

Introduction(week 1, part 1)

4 Sep 2023

Basics

Land acknowledgement

McMaster University (and this class) is on the traditional territories of the Mississauga and Haudenosaunee nations, and within the lands protected by the “Dish with One Spoon” wampum agreement. (Why? See [“Beyond Territorial Acknowledgements”](#))

Logistics

- (almost) everything at the [course web page](#)
- communication/forums ([Piazza](#)); e-mail if necessary
- assignment marks (Avenue)
- Zoom/recordings (by request)

Integrity

- [notes on honesty](#)
- why copying code is good
- Stack Overflow, ChatGPT, and all that
- group work

Prerequisites

From the course outline:

- basics of linear models (as in [STATS 3A03](#)), with associated linear algebra
- basics of generalized linear models (as in [STATS 4C03/6C03](#)), including knowledge of exponential family distributions
- inferential statistics: sampling distributions, Central Limit theorem, hypothesis testing, Wald tests, maximum likelihood estimation
- ideally, *basic* knowledge of Bayesian statistics and Markov chain Monte Carlo estimation
- intermediate knowledge of R

Goals

- principles/practices of statistical modeling
 - choosing a model
 - diagnostics and troubleshooting
 - interpreting and communicating results
- understanding the tools ((penalized) (G)L(A)(M)Ms; unifying principles of regression modeling, shrinkage/penalized estimators
 - both frequentist and Bayesian approaches
- awareness of computational foundations/scaling

Scope

- **regression** modeling
- the components:
 - linear model matrices/basis spaces
 - link functions
 - conditional distributions (GLM “families”)
 - penalization/shrinkage
- includes a vast range of useful models

Technical skills & tools

Not focal, but unavoidable and useful

- R (base + some [tidyverse](#))
- reproducibility (Jenny Bryan (2017); Jennifer Bryan (2017))
 - version control (Git/GitHub)
 - documents: [Quarto](#)

about me

- weird background (physics/math u/g, Zoology PhD, epidemiological modeling)
- math biology (ecology/evolution/epidemiology)
- computational statistics (mixed models, Bayesian stats)

things I like/obsess about

- scientific inference \gg pure prediction (but see Navarro (2019))
- data visualization
- solving problems in context, practical issues
- battling bad statistical practice (p-value abuse, snooping, dichotomania, imbalance handling, ...)

a tiny bit of philosophy

- frequentist vs. Bayesian
- *computational* or *pragmatic* Bayesian
- in a perfect world either choice would be available for any problem
 - Bayesian approaches generally more flexible but slower
 - priors!
- hierarchical/latent/mixed models make the boundaries fuzzy (*empirical Bayesian*)

strong feelings about p-values

- if you use frequentist methods, please use them correctly
- H_0 is almost never true
- never accept the null hypothesis
- a low p value doesn't mean the effect is large or important
- may help to frame results in terms of **clarity** (Dushoff, Kain, and Bolker 2018)

The modeling cycle

Before you start

- you need to know what the question is!
- this is hard for statisticians
 - when mining data, go back to the original paper(s)
- a low p -value is not inherently interesting!
- what is a large effect? what is an interesting effect?

Effect sizes

- standardized measures like Cohen's d $((\bar{x}_1 - \bar{x}_2)/s$, where s is some measure of pooled standard deviation: [Wikipedia](#)) are common ...
- but shouldn't be used mindlessly. Real-world, unstandardized effects are usually more meaningful
- effects estimated on the log or logit scale are unitless and hence *may* be easier to generalize
- scaling predictors and responses may help (Schielzeth 2010)

An iterative process ...

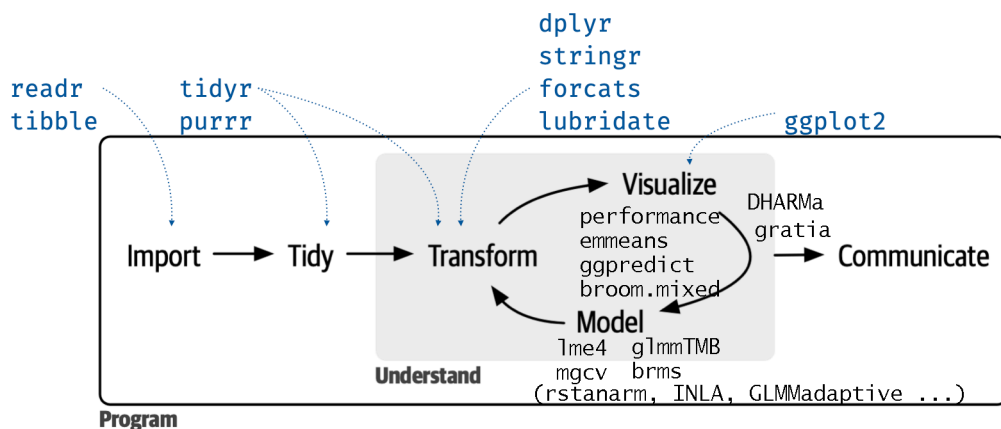


Figure 1: original from [Mine Çetinkaya-Rundel](#)

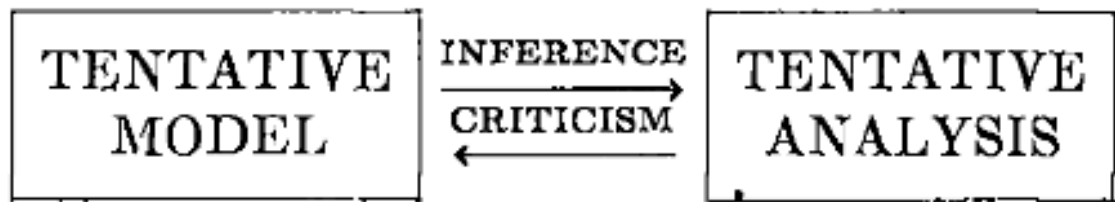


Figure 2: From Box (1976)

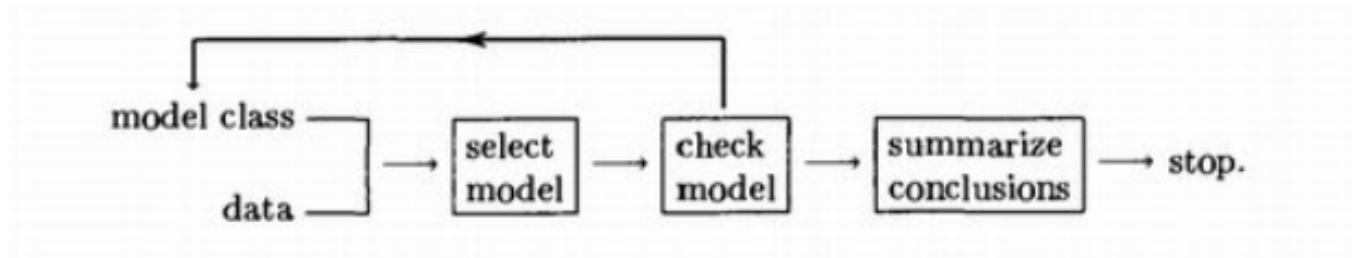


Figure 3: From McCullagh and Nelder (1989) p. 392: ‘The introduction of this loop changes profoundly the process of analysis and the reliability of the final models found.’

Beware the garden of forking paths!

- “researcher degrees of freedom”, “HARKing”, etc.
- Simmons, Nelson, and Simonsohn (2011); Gelman and Loken (2014)

Solutions?

- pre-registration (formal or informal); report deviations from planned analysis
- choose model complexity (see Harrell RMS ch. 3), do diagnostics etc., **without reference to response variable** or metrics of significance

Choosing model complexity

- see Harrell ch. 3
- rules of thumb for inferential models with adequate power
 - e.g. $p < n/10$ or $n/20$
 - effective n depends on data type (binary < small counts < continuous)
 - how does clustering/correlation in data change effective n ?



Figure 4: from [Art Share LA](#)

- simplest if done **a priori**
 - data-driven choice of model complexity (e.g. by cross-validation), *while maintaining valid inference*, is delicate

Comparing models with reality

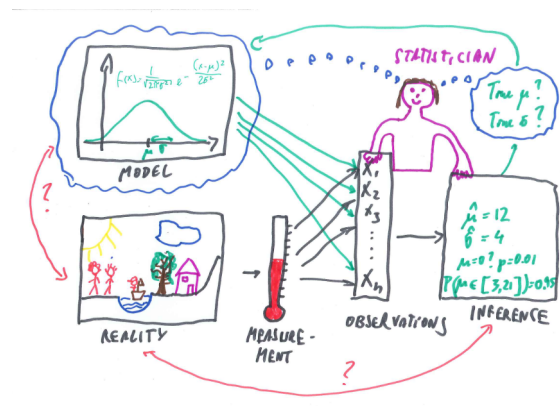


Figure 5: one view of models and reality, from Hennig (2022)

Model diagnostics

- all models make assumptions
- results *may* be sensitive to **misspecification**: bias, inefficiency, inflated/deflated type I error, poor coverage ...

- hypothesis tests (e.g. Shapiro-Wilk) are deprecated

Harvey Motulsky on [CrossValidated](#):

The question normality tests answer: Is there convincing evidence of any deviation from the Gaussian ideal? With moderately large real data sets, the answer is almost always yes.

The question scientists often expect the normality test to answer: Do the data deviate enough from the Gaussian ideal to “forbid” use of a test that assumes a Gaussian distribution? Scientists often want the normality test to be the referee that decides when to abandon conventional (ANOVA, etc.) tests and instead analyze transformed data or use a rank-based nonparametric test or a resampling or bootstrap approach. For this purpose, normality tests are not very useful.

- “is there a statistically significant deviation from the model assumptions?” vs. “are the violations of the assumptions large enough to mess up my conclusions?” (**never** “are the data normally distributed?”)
 - **data “too big”**: will reject assumptions even when it’s OK
 - **data “too small”**: will fail to reject assumptions even when they’re problematic (???)
- **two-stage testing** often has bad properties (H. Campbell and Dean 2014; Harlan Campbell 2021; Rochon, Gondan, and Kieser 2012; Zimmerman 2004; Shamsudheen and Hennig 2021)
- graphical diagnostics are often recommended
- but how do we judge whether deviations are too large? ([Q-Q plot survey](#))
- **open question**

Untestable issues are often the worst problems

- biased/non-representative sampling (Meng 2018)
- lack of randomization
- problems with causal inference: unobserved confounders, etc.
- non-independence
 - pseudo-replication (Hurlbert 1984)
 - temporal/spatial structure

More on model diagnostics

- assumptions apply to **conditional** distribution of the response (not the marginal distribution, not the predictor variables)
- test assumptions in order of importance (George Box: “It is inappropriate to be concerned about mice when there are tigers abroad.”)
 - bias/nonlinearity (residuals vs fitted plot)
 - heteroscedasticity (scale-location plot)
 - influential points/outliers (Cook’s distance, leverage, etc.)
 - distributional assumptions (Q-Q plot)
 - ¿¿ correlated predictors ??
- performance, DHARMA packages: more later

References

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