

Background

Medical research often involves hierarchical data structures requiring the use of mixed effects models

Primary outcomes are often categorical, such as whether a patient develops a disease or not

Generalized linear mixed models (GLMMs) account for hierarchical structure and categorical outcomes, but are instable for rare outcomes and small samples. Confidence intervals for predicted probabilities are not well defined.

Bayesian GLMMs incorporate prior information helping with small samples, and posterior predicted probabilities can be drawn from a distribution to easily obtain estimates and credible intervals

Method & Sample

- Patients (n = 1125) in critical care units in one health system had their treatment assessed over multiple days by treating physicians (n = 36)
- Physicians assessed treatment as *not futile*, *probably futile*, or *definitely futile* each day
- A total of 6,897 observations were cross classified in both patients and physicians
- Analyses used a cross classified random effects Bayesian ordered probit model using the MCMCglmm package (Hadfield, 2010) in R 2.15.2 (R Core Team, 2012)

Technical Details

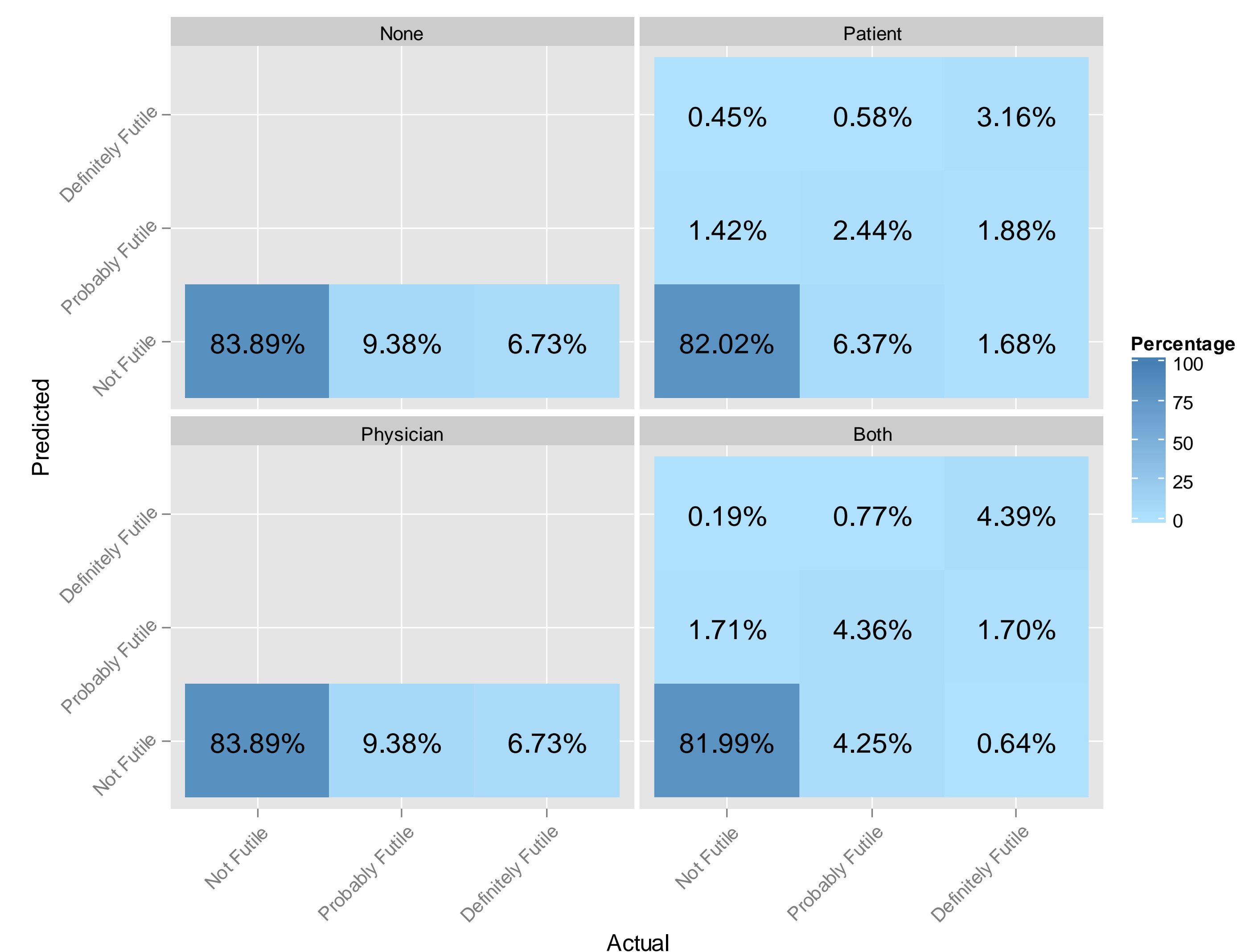
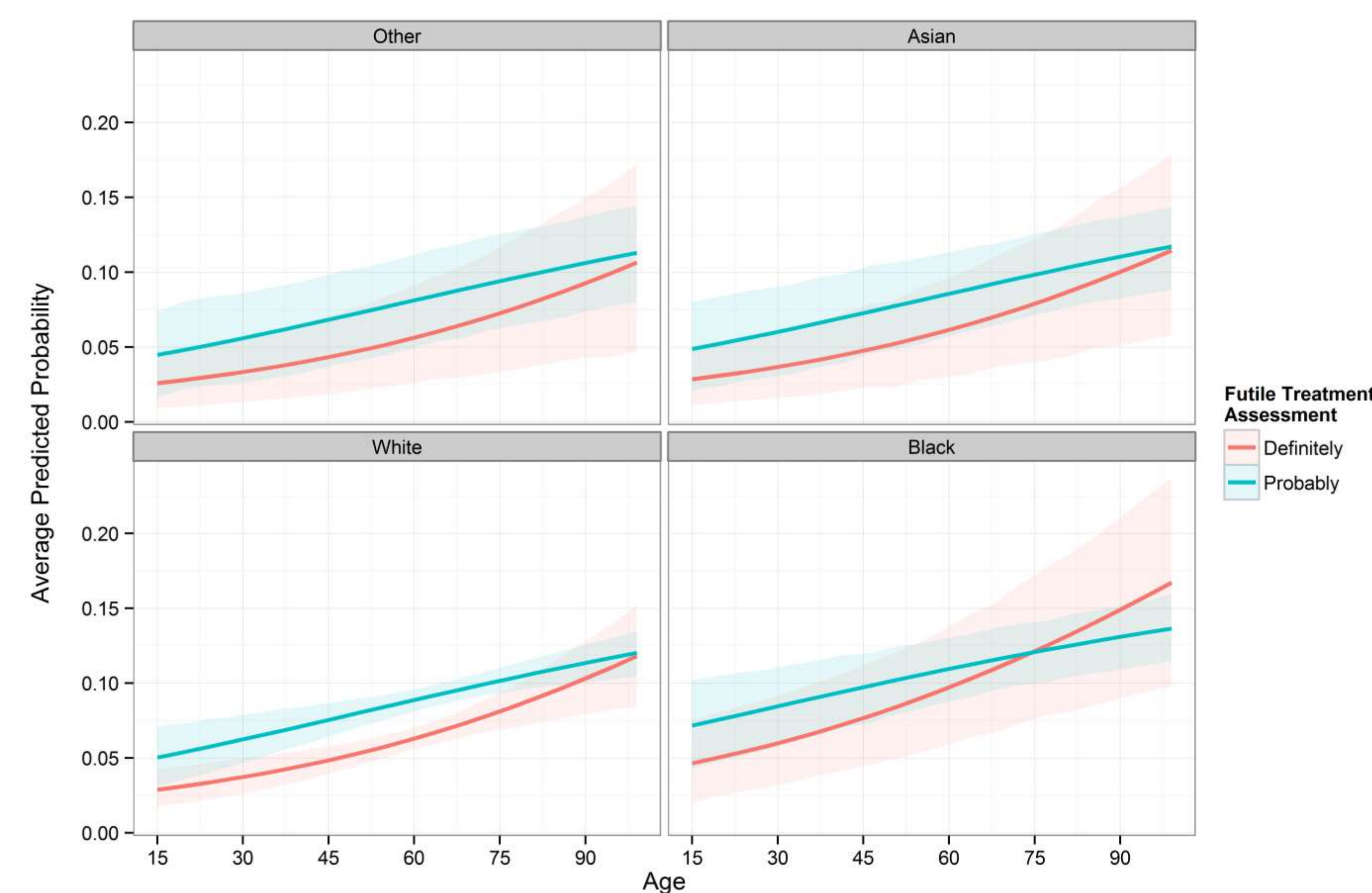
- Predictor effects** are presented as the average marginal change in predicted probability for a one unit change in the predictor with 95% highest posterior density intervals
$$\text{marginal effect} = E \left[\frac{P(Y|X = x + h) - P(Y|X = x)}{h} \right]$$
- Predicted values** are presented as predicted probabilities with 95% HPD intervals
- Model accuracy** is presented as the percentage of correctly predicted futility assessments. Classifications were based on the highest model predicted probability
- Random effects** are often presented as variance of log odds. We start with fixed effects associated with a probability of .5 of not receiving futile treatment. The random effects are added to these and then transformed to probabilities and the standard deviation of these computed

An R package, **postMCMCglmm** was developed to facilitate calculation of each of these measures. It is available from <https://github.com/jwiley/postMCMCglmm>

Average Marginal Predicted Probabilities

	Not Futile	Probably Futile	Futile
Age	-.02 [-.03, -.01]	.01 [.0, .01]	.01 [.01, .02]
Other	.02 [-.06, .09]	-.01 [-.04, .03]	-.01 [-.04, .04]
Asian	0 [-.06, .07]	-.00 [-.03, .02]	-.00 [-.04, .04]
Black	-.06 [-.12, .01]	.02 [-.00, .04]	.04 [-.01, .08]

Average Predicted Probabilities by Age and Ethnicity
Shaded regions indicate 95% HPD Intervals



- Top:* Predicted probabilities by race and age. Probabilities were calculated by setting everyone in the sample to each age and race, and then averaging across the probabilities
- Bottom:* Percent correctly and incorrectly classified varying whether patient and physician random effects are included.

Random Effects

For a baseline probability of non futile assessment, of .5 the average between patient variability on the probability scale was SD = .396, 95% HPD [.383, .409]

The average between physician variability on the probability scale was SD = .053, 95% HPD [.045, .061]

Including both random effects, the average variability on probability scale was SD = .399, 95% HPD [.386, .413]

Conclusion

For categorical data, commonly used probit or logit scales are difficult to interpret.

An interpretational difficulty with probabilities is that they are nonlinear, we address this by calculating the *average marginal change*, which is the average effect in the sample.

Drawing from the posterior distribution from bayesian GLMMs, it is conceptually straightforward to calculate point estimates such as average marginal change, or average predicted probabilities, and to provide credible intervals on how precise the estimates are.

For ordinal probit models in R, `recycler()`, `predict2()`, and `stdranef()` from the **postMCMCglmm** package make it easy to compute average marginal probabilities, predicted probabilities, and random effects, respectively, on the probability scale, instead of the default probit scale.

For categorical data, presenting results on the probability scale facilitates interpretation and is more familiar to researchers and practitioners

References

- Hadfield, J. D. (2010). *MCMC Methods for Multi-Response Generalized Linear Mixed Models: The MCMCglmm R package*. Journal of Statistical Software, 33(2), 1-22, URL <http://www.jstatsoft.org/v33/i02/>.
- Huynh, T. N., Kleerup, E. C., Wiley, J. F., Savitsky, T. D., Guse, D., Garber, B. J., & Wenger, N. S. (submitted for review). *The frequency and cost of futile treatment in critical care*.
- R Core Team (2012). *R: A language and environment for statistical computing*. R Foundation for Statistical Computing, Vienna, Austria. ISBN 3-900051-07-0, URL <http://www.R-project.org/>.