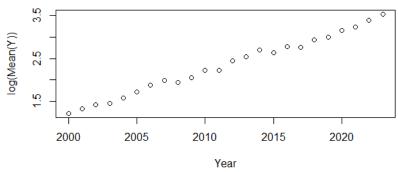
If cant get linear, its time for GAM



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



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

Distribution

Data Type

Assumptions

Application



Binomial

Beta

Beta-Binomial

Binary, presence-absence, proportions

<u>Proportion</u>: not based on Bernouilli 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



Poisson

Negative Binomial

Counts: abundance

Counts with overdispersion: abundance

Variance is equal to the mean

Relationship between variance and mean:

nbinom1: linear

nbinom2: quadratic

Family = poisson Link = log, identity Family = nbinom1, nbinom2 Link = log, identity



Gaussian

Gamma

Tweedie

Continuous metric: temperature, time

Positive continuous data: time

Continuous or count, <u>may contain</u> <u>zeros</u>: abundance, weight, ...

Linear mixed model
Variance is not a function of
the mean

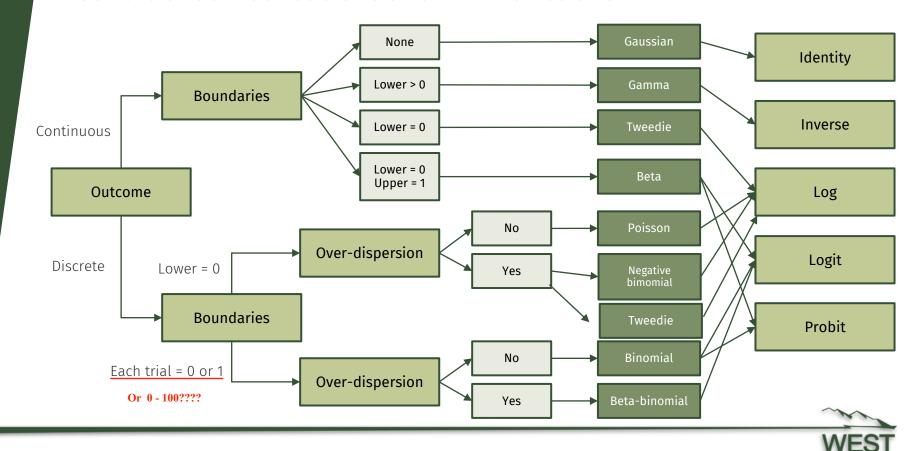
Mean is linear in scale parameter, variance is quadratic in scale parameter

Useful for continuous data with zeros

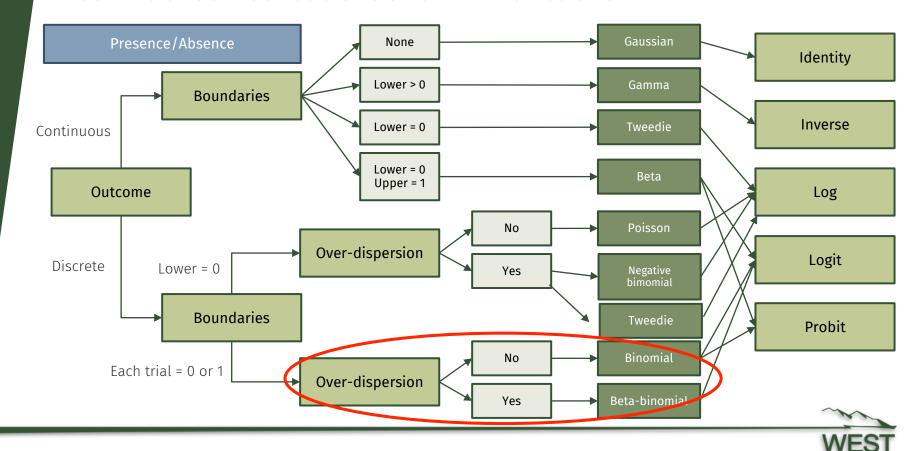
Family = gaussian Link = identity Family = gamma Link = inverse Family = tweedie Link = log



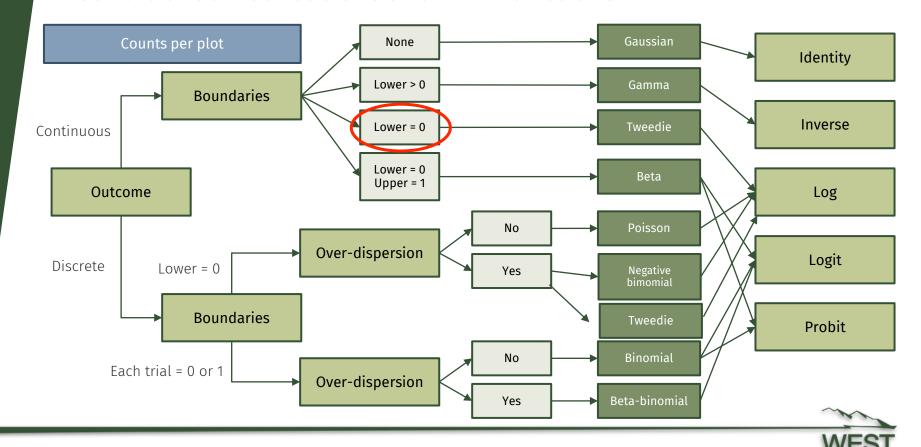
Conditional Distributions and Link Functions



Conditional Distributions and Link Functions



Conditional Distributions and Link Functions



GLMM Components

Conditional Distribution

Based on data structure Exponential family

Random Effect Structure

Reflects study design, correlation structure

Link Function

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

Fixed Effects Structure

Predictors of interest



Specifying Random Effects in R

Random intercept

Nested random effects

(1|Site)

(1|Site/Island)

Correlated random intercept (site) and slope (year)

Uncorrelated random intercept and Year slope

(1+WYear|Site)

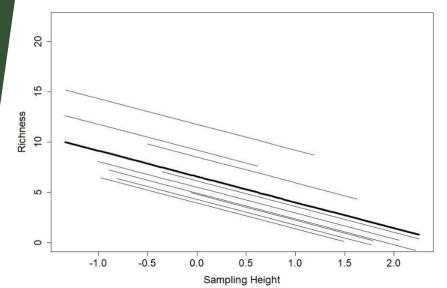
(1+WYear||Site) or (1|Site) + (-1+WYear|Site)



Random Intercept

Richness ~ Sampling Height + (1|Beach)

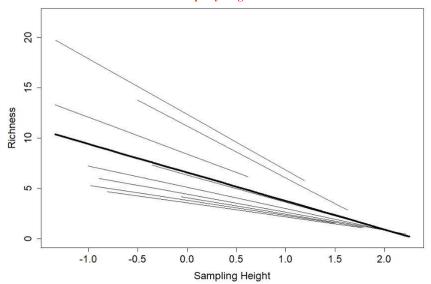
intercept varies slope does not



Random Slope and Intercept

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

random slope by height and beach





Build and Assess

Conditional Distribution and Link

Specify and Fit the Full Model

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

Check Model Assumptions

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

Conditional Distribution and Link

Specify and Fit the Full Model

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

Specify FE and RE
Fit saturated model with REML
Identify optimal RE structure
Include important interactions

Check Model Assumptions

Finalize Model

Residuals, residuals, residuals

DHARMa plots

Specify FE with ML
Issue: convergence, singularity
Fit final model with REML
Model Selection



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

<u>Common cause of overdispersion</u> 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

Independence

Check <u>linear relationship with link</u>transformed means Partial <u>autocorrelation plots</u> of residuals

RE are Independent of Residuals

Normality of Random Effects

Pearson's correlation tests

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

Conditional Distribution and Link

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

Check Model Assumptions

Residuals, residuals, residuals

DHARMa plots

Specify and Fit the Full Model

Specify FE and RE
Fit saturated model with REML
Identify optimal RE structure
Include Important interactions

Finalize Model

Specify FE with ML
Issue: convergence, singularity
Fit final model with REML
Model Selection



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

Nested random effects

(1|Site)

(1|Site/Island)

Correlated random intercept and Year slope

Uncorrelated random intercept and Year slope

(1+WYear|Site)

(1+WYear||Site) or (1|Site) + (-1+WYear|Site)



Importance of Variance Components

- Decomposing the error into variance components tells us something about the response variable
 - High variation from year-to-year ⇒ 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 ⇒ Adequate sample size of sites needed
 - High variation among trend lines at a site ⇒ Possible subpopulations responding differently to stressors
 - High site-by-year interaction variance ⇒ 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:
 - X REML <--> ML
 - ✓ REML <--> REML
 - ✓ ML <--> ML

do not compare aics between reml and 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:
 - X 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 >= 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



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