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Relative variable importance in Bayesian generalized linear mixed models with applications in quantitative genetics

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ABSTRACT

As one of the most widely used statistical methods, regression models have a fundamental position in statistics. Obtaining inference on the covariates used to model the response is a key part of regression analysis, and often it is desirable to assign the covariates with a *relative importance*. To do so, numerous methods from multiple perspectives exist. Despite this, a consensus has not been reached, and the traditional methods using p -values have created a reproduction crisis in the social and biomedical sciences. Our contribution to remedy this, is to put forth a Bayesian relative variable importance measure. The measure is designed to make researchers more thoroughly interpret the statistical model and its results, rather than blindly following a threshold to draw conclusions.

Our measure, coined *Bayesian Variable Importance* (BVI), is implemented by transferring the logic of more established, frequentist methods, to the Bayesian framework. The BVI method is applicable to generalized linear mixed models (GLMMs) with continuous, Binomial and Poisson distributed responses. The core of the method, is to utilize the relative weights method on the covariates of before fitting a Bayesian GLMM and performing calculations with respect the Bayesian framework. This produces posterior distributions of the relative importance of all covariates present in the model, as well as the estimated distributions of the marginal and conditional R^2 . To make the methodology and measure easily available for researchers across fields, an R package called `BayesianVariableImportance` was made.

Based on the authors previous work for linear mixed models (Arnstad 2024), a simulation study, case studies and a real world application, we have shown that the BVI method is a viable analogue to the existing frequentist methods. The method is able to produce plausible results for GLMMs with a complex covariance structure, while being simultaneously being computationally efficient. Hopefully, the BVI method can be used across various field and help researchers in their work. With relative variable importance being a topic of much interest and active research, recently also in the Bayesian framework, we believe that the BVI method can be further improved in the future.

SAMMENDRAG

Som en av de mest brukte statistiske metodene, har regresjonsmodeller en fundamental posisjon i statistikk. En nøkkeldel av regresjonsanalysen er å skaffe inferens om kovariatene som brukes til å modellere responsvariabelen, og tilegne kovariatene en *relativ viktighet*. For å gjøre dette, eksisterer flere metoder fra ulike perspektiver. Til tross for mange forskjellige metoder, har det ikke blitt oppnådd en konsensus, og den tradisjonelle fremgangsmåten med p -verdier har skapt en reproducerbarhetskrise i samfunns- og biomedisinsk forskning. Vårt bidrag for å bøte på dette, er å legge frem et Bayesiansk mål for relativ variabelviktighet. Dette målet er designet for at forskere skal tolke den statistiske modellen og dens resultater grundigere, i stedet for å slå seg til ro med konklusjoner basert på en forhåndsbestemt terskel.

Vårt mål, døpt *Bayesiansk Variabel Viktighet* (BVV), er implementert ved å overføre logikken fra mer etablerte, frekventistiske metoder, til det Bayesianske rammeverket. BVV er anvendbart på generaliserte lineære blandingsmodeller (GLBM) som har kontinuerlige, binomiske og Poisson fordelte responser. Kjernen i metoden er å benytte relativ vektning på kovariatene før en Bayesiansk GLBM blir konstruert. Dette produserer posteriore fordelinger av den relative viktigheten til alle kovariatene i modellen, samt de estimerte fordelingene til den marginale og betingede R^2 . For å gjøre metodikken og målet lett tilgjengelig for forskere på tvers av fagfelt, ble en R pakke kalt `BayesianVariableImportance` laget.

Basert på forfatterens tidligere verk Arnstad (2024) for lineære blandingsmodeller, en simulasjonsstudie, case studier og en anvendelse på reelle data, har vi vist at BVV metoden er en levedyktig analog til eksisterende frekventistiske metoder. Metoden er i stand til å produsere plausible resultater for GLBM med komplekse kovariansstrukturer, samtidig som den er beregningsmessig effektiv. Forhåpentligvis kan BVV metoden bli brukt på tvers av ulike fagfelt og hjelpe forskere i deres arbeid. Med tanke på at relativ variabelviktighet er et område av stor interesse og aktiv forskning, også i det Bayesiansks rammeverket, tror vi at BVV metoden kan bli ytterligere forbedret i fremtiden.

PREFACE

This master's thesis concludes the Master of Science degree obtained from the program Physics and Mathematics, with a specialization in Industrial Mathematics, at the Norwegian University of Science and Technology (NTNU). In combination with the project thesis (Arnstad 2024), the master's thesis constitutes 45ECTS, and has been developed during the spring of 2024.

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CONTENTS

Abstract	i
Sammendrag	ii
Preface	iii
Contents	iv
List of Figures	vii
List of Tables	ix
1 Introduction	1
2 Theory	7
2.1 Linear regression	7
2.1.1 Linear regression	7
2.1.2 Qualitative covariates	8
2.1.3 Correlation among covariates in linear regression	8
2.2 Variable importance in linear regression models	9
2.2.1 Relative importance measures	9
2.2.2 Naive decompositions of R^2	10
2.2.3 The Lindemann, Merenda and Gold(LMG) method	11
2.2.4 Relative weights method	12
2.3 Extensions of the linear regression model	13
2.3.1 Generalized linear models (GLMs)	13
2.3.2 Linear mixed models (LMMs)	15
2.3.3 Generalized linear mixed models(GLMMs)	16
2.4 Extending R^2 to GLMMs	16
2.4.1 R^2 for GLMs	17
2.4.2 R^2 for LMMs and random slope models	18
2.4.3 R^2 for GLMMs	21
2.4.4 Extensions of the LMG and relative weights method	22
2.5 The Bayesian framework	22
2.5.1 General idea	23
2.5.2 Prior and posterior distributions	23
2.5.3 Penalising complexity (PC) priors	23

2.5.4	Hierarchical Bayesian modelling	24
2.5.5	R^2 in the Bayesian framework	25
2.5.6	Variable importance measures in the Bayesian framework	26
2.6	The INLA framework	28
2.6.1	Introduction to INLA	28
2.6.2	Approximating the marginals	29
2.6.3	Parameter estimation and sampling procedure	30
2.7	Quantitative genetics and relative variable importance	31
2.8	The Animal Model as a Gaussian Markov Random Field	33
3	Methods	35
3.1	Variable importance in the Bayesian framework	35
3.2	Extending the R^2 to Bayesian GLMMs	36
3.3	Decomposing the R^2 value	37
3.3.1	Applying the relative weights method in the Bayesian frame-work	37
3.3.2	Random effects	38
3.3.3	Drawing samples	38
3.4	Gaussian simulation study	39
3.5	Heritability of phenotypic traits	40
3.5.1	Heritability as relative variable importance	41
3.5.2	House sparrow study	41
3.6	Non-Gaussian studies	42
3.6.1	Binomial and Poisson simulation studies	42
3.6.2	Binomial and Poisson case studies	45
3.7	Simulation study with R^2 -induced Dirichlet decomposition priors and Generalized Decomposition Priors on R^2	46
4	Results	49
4.1	Gaussian simulation study	49
4.1.1	Relative importance of the fixed effects	49
4.1.2	Relative importance of the random effects	52
4.1.3	Total variance explained - R^2 estimates	52
4.2	Heritability of house sparrow traits	54
4.3	Non-Gaussian simulation study	57
4.3.1	Binomial simulation	58
4.3.2	Poisson simulation	62
4.4	Comparison with <code>rptR</code> package	67
4.5	Comparing the BVI method with R^2 -induced Dirichlet decomposi-tion priors and Generalized Decomposition Priors on R^2	69
5	Discussion & Further work	73
6	Conclusions	81
	References	83
	Appendices	89

A GitHub repository	89
B Bayesian Variable Importance usage	90
C Supplementary Material	98
D Miscellaneous proofs	105

LIST OF FIGURES

2.1	Illustration of a pedigree as a GMRF	34
4.1	Relative importance of the fixed effects in Gaussian LMM	51
4.2	Relative importance of the random effect α in Gaussian LMM	52
4.3	Marginal and conditional R^2 in Gaussian LMM	53
4.4	Estimated heritability of body mass from grid and CCD strategy	55
4.5	Estimated heritability of wing length	56
4.6	Estimated heritability of tarsus length	57
4.7	Relative importance of the fixed effects in Binomial GLMM	59
4.8	Relative importance of the random effect α in Binomial GLMM	60
4.9	Marginal and conditional R^2 in Binomial GLMM	62
4.10	Relative importance of the fixed effects in Poisson GLMM	64
4.11	Relative importance of the random effect α in Poisson GLMM	65
4.12	Marginal and conditional R^2 in Poisson GLMM	66
4.13	Estimated repeatability of color in male beetles	68
4.14	Estimated repeatability of eggs laid by female beetles	69
4.15	Comparison of the relative importance from the BVI method and the shrinkage prior methods	70
4.16	Comparison of the marginal R^2 from the BVI method and the shrinkage prior methods	71
C.1	Posterior relative importance distributions of all fixed effects in body mass model for house sparrow study	98
C.2	Posterior relative importance distributions of all random effects in body mass model for house sparrow study	99
C.3	Posterior distributions of R^2 values in body mass model for house sparrow study	99
C.4	Posterior relative importance distributions of all fixed effects in wing length model for house sparrow study	100
C.5	Posterior relative importance distributions of all random effects in wing length model for house sparrow study	100
C.6	Posterior distributions of R^2 values in wing length model for house sparrow study	101
C.7	Posterior relative importance distributions of all fixed effects in tar- sus length model for house sparrow study	101

C.8	Posterior relative importance distributions of all random effects in tarsus length model for house sparrow study	102
C.9	Posterior distributions of R^2 values in tarsus length model for house sparrow study	102

LIST OF TABLES

3.1	Distribution-specific variance σ_d^2 for the Gaussian, Binomial and Poisson distributions	37
3.2	Expected R^2 for Gaussian LMM	40
3.3	Expected relative importance of independent covariates for non-Gaussian GLMMs	44
3.4	Expected R^2 for non-Gaussian GLMMs	45
3.5	Expected R^2 for comparison of BVI and shrinkage prior methods . .	46
4.1	Heritability estimates and confidence intervals	54
C.1	Summary statistics for binomial GLMM simulation study	103
C.2	Summary statistics for Poisson GLMM simulation study	104

CHAPTER ONE

INTRODUCTION

Statistics as a mathematical field has a long history as a tool for characterizing social, economic and scientific phenomena. One of the most used statistical methods is regression analysis (Grömping 2015), which is used to model the relationship between a response variable and one or more covariates. To understand this relationship, researchers often want to determine whether a covariate is associated with the response, and to what extent. The exploration of this fundamental question has lead to a number of statistical methods trying to answer it. An agreement on a single method has not been reached, with the topic still being debated and actively researched.

A pioneer in statistics, Ronald Fisher, introduced the concept of the *p*-value almost one hundred years ago (Fisher 1925). To this day, the *p*-value is arguably the most widespread and used method, to determine if a covariate is *statistically significant* with respect to the response. Popularly, to determine statistical significance, a hypothesis test is performed and the resulting *p*-value is compared to a threshold level α . Typically, if the *p*-value is smaller than α , the covariate is considered statistically significant. However, this way of determining statistical significance is very prone to misinterpretations, and is subject to great criticism (Benjamin et al. 2018).

Recently, the social and biomedical sciences have been subject to a reproducibility crisis, in which published results cannot be reproduced (Blakeley B. McShane & Gelman 2019). One possible solution is suggested by 72 authors in Benjamin et al. (2018), which is to lower the typical significance level from $\alpha = 0.05$ to $\alpha = 0.005$. This could be a solution, however Blakeley B. McShane & Gelman (2019) sees this as a quick fix, which will not solve the underlying problem. Instead, it is proposed to simply abandon the term statistical significance, and not force results to be based on a threshold value which gives rigid and binary answers. The remedy, according to Blakeley B. McShane & Gelman (2019), is to rather interpret the *p*-value as a continuous measure of evidence, among many others. Going forward, the thesis will not consider statistical significance but rather statistical evidence to avoid the hazards by using such rigid interpretations.

There exists many other measures to compliment the *p*-value when assessing a

statistical model. As in Arnstad (2024), we list some of the most common measures

- **Effect sizes:** By looking at the squared value of the standardized regression coefficients, one can determine the effect size of the covariates. For independent covariates, the effect size coincides with the proportion of variance explained by the covariate. The effect sizes are a good measure for uncorrelated covariates, but falls short when correlation makes the coefficient estimates unstable.
- **Confidence intervals:** Confidence can be calculated for the effect sizes, and the interval can be used to determine a range of values that can be seen as statistically consistent with the data. The downside to using confidence intervals is that one effectively relies on the p -value to determine it, and therefore it does not provide any additional information than using the p -value.
- **Information criteria:** The AIC (Akaike 1974) and the BIC (Schwarz 1978) are information criteria that use the likelihood function to compute a goodness of fit statistic. These can be used to assess the information contained in the model, and can be used to assess the unique information that one covariate contributes to the model. Again, the criteria fall short when correlation is present, as it cannot take shared information between covariates into account.
- **Bayes factor:** As an alternative to the p -value, Bayes factor can be used to assess the evidence from the data to either support a hypothesis or not. Bayes factor is therefore less rigid than the p -value, and is rather a continuous measure of evidence.
- **Decomposing the R^2 :** The R^2 measures the variance explained in the response by the covariates. As such, it is a goodness of fit measure and can be decomposed into a share from each covariate in the model to determine the variance explained by each covariate. The R^2 is widely used and intuitive, but a proper decomposition of the value with correlated covariates is not straightforward.

The methods listed all have in common that they become less reliable for correlated covariates. As correlation is a common feature of many datasets, especially from the real world, this is a general problem that regression models are not well suited to handle (Grömping 2015).

By decomposing the R^2 value and assigning each variable with a share of explained variance in the response, we have a measure of the relative importance of each covariate (Grömping 2007). The problem of decomposing the R^2 in a sensible manner has lead to various approaches. One of the most rigorous methods is the approach by Lindemann, Merenda and Gold (LMG) (Lindeman et al. 1980), which decomposes the R^2 value by considering all possible orderings of the covariates. As covariates are added to the null model, the average increase in R^2 for each permutation is calculated and each covariate is assigned a share of relative

variable importance. As the LMG method is very popular, it has been applied in dominance analysis (Budescu 1993) and coincides with the Shapley value in game theory (Shapley (1953), Lipovetsky & Conklin (2001)). The LMG method has proven to be consistent for the linear regression, but is computationally expensive and therefore in some cases not feasible.

The relative weights method (Johnson (1966), Fabbris (1980), Genizi (1993)) can be seen as an approximation of the LMG method, to remedy the computational burden. By projecting the covariates into an orthogonal space and then conducting the analysis on the projected covariates, before transforming them back to the original covariate space, the relative weights method efficiently decomposes the R^2 value. At the cost of approximating rather than being more rigorous, the relative weights method is able to handle larger models and is therefore preferred if the LMG is not feasible (Grömping 2007).

Both the LMG method and the relative weights method are originally designed to decompose the R^2 for the linear regression. However, for many scenarios, a linear regression is not sufficient to model the relationship between the response and the covariates. Both models were extended in Matre (2022), so that they could be applicable for the random intercept models, but we are not aware of any further extensions. Another problem is that there is little consensus on how to properly define the R^2 for more complex methods. A simple and intuitive definition of the R^2 for more general regression models was suggested by Nakagawa & Schielzeth (2013) and serves as the basis for the extensions of the LMG and relative weights method.

The Bayesian framework has significant advantages when compared to the frequentist framework (Robert 2007) and has had a surge in popularity due to the recent years advancements in computational capabilities (Hackenberger 2019). The treatment of parameters as random variables in the Bayesian framework, rather than point estimates, naturally includes moments such as variance in the posterior distribution. As the Bayesian framework has become more available, researchers can obtain more inference and thereby make more informed decisions. Therefore, we believe that the Bayesian framework provides a better platform for assessing the statistical evidence of covariates. One important application of statistical analysis is the decomposition of variance in regression models, a practice that has been common in various fields for a long time. For instance, in quantitative genetics, researchers often aim to determine the heritability of phenotypic traits. Heritability is a key measure used to explain how the mean value of a trait changes, and can thereby help us better understand evolution. Although point estimates of heritability are fairly straightforward to obtain, it is considerably more difficult to evaluate the uncertainty of the estimates. We believe that the Bayesian framework is the natural choice for quantifying this uncertainty, and this particular example has been a great motivation behind our attempt to develop a Bayesian relative variable importance measure.

In the Bayesian framework, the field of relative variable importance is small. The LMG and relative weights method are both based on the frequentist framework,

and the Bayesian framework has not been explored to the same extent. One possible line of action are the Generalized decomposition priors on R^2 (GDR2) (Aguilar & Bürkner 2024), which are based on the R^2 -induced Dirichlet decomposition (R2D2) priors (Zhang et al. 2020). By placing a prior on the R^2 value, the R2D2 priors proceed with a Dirichlet decomposition of the R^2 value to be able to assign each covariate with a share of relative variable importance. The GDR2 priors are a generalization of the R2D2 priors which performs the decomposition using logistic normal distributions (Aguilar & Bürkner 2024). At the time being, the R2D2 and GDR2 priors have been applied only to the linear regression, with a focus on obtaining trustworthy predictions. Therefore, they have not been used explicitly as relative importance measures, and it is not clear how they generalize to more complex models. Nonetheless, they serve as an important contribution to the Bayesian framework for relative variable importance.

To summarize, the field of relative variable importance offers a wide range of methods to determine the statistical evidence of covariates in the simple linear regression model. However, correlation between the covariates is troublesome and can lead to unreliable results. Further, for more complex models, there is little consensus on how computations should be carried out and therefore a lack of robust methods. Many of the existing methods have been developed for specific purposes and do not pose a general method for relative variable importance. In the Bayesian framework, there has been done little work on relative variable importance, despite its advantageous features for statistical inference.

This thesis aims to develop a relative variable importance measure in the Bayesian framework, which can be easily implemented by researchers in different fields. The measure should be applicable to a wide range of regression models with complex structures, with an emphasis on interpretable and reliable results. The method will use the relative weights method as a basis, and apply the method in the Bayesian framework. To calculate the relative variable importance measures, the method will merge the R^2 definition for GLMMs of (Nakagawa & Schielzeth 2013) and (Nakagawa et al. 2017), with the Bayesian R^2 for the linear regression model defined by Gelman et al. (2017). The results obtained will be in the form of approximated posterior distributions, allowing for uncertainty in the estimates to be quantified. Our hope is that these distributional results will be easy to interpret and allow researchers to interpret the uncertainty, rather than using a threshold based method. For the method to be easily used, it has been implemented as a package in R. The package is called `BayesianVariableImportance` and is available, along with installation and usage examples, on the authors GitHub <https://github.com/AugustArnstad/BayesianVariableImportance>.

As regression models are perhaps the most used statistical modelling tool, the span of applications for the BVI method is wide. Given that the regression model is a suitable choice, the BVI method can be useful in fields such as biology, economics, social sciences, medicine and more. Research in many of these fields is crucial if we are to reach the 17 goals for sustainability set by the United Nations (United Nations 2023). We highlight the application of the BVI method for use in quantitative genetics. The overarching question for such analysis is to better un-

derstand evolution and how species develop when subject to different and changing environments. Answers to such questions is very useful to reach goal 14 and 15 of the United Nations sustainability goals, which is about life in the ocean and on land. The aim of goal 14 is to *Conserve and sustainably use the oceans, seas and marine resources for sustainable development* and goal 15 is about *Protecting, restoring and promoting sustainable use of terrestrial ecosystems, sustainably manage forests, combat desertification, and halt and reverse land degradation and halt biodiversity loss* (United Nations 2023). Further, we believe that the BVI method can give insight that can be useful for numerous of these goals, such as goal 3 about *Good health and well-being*, goal 7 about *Affordable and clean energy* and goal 13 about *Climate action*.

The structure of the thesis is as follows. In Chapter 2 we look at some background theory and put forth the theoretical results that will be used in the method. To describe our calculations, Chapter 3 presents the methodology and logic of our contribution. Evaluating the method is done in Chapter 4, where we look at a simulation study, a case study and apply the method to a real dataset. We discuss the findings in Chapter 5 and conclude the thesis in Chapter 6. In Appendix A we give the link to the authors GitHub repository for the R package and the thesis, Appendix B contains a usage example of the R package and Appendix C has some supplementary material. A miscellaneous proof is found in Appendix D.

Please note that this thesis continues the work done by the author in Arnstad (2024), and therefore some sections overlap. Following the guidelines of the Institute of Mathematical Sciences, stating that sections need not be rewritten, some sections are the same, or slightly modified, as in the project thesis.

CHAPTER
TWO

THEORY

2.1 Linear regression

All regression models are based on the assumption that the response variable is influenced by one or more covariates. The relationship between the response and the covariate is assumed not to be deterministic, so we expect our modelling of the response to be influenced by some random error (Fahrmeir et al. 2013). This means that the response is treated as a random variable, and it is desirable to decompose the response into systematic components and random components.

2.1.1 Linear regression

Assuming that an observed response y_i has a linear relationship with a covariate x_i is the basis for the simple linear regression. This can be modeled by the equation

$$y_i = \beta_0 + \beta_1 x_i + \varepsilon_i , \quad (2.1)$$

where β_0 is the intercept, β_1 is the slope, and ε_i is the error term. The error term, or residuals, is assumed to be normally distributed with mean zero and variance σ^2 , i.e. $\varepsilon_i \sim \mathcal{N}(0, \sigma^2)$. Generalizing to multiple covariates is straightforward by defining the $n \times p$ matrix \mathbf{X} as a design matrix with, including an intercept, the p covariates in the columns and the n observations in the rows. With this definition, the linear regression model can be written as

$$\mathbf{y} = \mathbf{X}\boldsymbol{\beta} + \varepsilon , \quad (2.2)$$

where now $\mathbf{y} = (y_1, y_2, \dots, y_n)$ is a vector of n responses, $\boldsymbol{\beta} = (\beta_0, \beta_1, \dots, \beta_{p-1})$ is a vector of coefficients including the intercept β_0 , and $\varepsilon = (\varepsilon_1, \varepsilon_2, \dots, \varepsilon_n)$ is a vector of error terms. The error terms are assumed to be independent and identically distributed (i.i.d.) with $\varepsilon \sim \mathcal{N}(0, \sigma^2 \mathbf{I})$, where \mathbf{I} is the identity matrix of size $n \times n$. Consequently, the response \mathbf{y} is conditionally independent given the covariates \mathbf{X} , i.e.

$$\mathbf{y} | \mathbf{X} \sim \mathcal{N}_n(\mathbf{X}\boldsymbol{\beta}, \sigma^2 \mathbf{I}) . \quad (2.3)$$

In practice, the coefficients $\boldsymbol{\beta}$ are estimated from the maximum likelihood estimation (MLE) method, given by

$$\hat{\boldsymbol{\beta}} = (\mathbf{X}^T \mathbf{X})^{-1} \mathbf{X}^T \mathbf{y} . \quad (2.4)$$

2.1.2 Qualitative covariates

In many cases the covariates are qualitative, meaning they are categorical variables that can be grouped into different levels or factors. Qualitative covariates, unlike quantitative, cannot be measured numerically, and we must adjust our modelling to account for this. A common approach to model qualitative data is to include dummy variables, which are assigned a value 1 if the observation is in the respective category(factor) and 0 otherwise. Given N factors, it is standard practice to model $N - 1$ dummy variables and let one factor be captured by the intercept to uniquely determine the model. Dummy encoding in this way retains the properties of the linear regression, and are limited by the same assumptions. The model for the response y_i , assuming no quantitative covariates, from group j with dummy encoding is then given by

$$y_i = \beta_0 + \sum_{j=1}^{N-1} \beta_j x_{i,j} + \varepsilon_i , \quad (2.5)$$

where β_j denotes the factor coefficient of observation i and the dummy variable

$$x_{i,j} = \begin{cases} 1 & \text{if observation } i \text{ is in group } j \\ 0 & \text{otherwise} \end{cases} . \quad (2.6)$$

This way of modelling qualitative covariates is intuitive and easy to interpret, but it also assumes that factor specific effects are uniform and fixed across all levels and becomes cumbersome with many categorical covariates.

2.1.3 Correlation among covariates in linear regression

Correlation among covariates is to be expected, as it is natural in many scenarios. However, if the correlation is very strong, this poses some serious problems when interpreting the linear regression model. The covariates \mathbf{x}_i in a linear regression are assumed to be linearly independent, so that the design matrix \mathbf{X} has full rank. If the design matrix is not of full rank, that is one or more covariates are perfectly correlated, the model (2.2) is said to be *multicollinear* (Poole & O'Farrell 1971). From equation (2.4) one can see that if the matrix \mathbf{X} is not of full rank, the term $(\mathbf{X}^T \mathbf{X})^{-1}$ is not invertible and the MLE of $\boldsymbol{\beta}$ does not exist. Further, the variance of the MLE of $\boldsymbol{\beta}$ grows as the correlation between covariates grows (Fahrmeir et al. 2013, p. 116). A larger variance in $\hat{\boldsymbol{\beta}}$ also leads to larger standard errors and larger p -values for $\hat{\boldsymbol{\beta}}$, making it hard to assess the model. Both coefficients and covariates affect the total marginal model variance, which can be decomposed as

$$\text{Var}(\mathbf{y}) = \text{Var}(\mathbf{X}\boldsymbol{\beta}) + \text{Var}(\boldsymbol{\varepsilon}) = \boldsymbol{\beta}^T \mathbf{V} \boldsymbol{\beta} + \sigma_\varepsilon^2 = \sum_{j=1}^p \beta_j^2 v_j + \sum_{j=1}^{p-1} \sum_{k=j+1}^p \beta_j \beta_k \sqrt{v_j v_k} \rho_{jk} + \sigma_\varepsilon^2 , \quad (2.7)$$

(Grömping 2007) where $\mathbf{V} = \text{Cov}(\mathbf{X})$ is the $p \times p$ covariance matrix of the covariates which is assumed to be positive definite, $\boldsymbol{\beta}$ is the $p \times 1$ vector of regression coefficients, v_j the regressor variances for $j = 1, \dots, p$ found along the diagonal of \mathbf{V} and ρ_{jk} the inter-regressor correlations between regressor j and k . The middle term in 2.7 consist of the covariance between the covariates and the variance contribution from a single covariate is not immediately clear.

2.2 Variable importance in linear regression models

In a regression setting with multiple regression coefficients, it is often desirable to be able to assign each covariate with a measure of its relative importance with respect to the model. The relative importance of covariate \mathbf{x}_i is defined as the standardized contribution to explained variance in the response \mathbf{y} from \mathbf{x}_i (Grömping 2007). Assigning relative importance is no trivial task, as correlation among covariates poses a challenge in assessing the relative importance of each covariate.

2.2.1 Relative importance measures

The coefficient of determination, R^2 , is a widely used and intuitive summary statistic of goodness-of-fit and can also be used in model comparison. Conceptually, the R^2 quantifies how much variance in the response variable can be attributed to the covariates in the model. For the linear regression model, the R^2 is defined as

$$R^2 = 1 - \frac{(\mathbf{y} - \mathbf{X}\boldsymbol{\beta})^T(\mathbf{y} - \mathbf{X}\boldsymbol{\beta})}{(\mathbf{y} - \bar{\mathbf{y}})^T(\mathbf{y} - \bar{\mathbf{y}})} = \frac{\text{Var}(\mathbf{y}) - \sigma_\varepsilon^2}{\text{Var}(\mathbf{y})}, \quad (2.8)$$

where $\bar{\mathbf{y}}$ is the mean vector of responses \mathbf{y} . Instead of referring to the R^2 value alone, going forward this thesis will focus on decomposing of the R^2 value and allocate a proportion of R^2 to the model covariates. This decomposition is done in order to assess the relative importance, or variance explained, of each covariate in the model. The special case of uncorrelated covariates in \mathbf{X} gives

$$\text{Var}(\mathbf{y}) = \sum_{j=1}^p \beta_j^2 v_j + \sigma_\varepsilon^2, \quad (2.9)$$

and provides a natural decomposition of the R^2 in terms of contribution from each covariate, as each predictor \mathbf{x}_i contributes $\beta_i^2 v_i$ to the total response variance (Grömping 2007). In (2.7) however, the response variance is split into three parts, the first two sums which comes from the regressors and the latter term which is the variance of the error. As mentioned, it is the middle term that poses the problem of assigning importance to each covariate, since it is not immediately clear how to distribute the total response variance to each covariate, as some variance contributions in the response variance are shared among covariates. The literature has established some conditions that relative importance measures should fulfill, so that they can be interpreted and compared in a sensible manner (Grömping 2007). As listed in Grömping (2007), the methods should have

1. **Proper decomposition:** The model variance should be decomposed into shares for each regressor that sum up to the total variance, and the method shall allocate the shares to each regressor.
2. **Non-negativity:** Each share of the variance should be non-negative.

3. **Exclusion:** If a regressor is excluded from the model, $\beta_j = 0$, its share of the variance should be zero.
4. **Inclusion:** If a regressor is included in the model, $\beta_j \neq 0$, its share of the variance should be positive.

Further, Grömping (2007) introduces some useful notation for considering variable importance, for the explained variance of a subset of regressors S as

$$\text{evar}(S) = \text{Var}(\mathbf{y}) - \text{Var}(\mathbf{y}|\mathbf{x}_j, j \in S), \quad (2.10)$$

and the added explained variance of a subset M to a model that already contains the covariates in S as

$$\text{svar}(M|S) = \text{evar}(M \cup S) - \text{evar}(S). \quad (2.11)$$

2.2.2 Naive decompositions of R^2

To make it clear that some simple decompositions fail the conditions of relative importance measures, we will consider two naive approaches for decomposing the R^2 . We denote the R^2 of a linear regression with regressors X_1, \dots, X_p as $R^2(\{1, \dots, p\})$ and the relative importance of regressor X_i as $\text{RI}(\{i\})$.

The first naive method is to fit a model with all regressors p , and then fit a model with all regressors excluding regressor i . The relative importance of X_i is then the difference $R^2(\{1, \dots, p\}) - R^2(\{1, \dots, p\} \setminus i)$. To show how this fails the conditions of relative importance measures, an example from Matre (2022) is discussed. The example considers the simple case

$$Y = X_1 + X_2, \quad \text{Var}(X_1) = \text{Var}(X_2) = 1, \quad \text{Cov}(X_1, X_2) = 0.9. \quad (2.12)$$

The R^2 of the model with both covariates is $R^2(\{1, 2\}) = 1$, since the covariates X_1, X_2 explain fully the response Y . Then one would expect that the importance of X_1 and X_2 is 0.5 each, since they both explain half of the response variance. Using the proposed decomposition, one would calculate

$$\text{RI}(\{2\}) = R^2(\{1, 2\}) - R^2(\{1\}) = 1 - \frac{\text{Cov}(Y, X_1)^2}{\text{Var}(Y)\text{Var}(X_1)} = 1 - \frac{1.9^2}{3.8} \approx 0.05, \quad (2.13)$$

where it is used that for the simple linear regression, the R^2 is given by the squared correlation coefficient between the response and the regressor. By symmetry $\text{RI}(\{1\}) = \text{RI}(\{2\})$, so the sum of the relative importances is 0.1. However, the total explained variance of the model is 1, so this decomposition violates the proper decomposition condition. This decomposition only assign importances to the regressor based on the information that the regressor does not share with any other regressors. Therefore, it does not take into account the shared information and the importance estimated is too low.

Another naive decomposition would be to compare the relative importance of

a model with one regressor i to the empty model, *i.e.* the model with no covariates. The empty model has an $R^2 = 0$ and therefore for X_1 in the above example we would have

$$\text{RI}(\{1\}) = R^2(\{1\}) - R^2(\{\emptyset\}) = \frac{\text{Cov}(Y, X_1)^2}{\text{Var}(Y)\text{Var}(X_1)} = \frac{1.9^2}{3.8} \approx 0.95. \quad (2.14)$$

Once more by symmetry we have $\text{RI}(\{2\}) = \text{RI}(1)$, so the sum of the relative importances is 1.9, violating the proper decomposition condition. In contrast to the first naive approach, this decomposition assigns importances based on the full information contained in the regressor. Therefore, it overestimates the importance of each variable, since the shared information is accounted for twice.

As we have seen from these naive approaches, the task of decomposing the R^2 value is far from trivial, and calls for more sophisticated methods.

2.2.3 The Lindemann, Merenda and Gold(LMG) method

A method that handles correlation among covariates, and is frequently reinvented (Grömping 2007) from different approaches, is the Lindemann, Merenda and Gold (LMG) method. We shall therefore discuss it, as it serves an important role as a leading method for assigning relative variable importance. The LMG method takes use of averaging over orders, meaning that it permutes the index set $\{1, \dots, p\}$ of the regressors $(p - 1)!$ times, excluding the intercept, and sequentially adds the regressors to the model for each permuted index set. By adding regressors sequentially for each permutation, one can investigate how the importance of the regressors vary depending on what other regressors are included, which is useful when they are correlated. This is justified by the assumption that there is no relevant ordering of the regressors in the index set (Kruskal 1987). For each regressor added, starting with none, it allocates a share of explained variance, or importance, and then adds a new regressor. The final allocated share to the regressor is the average of the allocated shares to that regressor for all permutations of the set of regressors indices. This would mean that for two correlated regressors whose importance share varies depending on which is added first, would receive an averaged importance. Averaging over orders is a statistical tradition (Kruskal 1987) and gives a robust assessment of each regressor's importance by considering different orderings of how they are added to the model. The iterative process for the regressors $\{X_0, X_1, X_2, X_3\}$, where X_0 is the intercept, would be

1. Considering $\{X_1, X_2, X_3\}$, X_1 is added to the model, and the share of explained variance allocated to X_1 is $\text{svar}(\{1\}|\emptyset)$. X_2 is added and allocated a share of $\text{svar}(\{2\}|\{1\})$, and lastly X_3 is added and allocated a share of $\text{svar}(\{3\}|\{1, 2\})$.
2. Considering $\{X_1, X_3, X_2\}$, X_1 is added to the model, and the share of explained variance allocated to X_1 is $\text{svar}(\{1\}|\emptyset)$. X_3 is added and allocated a share of $\text{svar}(\{3\}|\{1\})$, and lastly X_2 is added and allocated a share of $\text{svar}(\{2\}|\{1, 3\})$.

The above iteration is repeated for all 6 possible permutations of orderings among regressors to obtain the final result. This iterative process gives rise to the general

formula for share of explained variance allocated to X_1 by the LMG method with p regressors (Grömping 2007),

$$\text{LMG}(1) = \frac{1}{p!} \sum_{S \subseteq \{2, \dots, p\}} n(S)!(p - n(S) - 1)! \text{svar}(\{1\}|S) , \quad (2.15)$$

where $n(S)$ is the number of regressors in S . Equation (2.15) averages the increase in R^2 , $\text{svar}(\{X_i\})$, when adding the covariate of interest, X_i , over all possible orderings of covariates. This mean increase over orderings is assigned as the proportion of R^2 explained by X_i . The LMG method fulfills all but the exclusion criteria described previously (Grömping 2007), but Grömping (2007) argues that this "*must be seen as a natural result of model uncertainty*" and therefore that this criterion is not indispensable. Therefore, we find it also suitable for our purposes to focus on the three other criteria. The setback of the LMG method is the great computational expense that the permutations require when p is large. The complexity is 2^{p-1} summations (Grömping 2007), and therefore, the LMG is not suitable for high dimensional models.

2.2.4 Relative weights method

A method that takes advantage of the straightforward decomposition of the variance when the fixed covariates are uncorrelated is the relative weights method (Johnson 2000), which will now be discussed.

The relative weights method proposes an alternative to the LMG, which is significantly less computationally expensive. Intuitively, the relative weights method projects the design matrix \mathbf{X} of the fixed effects into an orthogonal column space, resulting in a matrix \mathbf{Z} with orthogonal columns. The matrix \mathbf{Z} is then an approximation of \mathbf{X} and will be used as the design matrix in the regression. Since the columns of the design matrix \mathbf{Z} are orthogonal, each covariate is uncorrelated. This allows us to decompose the variance in the straightforward manner as in equation (2.9).

In relative weights one uses the singular value decomposition (Nimon & Oswald 2013), to project the real-valued design matrix \mathbf{X} into an orthonormal matrix $\mathbf{U} \in \mathbb{R}^{n \times n}$ containing the eigenvectors of $\mathbf{X}\mathbf{X}^T$, an $n \times p$ diagonal matrix \mathbf{D} containing the singular values of \mathbf{X} and another orthonormal matrix $\mathbf{V} \in \mathbb{R}^{p \times p}$ containing the eigenvectors of $\mathbf{X}^T\mathbf{X}$ such that

$$\mathbf{X} = \mathbf{UDV}^T . \quad (2.16)$$

From the Eckhart-Young-Mirsky theorem (Mirsky 1960) and following the derivations of Johnson (1966), one can state that the matrix \mathbf{X} , of rank r , can be approximated by a matrix $\mathbf{Z} = \mathbf{UV}^T$ of rank $k \leq r$ such that the difference under the squared Frobenius norm

$$\|\mathbf{X} - \mathbf{Z}\|_F^2 = \text{tr}((\mathbf{X} - \mathbf{Z})^T(\mathbf{X} - \mathbf{Z})) , \quad (2.17)$$

is minimized. The relative weights approximation now utilizes the matrix (Johnson 2000) $\frac{1}{\sqrt{n-1}}\mathbf{Z}$, where the factor $\frac{1}{\sqrt{n-1}}$ is the standardization factor for \mathbf{Z} (Matre

2022), and regresses on \mathbf{Z} to find the MLE $\boldsymbol{\beta}_\mathbf{Z}$ as

$$\begin{aligned}\boldsymbol{\beta}_\mathbf{Z} &= (\mathbf{Z}^T \mathbf{Z})^{-1} \mathbf{Z} \mathbf{y} \\ &= ((n-1) \mathbf{V} \mathbf{U}^T \mathbf{U} \mathbf{V}^T)^{-1} \sqrt{n-1} \mathbf{V} \mathbf{U}^T \mathbf{y} \\ &= \frac{1}{\sqrt{n-1}} \mathbf{V} \mathbf{U}^T \mathbf{y}.\end{aligned}\tag{2.18}$$

As \mathbf{Z} is orthogonal, the relative importance for each column \mathbf{z}_i with respect to the response \mathbf{y} can be found as the square of $\beta_{Z,i}^2$, denoted as $\boldsymbol{\beta}_\mathbf{Z}^{[2]}$. The notation $\boldsymbol{\xi}^{[2]}$ for some $\boldsymbol{\xi}$ represents the Schur product of $\boldsymbol{\xi}$ with itself, *i.e.* element wise squaring of each element in $\boldsymbol{\xi}$. Once these importances are obtained, Johnson (2000) argues that we should regress \mathbf{X} on \mathbf{Z} to obtain the weights that relate the importance of each column of \mathbf{Z} to each column of \mathbf{X} . These weights can be calculated as the matrix

$$\boldsymbol{\Lambda} = (\mathbf{Z}^T \mathbf{Z})^{-1} \mathbf{Z}^T \mathbf{X} = (\mathbf{V} \mathbf{U}^T \mathbf{U} \mathbf{V}^T)^{-1} \mathbf{V} \mathbf{U}^T \mathbf{U} \mathbf{D} \mathbf{V}^T = \mathbf{V} \mathbf{D} \mathbf{V}^T,\tag{2.19}$$

and since \mathbf{Z} is orthogonal, the contribution from a column of \mathbf{z}_i with respect to a column \mathbf{x}_j is the squared entry Λ_{ij}^2 . The contribution from a column \mathbf{x}_j with respect to the response \mathbf{y} , *i.e.* the relative importance, is then estimated as the matrix product (Johnson 2000)

$$\text{RI}(\mathbf{X}) = \boldsymbol{\Lambda}^{[2]} \boldsymbol{\beta}_\mathbf{Z}^{[2]},\tag{2.20}$$

with RI as a column vector where each entry j contains the estimate of the relative importance corresponding to column j of \mathbf{X} . In Matre (2022, section 2.5.3) it is shown that the relative weights method fulfills the criteria same three criteria as the LMG method, because \mathbf{Z} and \mathbf{X} are linear combinations of each other and due to the properties of $\boldsymbol{\Lambda}$.

2.3 Extensions of the linear regression model

The linear regression model is a popular tool in many sciences, but it has limitations when one wants to model more complex structures between the response and covariates. We now generalize the concept of linear regression to be able to model more complex data structures.

2.3.1 Generalized linear models (GLMs)

The first step in expanding the linear regression model, is to allow the responses to be non-Gaussian. Instead of considering only the normal distribution as the distribution of the response, one can consider general responses belonging to the exponential family. Assume that each we have N observations of the response y_i , where $i = 1, \dots, N$, that are conditionally independent given the fixed effects. Then, y_i belongs to the univariate exponential family if

$$f(y_i | \theta_i, \phi) = \exp \left(\frac{(y_i \theta_i - b(\theta_i))}{a(\phi)} + c(y, \phi) \right),\tag{2.21}$$

for some functions $a(\cdot)$, $b(\cdot)$ and $c(\cdot)$, where θ_i is the parameter of the distribution, ϕ is a dispersion parameter and θ_i is a canonical parameter if ϕ is known (McCullagh & Nelder 1989). It is required that the function $b(\cdot)$ is twice differentiable, that the density function $f(y_i|\theta_i, \phi)$ is normalizable and that the support of $f(y_i|\theta_i, \phi)$ is not dependent on θ . Two key properties, expectation and variance, of the exponential family are given by

$$\begin{aligned}\mathbb{E}(Y|\theta) &= b'(\theta) \\ \text{Var}(Y|\theta) &= a(\phi)b''(\theta),\end{aligned}\tag{2.22}$$

where $b''(\theta)$ may also be referred to as the variance function (Fahrmeir et al. 2013) we have left out indexing, and a proof can be found in Appendix D. In the canonical form, the parameter θ_i coincides with the linear predictor η_i defined as

$$\theta_i = \eta_i = \mathbf{x}_i^T \boldsymbol{\beta}.\tag{2.23}$$

To connect the linear predictor η_i to the response, we define a monotonic, differentiable link function $g(\cdot)$ such that

$$\eta_i = g(\mu_i) = g(\mathbb{E}(Y_i)).\tag{2.24}$$

For normally distributed responses, one typically uses the identity function as the link function, which yields the linear regression model. If one considers a binary response, the perspective changes. In a binary regression, one wishes to analyse how the covariates influence the probability

$$\pi_i = \mathbb{P}(y_i = 1|\mathbf{x}_i) = \mathbb{E}[y_i].\tag{2.25}$$

This requires that $\mathbb{E}[y_i]$ lies in the interval $[0, 1]$ as it represents a probability measure. Therefore, the inverse of the link function must transform the linear predictor in such a way that the expectation fulfills this criterion (Fahrmeir et al. 2013). A popular choice of inverse link function is the logistic response function

$$\pi_i = h(\eta_i) = \frac{\exp(\eta_i)}{1 + \exp(\eta_i)},\tag{2.26}$$

yielding the logit-link function

$$g(\pi_i) = \log\left(\frac{\pi_i}{1 - \pi_i}\right) = \eta_i,\tag{2.27}$$

which will be further investigated later on. An intuitive interpretation of the coefficients can be made by noticing that odds

$$\frac{\pi_i}{1 - \pi_i} = \exp(\eta_i) = \exp(\beta_0) \exp(\beta_1 x_{1,i}) \dots \exp(\beta_p x_{1,p}),\tag{2.28}$$

is affected by the covariates in an exponential-multiplicative form (Fahrmeir et al. 2013). Another common regression type is regressing on count data. The most common way of modelling count data is by using the Poisson distribution, which assumes that the events occurring in a time interval or spatial region follow a Poisson process (McCullagh & Nelder 1989). The count of how many events y_i that happen in this time interval or region is said to follow a Poisson distribution

with some rate $\lambda_i = \mathbb{E}[y_i]$. As the number of events occurring cannot be negative, the rate is also restricted to positive values. The common choice of inverse link function is therefore

$$\lambda_i = \exp(\eta_i) = \exp(\beta_0) \exp(\beta_1 x_{1,i}) \dots \exp(\beta_p x_{1,p}) , \quad (2.29)$$

which means that the link function is then the logarithm of the rate (Fahrmeir et al. 2013), i.e.

$$\ln(\lambda_i) = \eta_i . \quad (2.30)$$

2.3.2 Linear mixed models (LMMs)

Data often comes in clustered form, for example due to repeated measurements of the covariate over time. Clustered data violate with the assumption of independent responses in linear regression and must be properly accounted for. One solution to this is to introduce random effects that are cluster specific, but independent of the fixed effects and the other clusters. Let the population contain m underlying clusters, with n_j , $j = 1, \dots, m$ observations in each cluster, so that $\mathbf{y} \in \mathbb{R}^{(N \times 1)}$ where $N = \sum_{j=1}^m n_j$. Assume that we investigate q random effects, including a random intercept and $q - 1$ random slopes, such that the random effects vector can be written as

$$\boldsymbol{\alpha} = (\boldsymbol{\alpha}_1, \dots, \boldsymbol{\alpha}_m)^T , \quad (2.31)$$

where each $\boldsymbol{\alpha}_j \in \mathbb{R}^{q \times 1}$ is assumed independent and represents the random effects for cluster j and has length q . For a cluster j the vector $\boldsymbol{\alpha}_j \sim \mathcal{N}_q(\mathbf{0}, \boldsymbol{\Sigma}) = \mathcal{N}_q(\mathbf{0}, \mathbf{Q}^{-1})$ where $\boldsymbol{\Sigma}$ is the $q \times q$ unknown covariance for the random effects assumed to be positive definite and $\mathbf{Q} = \boldsymbol{\Sigma}^{-1}$ the corresponding precision matrix. If the random effects for each cluster are independent of each other, the covariance matrix $\boldsymbol{\Sigma} = \text{diag}(\sigma_0^2, \dots, \sigma_q^2)$. The linear mixed model now takes the form

$$\mathbf{y} = \mathbf{X}\boldsymbol{\beta} + \mathbf{U}\boldsymbol{\alpha} + \boldsymbol{\varepsilon} , \quad (2.32)$$

where $\mathbf{X} \in \mathbb{R}^{N \times p}$ is the design matrix for the fixed effects, $\boldsymbol{\beta} \in \mathbb{R}^{p \times 1}$ are the regression coefficients for the fixed effects, $\mathbf{U} = \text{diag}(\mathbf{U}_j), \in \mathbb{R}^{N \times q}$ is the design matrix for the random effects and $\mathbf{U}_j \in \mathbb{R}^{n_j \times q}$ is the design matrix for cluster j . Since $\boldsymbol{\alpha}$ is a random variable, the parameter to estimate is the variance of each random effect $\boldsymbol{\Sigma}_{kk} = \sigma_k^2$ and their covariance $\boldsymbol{\Sigma}_{k,l} = \sigma_{k,l}$, where $k, l = 1, \dots, q$. In practice, it is often easier to estimate the precision rather than the variance, so calculations often involve the precision matrix \mathbf{Q} rather than the covariance matrix $\boldsymbol{\Sigma}$. In this model the independence between clusters are conserved for the response as a whole, but it expresses the correlation that observations of the same cluster have through the random effects. As for the simple linear regression it is assumed that $\mathbf{X}\boldsymbol{\beta}$ is fixed, and that \mathbf{U} is given, so they do not contribute to the model's variance. Therefore, the conditional expectation $\mathbb{E}(\mathbf{y}|\mathbf{X}, \mathbf{U}) = \mathbf{X}\boldsymbol{\beta}$ is easily obtained, and the conditional variance can be calculated as

$$\text{Var}(\mathbf{y}|\mathbf{X}, \mathbf{U}) = \text{Var}(\mathbf{X}\boldsymbol{\beta} + \mathbf{U}\boldsymbol{\alpha} + \boldsymbol{\varepsilon}) = \mathbf{U}\text{Var}(\boldsymbol{\alpha})\mathbf{U}^T + \sigma^2\mathbf{I} = \mathbf{U}\mathbf{G}\mathbf{U}^T + \sigma^2\mathbf{I} , \quad (2.33)$$

where $\mathbf{I} \in \mathbb{R}^{N \times N}$ and $\mathbf{G} \in \mathbb{R}^{mq \times mq}$ is the block diagonal covariance matrix of the random effects, with $\boldsymbol{\Sigma}_j$ along the diagonal for $j = 1, \dots, m$. As we assume that

the random effects are independent of the fixed effects, and that the random error term is i.i.d. for each observation, the conditional distribution of \mathbf{y} follows that of a sum of independent normal distributions, *i.e.*

$$\mathbf{y}|\mathbf{X}, \mathbf{U} \sim \mathcal{N}_n(\mathbf{X}\boldsymbol{\beta}, \mathbf{U}\mathbf{G}\mathbf{U}^T + \sigma^2\mathbf{I}) . \quad (2.34)$$

2.3.3 Generalized linear mixed models(GLMMs)

Now that we have expanded the linear regression in two different ways, the final step to complete the regression framework is to combine the LMM and GLM to obtain the GLMM. This is done by adding random effects to the linear predictor, such that

$$\theta_{i,j} = \eta_{i,j} = \mathbf{x}_{i,j}^T \boldsymbol{\beta} + \mathbf{u}_{i,j}^T \boldsymbol{\alpha}_j , \quad (2.35)$$

where $j = 1, \dots, m$ denotes the cluster and $i = 1, \dots, n_j$ denotes the observations in cluster j , $\mathbf{x}_{i,j}$ and $\mathbf{u}_{i,j}$ are the i -th columns of the submatrices \mathbf{X}_j and \mathbf{U}_j of the larger design matrices \mathbf{X} and \mathbf{U} respectively, for cluster j . The assumption of conditional independent observations $y_{i,j}$ is now conditional on the random effect as well as the covariates, and the conditional distribution of $y_{i,j}$ is still assumed to belong to the exponential family. The conditioning on the random effects is also present when choosing the appropriate link function, since one must now, in general, relate $\mathbb{E}[y_{i,j}|\mathbf{x}_{i,j}, \mathbf{u}_{i,j}, \boldsymbol{\alpha}_j]$ to the linear predictor $\eta_{i,j}$ (Fahrmeir et al. 2013). For the binary regression, this now means that the link function takes the form

$$\ln \left(\frac{\pi_{i,j}}{1 - \pi_{i,j}} \right) = \ln \left(\frac{\mathbb{P}(y_{i,j} = 1|\mathbf{x}_{i,j}, \mathbf{u}_{i,j}, \boldsymbol{\alpha}_j)}{\mathbb{P}(y_{i,j} = 0|\mathbf{x}_{i,j}, \mathbf{u}_{i,j}, \boldsymbol{\alpha}_j)} \right) = \eta_{i,j} . \quad (2.36)$$

For the Poisson random intercept model with log-link however, it is possible to define the model without conditioning on the random effects (Fahrmeir et al. 2013). This is done by noting that

$$\lambda_j = \exp(\mathbf{x}_j \boldsymbol{\beta} + \alpha_{0,j}) , \quad (2.37)$$

where $\alpha_j \sim \mathcal{N}(0, \tau_0^2)$, has a log-normal distribution. This is a special case, in which the marginal model can be determined analytically. In general however, the marginal model is not analytically tractable and so obtaining statistical inference on the GLMMs become increasingly complex when compared to the LMM. Parameter estimation therefore calls for numerical methods such as iterated reweighted least squares in the likelihood framework, or MCMC methods in the Bayesian framework, to obtain inference.

2.4 Extending R^2 to GLMMs

As we generalized the linear regression to LMMs, GLMs and GLMMs, we have to find a generalization of the concept of R^2 in order to generalize the concept of variable importance. This is fundamental to be able to propose a method for decomposing the R^2 and thereby assigning relative importance to covariates. However, the task of determining the R^2 , and decomposing it, is not a trivial task in the linear regression case and becomes even more complex in the case of GLMMs. Many extensions have been proposed, but due to a variety of theoretical

problems and/or computational difficulties, no consensus has been reached on a framework for calculating the R^2 for GLMMs (Nakagawa & Schielzeth 2013). To get an overview of the status quo for R^2 , we will follow the paper by Nakagawa & Schielzeth (2013) and go through the different components added to the linear regression to compose the GLMMs.

2.4.1 R^2 for GLMs

Recalling the definition of the R^2 from Equation (2.8), we now generalize this to the GLMs. This topic has been subject to significant research, (see for example Maddala (1983), Cameron & Windmeijer (1997), Menard (2000), Nakagawa & Schielzeth (2013)). The methods first suggested was based on the likelihood function of the model to be analysed. We will not implement such methods, as they are not suitable for the full generalization to be made later on, however they are important in building a framework for the R^2 value and are therefore included. To illustrate the likelihood based generalization of the R^2 value to GLMs, consider the deviance $\mathcal{D}(\mathbf{y}|\theta)$ function which is defined as twice the difference between the log likelihood of the **saturated model** and the log-likelihood of the model of interest (McCullagh & Nelder 1989). The saturated model denotes the model of the maximum achievable log likelihood, and therefore fits the data perfectly. For a linear regression, with $\theta = (\boldsymbol{\beta}, \sigma^2)$, we would therefore obtain

$$\begin{aligned}\mathcal{D}(\mathbf{y}|\hat{\theta}) &= -2 \left(\ln(\mathcal{L}(\boldsymbol{\beta}, \sigma^2 | \mathbf{y})) - \ln(\hat{\mathcal{L}}(\hat{\boldsymbol{\beta}}, \hat{\sigma}^2 | \mathbf{y})) \right) = -2 \left(l(\boldsymbol{\beta}, \sigma^2 | \mathbf{y}) - l(\hat{\boldsymbol{\beta}}, \hat{\sigma}^2 | \mathbf{y}) \right) \\ &= -2 \left(-\frac{n}{2} \ln(2\pi\sigma^2) - \frac{1}{2\sigma^2} (\mathbf{y} - \mathbf{X}\boldsymbol{\beta})^T (\mathbf{y} - \mathbf{X}\boldsymbol{\beta}) + \frac{n}{2} \ln(2\pi\sigma^2) \right) \\ &= \frac{1}{\sigma^2} (\mathbf{y} - \mathbf{X}\boldsymbol{\beta})^T (\mathbf{y} - \mathbf{X}\boldsymbol{\beta}) \\ &= 1 - R^2 ,\end{aligned}\tag{2.38}$$

where $\hat{\mathcal{L}}$ denotes the saturated model. Optimally, it is desirable to have as small deviance as possible while at the same time having a model that is not too complex. The best practice of the deviance is not as model fit, but rather model comparison, where one compares models through the reduction in deviance (McCullagh & Nelder 1989). Since the model of interest is nested within the saturated model, the deviance coincides with the likelihood ratio test. By comparing the model of interest to the **null model**, the simplest fit possible, one obtains for the linear regression

$$\begin{aligned}\mathcal{D}(\mathbf{y}|\hat{\theta}) - \mathcal{D}(\mathbf{y}|\theta_0) &= -2 \left(l(\boldsymbol{\beta}, \sigma^2 | \mathbf{y}) - l(\hat{\boldsymbol{\beta}}, \hat{\sigma}^2 | \mathbf{y}) \right) + 2 \left(l(\boldsymbol{\beta}_0, \sigma_0^2 | \mathbf{y}) - l(\hat{\boldsymbol{\beta}}, \hat{\sigma}^2) \right) \\ &= -2 (l(\boldsymbol{\beta}, \sigma^2 | \mathbf{y}) - l(\boldsymbol{\beta}_0, \sigma_0^2 | \mathbf{y})) \\ &= -\frac{2}{2\sigma^2} ((\mathbf{y} - \mathbf{X}\boldsymbol{\beta})^T (\mathbf{y} - \mathbf{X}\boldsymbol{\beta}) - (\mathbf{y} - \bar{\mathbf{y}})^T (\mathbf{y} - \bar{\mathbf{y}})) \\ &= 1 - \frac{(\mathbf{y} - \mathbf{X}\boldsymbol{\beta})^T (\mathbf{y} - \mathbf{X}\boldsymbol{\beta})}{\sigma^2} \\ &= R^2 .\end{aligned}\tag{2.39}$$

This is the foundation for the definitions of the generalization of R^2 to GLMs (Nakagawa & Schielzeth 2013), which primarily rely on a ratio of the maximum

likelihood of the model of interest and null model. However, in Nakagawa & Schielzeth (2013), two different R^2 measures are proposed as

$$R_G^2 = \left[1 - \left(\frac{\mathcal{L}_0}{\mathcal{L}_M} \right)^{2/n} \right] \frac{1}{1 - (\mathcal{L}_0)^{2/n}} \quad (2.40)$$

and

$$R_D^2 = 1 - \frac{-2 \ln(\mathcal{L}_M)}{-2 \ln(\mathcal{L}_0)} \quad (2.41)$$

where n denotes the total sample size, \mathcal{L}_0 is the likelihood of the null model and \mathcal{L}_M is the likelihood of the model of interest. The reason why we will not apply likelihood based R^2 measures is that when generalizing to the larger class of GLMMs, it is often desirable to do parameter estimation using the restricted maximum likelihood (REML) instead of the maximum likelihood (ML) (Fahrmeir et al. 2013). The REML estimator transforms the data, meaning that models cannot be compared when fitted, and therefore the proposed measure of R^2 is not applicable to the REML framework (Nakagawa & Schielzeth 2013). However, the extension of the R^2 measure to the larger class GLMMs will also cover the special case of GLMs, and is discussed further below in Section 2.4.3.

2.4.2 R^2 for LMMs and random slope models

In the LMMs, as opposed to the linear regression, one wishes to estimate two or more variance components instead of only the residual error variance. This increases complexity and makes the task of assigning relative importance to the covariates even more challenging. Initially, a definition was proposed for the R^2 in the LMMs that included fixed effects separately and then estimated the reduction in each variance component (Nakagawa & Schielzeth 2013, refering to Raudenbush & Bryk 1986, 1992). This violated a key condition, as adding a covariate could decrease σ_ε^2 while at the same time increasing σ_α^2 , which can lead to a negative R^2 . To handle this problem, Snijders & Bosker (1994) (Nakagawa & Schielzeth 2013) proposed a new definition of the R^2 , dividing it into two components R_1^2 and R_2^2 . Considering the simple random intercept model in scalar form;

$$y_{i,j} = \beta_0 + \mathbf{x}_{i,j}^T \boldsymbol{\beta} + \alpha_j + \varepsilon_{i,j}, \quad (2.42)$$

where $y_{i,j}$ denotes the i th observation in cluster j , β_0 is the fixed intercept, $\mathbf{x}_{i,j}$ is the column vector containing the covariates for the i th observation in cluster j , $\boldsymbol{\beta}$ is the $p \times 1$ vector of fixed effects, α_j is the random intercept for cluster j and $\varepsilon_{i,j}$ is the residual error for the i th observation in cluster j , the two R^2 components can be expressed in two ways, with the first being

$$\begin{aligned} R_1^2 &= 1 - \frac{\text{Var}(y_{i,j} - \hat{y}_{i,j})}{\text{Var}(y_{i,j})} = 1 - \frac{\sigma_\varepsilon^2 + \sigma_\alpha^2}{\sigma_{\varepsilon 0}^2 + \sigma_{\alpha 0}^2} \\ \hat{y}_{i,j} &= \beta_0 + \mathbf{x}_{i,j}^T \boldsymbol{\beta}, \end{aligned} \quad (2.43)$$

where $\sigma_{\varepsilon 0}^2$ and $\sigma_{\alpha 0}^2$ denote the residual and random effect variances of the null model respectively (Nakagawa & Schielzeth 2013) and $\hat{y}_{i,j}$ denotes the fitted value

of observation i in the j th cluster. Similarly, the second component is defined as

$$\begin{aligned} R_2^2 &= 1 - \frac{\text{Var}(y_j - \hat{y}_j)}{\text{Var}(\bar{y}_j)} = 1 - \frac{\sigma_\varepsilon^2 + \sigma_\alpha^2/k}{\sigma_{\varepsilon 0}^2 + \sigma_{\alpha 0}^2/k} \\ k &= \frac{M}{\sum_{j=1}^M \frac{1}{m_j}}, \end{aligned} \quad (2.44)$$

where \bar{y}_j is the mean for each observed value of the j th cluster, \hat{y}_j is the mean of the fitted values for the j th cluster, k is the harmonic mean of the number of observations per cluster, m_j is the number of observations for the j th cluster and M is the total number of clusters (Nakagawa & Schielzeth 2013). Note that we have formulated the above definitions in a notation corresponding to our previous formulation of the LMM, and therefore uses clusters in general, whereas Nakagawa & Schielzeth (2013) refers to a cluster as being individuals with repeated measurements. The reason for dividing the R^2 into two components, is that intuitively the R_1^2 measures the within cluster variance explained and the R_2^2 measures the between cluster variance explained (Nakagawa & Schielzeth 2013). However, three problems arise when using this definition of the R^2 for LMMs. Firstly, the R_1^2 and R_2^2 can decrease in large models, secondly, R_1^2 and R_2^2 have not been generalized to more complex LMMs with more than one random effect and lastly, it is not clear how to generalize the R_1^2 and R_2^2 to GLMMs (Nakagawa & Schielzeth 2013). To overcome these obstacles, Nakagawa & Schielzeth (2013) proposes a new formulation of the R^2 measure. Consider a general random intercept model as defined in Section 2.3.2, with q random intercepts, as

$$\mathbf{y} = \mathbf{X}\boldsymbol{\beta} + \mathbf{U}\boldsymbol{\alpha} + \boldsymbol{\varepsilon}, \quad (2.45)$$

with the parameters of interest being $\boldsymbol{\beta}$ and the variance components σ_ε^2 and σ_i^2 for the $i = 1, \dots, q$ clusters. Then define the variance of the fixed effects as

$$\sigma_f^2 = \text{Var}(\mathbf{X}\boldsymbol{\beta}) = \boldsymbol{\beta}^T \text{Var}(\mathbf{X}) \boldsymbol{\beta}, \quad (2.46)$$

and further define the R^2 for the LMM as

$$R_{\text{LMM(m)}}^2 = \frac{\sigma_f^2}{\sigma_f^2 + \sum_{i=1}^q \sigma_i^2 + \sigma_\varepsilon^2}. \quad (2.47)$$

This definition of the R_{LMM}^2 represents the marginal R_{LMM}^2 , denoted by (m) , as it measures the proportion of the variance explained by the fixed effects alone, whereas the conditional R_{LMM}^2 can be defined as

$$R_{\text{LMM(c)}}^2 = \frac{\sigma_f^2 + \sum_{i=1}^q \sigma_i^2}{\sigma_f^2 + \sum_{i=1}^q \sigma_i^2 + \sigma_\varepsilon^2}. \quad (2.48)$$

By inspection, it is clear that this definition will never lead to negative values of the R_{LMM}^2 . It may occur that the R_{LMM}^2 value may decrease when adding more covariates to the model, although Nakagawa & Schielzeth (2013) argues that this is unlikely. This definition now covers the random intercept model, but has not taken into account the possibility of having a LMM with a random slope. To further extend the R^2 to the random slope model, Johnson (2014) proposes a

method for computing the mean random effect variance. Consider the simple random intercept and slope model,

$$y_{i,j} = \beta_0 + \mathbf{x}_{i,j}^T \boldsymbol{\beta} + \alpha_{0,j} + \alpha_{1,j}x_{i,j} + \varepsilon_{i,j}, \quad (2.49)$$

where the same notation is used as in (2.42) with $\boldsymbol{\alpha}_j = (\alpha_{0,j}, \alpha_{1,j})$ being the random effect, $\alpha_{0,j}$ denoting the random intercept and $\alpha_{1,j}$ now denoting the random deviation from the global slope β_1 , for cluster j . The general assumption on the random effects are that

$$\begin{pmatrix} \alpha_0 \\ \alpha_1 \end{pmatrix} \sim \mathcal{N} \left(\begin{pmatrix} 0 \\ 0 \end{pmatrix}, \Sigma = \begin{pmatrix} \sigma_{\alpha_0}^2 & \sigma_{\alpha_0, \alpha_1} \\ \sigma_{\alpha_0, \alpha_1} & \sigma_{\alpha_1}^2 \end{pmatrix} \right), \quad (2.50)$$

where $\sigma_{\alpha_0}^2$ and $\sigma_{\alpha_1}^2$ are the variances of the random intercept and random slope respectively, and $\sigma_{\alpha_0, \alpha_1}$ is the covariance between the random intercept and random slope. Thus, we have three variance components of interest ($\frac{q(q+1)}{2}$ for q random effects) to estimate. When inspecting the variance of the random part in the model, we see that it has a dependence on the covariates, as illustrated by

$$\begin{aligned} \text{Var}(\alpha_{0,j} + \alpha_{1,j}x_{i,j}) &= \text{Var}(\alpha_{0,j}) + 2x_{i,j}\text{Cov}(\alpha_{0,j}, \alpha_{1,j}) + x_{i,j}^2\text{Var}(\alpha_{1,j}) \\ &= \sigma_{\alpha_0}^2 + 2x_{i,j}\sigma_{\alpha_0, \alpha_1} + x_{i,j}^2\sigma_{\alpha_1}^2 =: \sigma_{r,i,j}^2, \end{aligned} \quad (2.51)$$

where we define $\sigma_{r,i,j}^2$ as the variance of the random effect $\boldsymbol{\alpha}$ for observation i in the j th cluster. The method proposed by Johnson (2014) is to first estimate all the variance components, and then view the specific random effect as a normal mixture distribution of the random intercept and random slope. This mixture distribution is characterized as having a common mean of zero, and, if all values of the associated covariate $x_{i,j}$ are unique, having N different variances with N being the total number of observations. A mixture distribution with constant mean, has a variance which equals the mean of the individual variances in the distribution (Johnson 2014, citing Behboodian 1970). The proposed variance of the random effect $\boldsymbol{\alpha}$, is therefore the mean of the variance components in $\boldsymbol{\alpha}$, *i.e.*

$$\bar{\sigma}_r^2 = \frac{1}{N} \sum_{j=1}^q \sum_{i=1}^{n_j} (\sigma_{r,i,j}^2). \quad (2.52)$$

This formulation can be generalized in the case of q random effects, where each random effect has an associated design matrix \mathbf{U}_j and covariance matrix \mathbf{Q} as in Section 2.3.2, so that for each random effect r we have

$$\bar{\sigma}_r^2 = \text{Tr}(\mathbf{U}_j \mathbf{Q} \mathbf{U}_j^T), \quad r = 1, \dots, q. \quad (2.53)$$

To finally obtain the proposed R^2 for the general LMM, Johnson (2014) uses this estimate in the definition given by Nakagawa & Schielzeth (2013), to obtain

$$R_{\text{LMM(m)}}^2 = \frac{\sigma_f^2}{\sigma_f^2 + \sum_{r=1}^q \bar{\sigma}_r^2 + \sigma_\varepsilon^2}, \quad (2.54)$$

and

$$R_{\text{LMM(c)}}^2 = \frac{\sigma_f^2 + \sum_{r=1}^q \bar{\sigma}_r^2}{\sigma_f^2 + \sum_{i=1}^q \bar{\sigma}_r^2 + \sigma_\varepsilon^2}, \quad (2.55)$$

as the marginal and conditional R^2_{LMM} respectively. For the random intercept model with $\sigma_{r,i,j}^2 = \sigma_r^2$, this definition corresponds to the definition by Nakagawa & Schielzeth (2013) as

$$\overline{\sigma_r^2} = \frac{1}{N} \sum_{j=1} \sum_{i=1} (\sigma_{r,i,j}^2) = \sigma_{r,i,j}^2 = \sigma_r^2 . \quad (2.56)$$

The R^2_{LMM} proposed by Johnson now lets us compute the R^2 for general LMMs, however it is argued in Johnson (2014) whether the improved R^2 estimate by taking the random slope into account is worth the added complexity and computational cost.

2.4.3 R^2 for GLMMs

The final step towards a complete generalization for the R^2 value of regression models is to extend it to the GLMMs. When considering non-normal responses, the link function introduces an aspect not yet discussed, which is to define the residual variance. One can divide the residual variance σ_ε^2 into three components, namely distribution specific variance, multiplicative dispersion and additive dispersion (Nakagawa & Schielzeth 2013). The distribution specific variance is inherited from the link function used, and is therefore known before analysis is done. However, the multiplicative and additive dispersion is modelled to account for the variance present that exceeds the distribution specific variance, *i.e.* overdispersion (Nakagawa & Schielzeth 2010). Therefore, one must specify upon implementation on what scale the overdispersion is to be modelled. The multiplicative dispersion, denoted by ω , is overdispersion on the response (data) scale and modelled as a distinct parameter of the assumed distribution of the response \mathbf{y} (Nakagawa & Schielzeth 2010). Conversely, the additive dispersion, denoted by e , is overdispersion on the latent scale and introduced to the model as an additional random effect in the linear predictor (Nakagawa & Schielzeth 2010). Defining the residual variance now depends on the choice of dispersion modelling, and is either defined as

$$\sigma_\varepsilon^2 = \omega \sigma_d^2 \quad (2.57)$$

or

$$\sigma_\varepsilon^2 = \sigma_d^2 + \sigma_e^2 , \quad (2.58)$$

for multiplicative and additive dispersion respectively. With the residual variance defined, the generalization to of the R^2 to GLMMs (thereby also the GLMs) follows the same logic as the LMMs, and $R^2_{\text{GLMM (m)}}$ is defined as

$$R^2_{\text{GLMM(m, m)}} = \frac{\sigma_f^2}{\sigma_f^2 + \sum_{r=1}^q \overline{\sigma_r^2} + \sigma_\varepsilon^2} = \frac{\sigma_f^2}{\sigma_f^2 + \sum_{r=1}^q \overline{\sigma_r^2} + \omega \sigma_d^2} , \quad (2.59)$$

and

$$R^2_{\text{GLMM(m, a)}} = \frac{\sigma_f^2}{\sigma_f^2 + \sum_{r=1}^q \overline{\sigma_r^2} + \sigma_\varepsilon^2} = \frac{\sigma_f^2}{\sigma_f^2 + \sum_{r=1}^q \overline{\sigma_r^2} + \sigma_d^2 + \sigma_e^2} , \quad (2.60)$$

where the same notation as before is used and the subscripts (m, m) and (m, a) denote the multiplicative and additive dispersion respectively. The conditional R^2_{GLMM} can be defined similarly as,

$$R^2_{\text{GLMM(c, m)}} = \frac{\sigma_f^2 + \sum_{r=1}^q \bar{\sigma}_r^2}{\sigma_f^2 + \sum_{r=1}^q \bar{\sigma}_r^2 + \omega \sigma_d^2}, \quad (2.61)$$

and

$$R^2_{\text{GLMM(c, a)}} = \frac{\sigma_f^2 + \sum_{r=1}^q \bar{\sigma}_r^2}{\sigma_f^2 + \sum_{r=1}^q \bar{\sigma}_r^2 + \sigma_d^2 + \sigma_e^2}, \quad (2.62)$$

completing the generalization.

2.4.4 Extensions of the LMG and relative weights method

Extensions for both the LMG and the relative weights method have been proposed so that they can decompose the R^2 also for random intercept models (Matre 2022). The extended LMG, denoted as the ELMG, method uses the same permutations as described for the regular LMG, and now decomposes the R^2 value for the random intercept model instead. From this decomposition, the only extension needed for the LMG formula is to include also the random intercepts as model components, which for covariate i gives

$$\text{LMG}(i) = \frac{1}{(p+q)!} \sum_{\substack{S \subseteq \{2, \dots, p \\ p+1, \dots, p+q\} \setminus i}} n(S)!((p+q) - n(S) - 1)! \text{svar}(\{i\}|S), \quad (2.63)$$

where p denotes fixed effects and q denotes random effects (Matre 2022). It is equivalent to the original LMG method (2.15) except that here the random intercepts are treated as categorical fixed effects, where we do not consider the columns but rather the whole predictor, either completely in the model or not.

To create the extended relative weights (ERW) method, Matre (2022) uses the same transformation of data as for the relative weights method to project the covariates into an orthogonal space. Then the fixed effects are treated as one separate block, either in the model or not, and then uses the LMG approach to distribute a share of R^2 to each random intercept. The fixed effects will receive a joint share, which is distributed by using the relative weights method. Since the LMG approach is used for the q random effects and the block of fixed effects, the complexity of the ERW will follow that of the LMG method for $q+1$ covariates.

Both extensions described comes with new considerations (Matre 2022, for full details), for example that the inclusion criteria now should read "If a regressor $\beta_j \neq 0$, or a random intercept α with $\sigma^2(\alpha) > 0$, is included in the model then its share of the variance should be positive".

2.5 The Bayesian framework

So far, we have introduced statistical concepts without considering the framework in which they are used. We now expand the theory to consider the Bayesian framework, which is the framework used in this thesis.

2.5.1 General idea

The Bayesian framework stems from the notorious theorem developed by Thomas Bayes, (Bayes & Price 1763), which states that for events A and B , with nonzero probability of occurring, we have

$$\mathbb{P}(A|B) = \frac{\mathbb{P}(B \cap A)}{\mathbb{P}(B)} = \frac{\mathbb{P}(B|A)\mathbb{P}(A)}{\mathbb{P}(B)} . \quad (2.64)$$

This can be generalized to also apply to distributions of continuous random variables, namely that

$$\pi(\boldsymbol{\theta}|\mathbf{y}) = \frac{\pi(\mathbf{y}|\boldsymbol{\theta})\pi(\boldsymbol{\theta})}{\pi(\mathbf{y})} , \quad (2.65)$$

where $\pi(\boldsymbol{\theta}|\mathbf{y})$ is called the posterior distribution of $\boldsymbol{\theta}$, $\pi(\mathbf{y}|\boldsymbol{\theta})$ is the likelihood, or sampling, distribution of \mathbf{y} , $\pi(\boldsymbol{\theta})$ is the prior distribution of the parameters and $\pi(\mathbf{y}) = \int \pi(\mathbf{y}|\boldsymbol{\theta})\pi(\boldsymbol{\theta}) < \infty$ is the marginal distribution of the data, which is required to be finite in order to have a proper posterior distribution (Gelman et al. 2015). In practice, the marginal distribution is often omitted and one only consider the proportionality of (2.65), i.e.

$$\pi(\boldsymbol{\theta}|\mathbf{y}) \propto \pi(\mathbf{y}|\boldsymbol{\theta})\pi(\boldsymbol{\theta}) . \quad (2.66)$$

In the context of statistical analysis, with $\boldsymbol{\theta}$ being the parameter vector of the family of models for the random variable Y under investigation, $\pi(\boldsymbol{\theta}|\mathbf{y})$ is interpreted as the distribution of the parameters given the data \mathbf{y} . This is the key element that separates the Bayesian framework from the frequentist framework, as the parameter $\boldsymbol{\theta}$ is now treated as random variable instead of being point estimates.

2.5.2 Prior and posterior distributions

Generally, a Bayesian model is built by first introducing some prior knowledge through the prior distribution $\pi(\boldsymbol{\theta})$ and supplementing this with the likelihood function $\pi(\mathbf{y}|\boldsymbol{\theta})$. The prior distribution must be chosen based on the prior knowledge available, and can either be informative, noninformative or weakly informative (Gelman et al. 2015). As a compromise of the information in the prior and the likelihood of the data, the posterior distribution is obtained. The resulting posterior will be different from analysis to analysis, but some general relations between the prior and posterior are discussed in Gelman et al. (2015). In particular, it is stated that "*the posterior variance is on average smaller than prior variance by an amount that depends on the variation in posterior means over the distribution of possible data*" (Gelman et al. 2015). This further means that if one wishes to reduce the variability in the posterior, the potential for this lies in reducing the variation of possible posterior means. The posterior distribution will therefore, in general, be a compromise between the prior and the likelihood, which with increasing sampling size will be increasingly influenced by the likelihood (Gelman et al. 2015).

2.5.3 Penalising complexity (PC) priors

Prior distributions pose a great feature by allowing for inclusion of prior information, but also a great challenge in that they must be chosen with care. As the

theory of this is vast and out of the scope for this thesis, we will be mostly concerned with the penalising complexity priors proposed in Simpson et al. (2017). In this paper, four main principles are desirable to follow when choosing a prior distribution, namely

1. **Occam's razor** - If there is no evidence for a complex mode, a base model should be preferred.
2. **Measure of complexity** - The measure of model complexity is defined as $d(f||g) = \sqrt{2\text{KLD}(f||g)}$ where $\text{KLD}(f||g)$ denotes the Kullback-Leibler divergence (Simpson et al. 2017, for more information).
3. **Constant rate penalisation** - The penalisation, i.e. the decay of prior mass, grows as the complexity grows, but it is desirable that this growth is constant.
4. **User defined scaling** - Assuming that the user has an idea of the magnitude of the parameter of interest, the user should be able to scale the prior accordingly.

The PC priors therefore pose interpretable, applicable priors which are consistent with the above principles, and are therefore a practical choice for the Bayesian framework (Simpson et al. 2017). Particularly, for the case of a linear mixed model with a Gaussian random effect $\alpha \sim \mathcal{N}(0, \sigma^2 \mathbf{R}) = \mathcal{N}(0, \tau^{-1} \mathbf{Q}^{-1})$, the base model of the PC priors corresponds to the case where the precision $\tau = 0$ and the prior for τ takes the form

$$\pi(\tau) = \frac{\lambda}{2} \tau^{-3/2} \exp(-\lambda \tau^{-1/2}), \quad \tau, \lambda > 0. \quad (2.67)$$

To specify λ , the user is required to supply the values (U, a) such that $\mathbb{P}(1/\sqrt{\tau} > U) = a$. This defines the scaling parameter of principle 4 and leads to $\lambda = -\ln(a)/U$ (Simpson et al. 2017). When fitting additive models, thereby modelling additive overdispersion, using PC priors is a natural choice (Gómez-Rubio 2020).

2.5.4 Hierarchical Bayesian modelling

When modelling in the Bayesian framework, the posterior distribution of the parameter $\boldsymbol{\theta}$ given the data is what one wants to infer. For many applications, $\boldsymbol{\theta}$ is a high dimensional vector, with naturally connected entries (Gelman et al. 2015). It may therefore be reasonable to assume that the parameters themselves are drawn from a population distribution, which can further be modelled by what is called hyperparameters. The main idea is that the prior $\pi(\boldsymbol{\theta})$ itself contains a hierarchical structure and can be split into levels of conditional prior distributions, i.e. $\pi(\boldsymbol{\theta}) = \pi(\boldsymbol{\theta}|\boldsymbol{\phi})\pi(\boldsymbol{\phi})$ for some hyperparameter $\boldsymbol{\phi}$ (Robert 2007). Assuming that the data \mathbf{y} depends only on the parameter $\boldsymbol{\theta}$, and that $\boldsymbol{\theta}$ depends on the hyperparameters $\boldsymbol{\phi}$, we can write the joint posterior distribution of $(\boldsymbol{\theta}, \boldsymbol{\phi})$ as

$$\pi(\boldsymbol{\theta}, \boldsymbol{\phi}|\mathbf{y}) \propto \pi(\mathbf{y}|\boldsymbol{\theta}, \boldsymbol{\phi})\pi(\boldsymbol{\theta}|\boldsymbol{\phi})\pi(\boldsymbol{\phi}) = \pi(\mathbf{y}|\boldsymbol{\theta})\pi(\boldsymbol{\theta}|\boldsymbol{\phi})\pi(\boldsymbol{\phi}), \quad (2.68)$$

where $\pi(\boldsymbol{\phi})$ is a prior placed on the hyperparameters. This hierarchical structure allows us to first estimate the population distribution using the hyperparameters,

and then estimate the parameters of interest using the population distribution, instead of estimating each component of $\boldsymbol{\theta}$ separately (Gelman et al. 2015). It may be practical to view the model in three parts and consider an example with a tractable posterior distribution. Let the observational model be $\pi(\mathbf{y}|\boldsymbol{\theta})$ be defined as

$$y_i|\theta_i \sim \text{Po}(\theta_i), i = 1, \dots, n, \quad (2.69)$$

for conditionally independent observations y_i given the parameters θ_i . Define then the latent model $\pi(\boldsymbol{\theta}|\boldsymbol{\phi})$ as

$$\theta_i|\boldsymbol{\phi} \sim \text{Gamma}(\alpha, \beta), \quad (2.70)$$

for conditionally independent parameters θ_i given the hyperparameters α, β . Lastly, consider the hyperpriors $\pi(\boldsymbol{\phi})$ as

$$\alpha \sim \text{Exp}(a) \text{ and } \beta \sim \text{Gamma}(b, c), \quad (2.71)$$

The full posterior density now reads

$$\pi(\boldsymbol{\theta}, \alpha, \beta | \mathbf{y}) \propto \underbrace{\prod_{i=1}^n \theta_i^{y_i} e^{-\theta_i}}_{\text{Po}(\theta_i)} \underbrace{\prod_{i=1}^n \frac{\beta^\alpha}{\Gamma(\beta)} \theta_i^{\alpha-1} e^{-\beta\theta_i}}_{\text{Gamma}(\alpha, \beta)} \underbrace{\alpha^{a-1} e^{-\alpha}}_{\text{Exp}(a)} \underbrace{\beta^{b-1} e^{-c\beta}}_{\beta \sim \text{Gamma}(b, c)}, \quad (2.72)$$

which can be used to make inference about the parameters of interest. This hierarchical structure is similar to that of the GLMM and is therefore a natural way of modelling a Bayesian GLMM. To set up a Bayesian GLMM, consider again observations \mathbf{y} having the density function in (2.21) with dispersion parameter ϕ and associated linear predictor

$$\boldsymbol{\eta} = \mathbf{X}\boldsymbol{\beta} + \mathbf{U}\boldsymbol{\alpha}, \quad (2.73)$$

where we assume that $\boldsymbol{\alpha} \sim \mathcal{N}(0, \mathbf{Q}^{-1})$ for some precision matrix $\mathbf{Q} = \mathbf{Q}(\boldsymbol{\rho})$ dependent on the hyperparameter $\boldsymbol{\rho}$. Then, to define the model, a prior must be assigned to the likelihood specific parameter ϕ , the fixed effects coefficients $\boldsymbol{\beta}$, and the variance components of the random effects $\boldsymbol{\rho}$. For a general GLMM belonging to the exponential family defined in (2.21), the posterior can be written out as

$$\begin{aligned} \pi(\boldsymbol{\beta}, \boldsymbol{\alpha}, \phi, \boldsymbol{\rho} | \mathbf{y}) &\propto \left(\prod_{j=1}^m \pi(\mathbf{y}_j | \boldsymbol{\beta}, \boldsymbol{\alpha}, \phi, \boldsymbol{\rho}) \right) \pi(\boldsymbol{\alpha} | \boldsymbol{\rho}) \pi(\boldsymbol{\beta}) \pi(\phi) \pi(\boldsymbol{\rho}), \\ &\propto \exp \left(-\frac{1}{2} \boldsymbol{\alpha}^T \mathbf{Q}(\boldsymbol{\rho}) \boldsymbol{\alpha} + \sum_{j=1}^m \ln \pi(\mathbf{y}_j | \boldsymbol{\beta}, \boldsymbol{\alpha}, \phi) \right) |\mathbf{Q}(\boldsymbol{\rho})|^{1/2} \pi(\boldsymbol{\beta}) \pi(\phi) \pi(\boldsymbol{\rho}), \end{aligned} \quad (2.74)$$

where the vector \mathbf{y}_j denotes the j th cluster of observations (Fong et al. 2010).

2.5.5 R^2 in the Bayesian framework

When working in the Bayesian framework, the definition of R^2 for the linear regression is not as straightforward as in the classical framework. As parameters are not treated as fixed, but as random variables, the R^2 value will also be a random

variable. A possible remedy to this could be to use the posterior mode of the parameters β in (2.8), however Gelman et al. (2017) states two conflicts that this poses. Firstly, the use of point estimates to calculate statistics in the Bayesian framework rejects the fundamental uncertainty of the Bayesian framework. Secondly, when the parameters are estimated in a Bayesian framework, there is no guarantee that the $R^2 \in [0, 1]$, reducing its intuitive interpretability. In Gelman et al. (2017) a definition of the R^2 for the Bayesian linear regression is proposed. Consider a draw s of the parameters β from the posterior distribution. Then, the proposed definition is

$$R_s^2 = \frac{\beta_s^T \Sigma_{\mathbf{X}^T \mathbf{X}} \beta_s}{\beta_s^T \Sigma_{\mathbf{X}^T \mathbf{X}} \beta_s + \sigma_s^2}, \quad (2.75)$$

where $\Sigma_{\mathbf{X}^T \mathbf{X}}$ is the covariance matrix of the design matrix \mathbf{X} and σ_s^2 is the variance of the error term which can be sampled from the posterior distribution. Contrary to the classical definition this definition of R^2 contains only the estimated values from our model and not the observed values. The reasoning behind this is to carry this inherent uncertainty in the Bayesian framework by not using point estimates from the posterior mean, but rather averaging over a posterior distribution. Drawing enough samples from (2.75) one would eventually obtain also an approximation of the distribution for the R^2 value (Gelman et al. 2017).

2.5.6 Variable importance measures in the Bayesian framework

Although the field of Bayesian variable importance is small, there are some methods that can be contextualized so that they can be used as Bayesian variable importance measures. More specifically, the R^2 -induced Dirichlet decomposition (R2D2) priors are originally applied as shrinkage priors and used to obtain reliable predictions in high dimensional linear regression models, but can be interpreted as a variable importance measure (Zhang et al. 2020). Using the same definition of R^2 as that of Gelman et al. (2017), the R2D2 prior directly places a prior on the marginal or conditional R^2 value. We will consider the marginal R^2 prior, which assumes the marginal R^2 value to follow a Beta distribution (Zhang et al. 2020). The scenario introduced in Zhang et al. (2020) is a linear regression model where we consider a prior for β such that $\mathbb{E}[\beta] = 0$ and $\text{cov}(\beta) = \sigma^2 \Lambda$ where Λ is a diagonal matrix with the diagonal elements $\lambda_1, \dots, \lambda_p$. From the calculations in Zhang et al. (2020) and following the definition of the R^2 in Gelman et al. (2017), one can write

$$\text{Var}(\mathbf{x}^T \beta) = \sigma^2 \sum_{j=1}^p \lambda_j \quad (2.76)$$

and

$$R^2 = \frac{\text{Var}(\mathbf{x}^T \beta)}{\text{Var}(\mathbf{x}^T \beta) + \sigma^2} = \frac{\sum_{j=1}^p \lambda_j}{\sum_{j=1}^p \lambda_j + 1} := \frac{W}{W + 1} \quad (2.77)$$

where W is the sum of the diagonal elements of Λ . Assuming that $W \sim BP(a, b)$ where BP denotes the Beta prime distribution, is equivalent to assuming that $R^2 \sim Beta(a, b)$ value follows a Beta distribution (Zhang et al. 2020). Further, Zhang et al. (2020) expresses each $\lambda_j = \phi_j \omega$ with $\sum_{j=1}^p \lambda_j = 1$ such that

$W = \sum_{j=1}^p \phi_j \omega = \omega$. This is the key element that makes the R2D2 prior analogous to a variable importance measure, because ω represents the total prior variability and ϕ_j represents the proportion of the total variability allocated to covariate j for $j = 1, \dots, p$ (Zhang et al. 2020). Now, to decompose the model R^2 based on these priors, it is proposed to let $\boldsymbol{\phi} = (\phi_1, \dots, \phi_p) \sim Dir(a_\pi, \dots, a_\pi)$ where Dir denotes the Dirichlet distribution, and a_π is a concentration parameter (Zhang et al. 2020). This means that ϕ_j is estimated via a Dirichlet decomposition which is analogous to assigning each covariate with a share of relative variable importance. The concentration parameters can be seen as an a priori importance for the covariates (Aguilar & Bürkner 2024) and larger a_π leads to smaller variance of ϕ_j and produces a more uniform $\boldsymbol{\phi}$. Conversely, a smaller a_π leads $\boldsymbol{\phi}$ to have some components with a larger ϕ_j (Zhang et al. 2020). The authors then show the prior on $\boldsymbol{\beta}$ that is induced by the prior on R^2 and further develop the theory and list properties of the R2D2 priors (Zhang et al. 2020). An obvious advantage with the R2D2 priors for the marginal R^2 is that they allow for more extensive asymptotic analysis of both the prior and posterior distributions (Zhang et al. 2020). However, the Dirichlet distribution has some limitations that make it unsuitable for many applications. Firstly, when multiple covariates compete for the shares of importance, the Dirichlet distribution has a tendency to gravitate this competition towards a negative dependency structure, which is completely determined by the mean of each component (Aguilar & Bürkner 2024). Further, the Dirichlet distribution is not very flexible, and therefore struggles to model correlation structures between covariates (Aguilar & Bürkner 2024, and references therein). The Dirichlet distribution enforces a high number of constraints on the covariance structure, making some covariance structures impossible to model (Aguilar & Bürkner 2024). Therefore, Aguilar & Bürkner (2024) propose what they call Generalized Decomposition Priors on R^2 (GDR2). The GDR2 priors address some limitations in the R2D2 priors, and suggests to rather place a Logistic Normal (LN) prior on the parameters $\boldsymbol{\phi}$. This means that $\boldsymbol{\phi} \sim LN(\boldsymbol{\mu}, \boldsymbol{\Sigma})$ and instead of specifying the parameter a_π , we must now specify the mean $\boldsymbol{\mu}$ and the covariance matrix $\boldsymbol{\Sigma}$ (Aguilar & Bürkner 2024). The authors suggest automating the process of choosing prior values for the mean and covariance of $\boldsymbol{\phi}$ by what they call *prior matching*. By letting $f \sim Dir(\alpha)$ for a fixed $\alpha = a_\pi$ and $g \sim LN(\boldsymbol{\mu}, \boldsymbol{\Sigma})$, the Kullback-Leibler divergence (Kullback & Leibler 1951) between f and g can be minimized. In Aguilar & Bürkner (2024) a closed form for the minimizers is obtained as

$$\begin{aligned}\boldsymbol{\mu}_k^* &= \delta(\alpha_k) - \delta(\alpha_K) \\ \sigma_{kk}^* &= \varepsilon(\alpha_k) + \varepsilon(\alpha_K) \\ \sigma_{kj}^* &= \varepsilon(\alpha_k) \quad k \neq j ,\end{aligned}\tag{2.78}$$

for $k = 1, \dots, K$ where α_K is a reference unit, $\delta(\alpha_k)$ denotes the digamma function and $\varepsilon(\alpha_k) = \delta'(\alpha_k)$ denotes the first polygamma function (Abramowitz & Stegun 1972, pages 258-260). We will not go further into detail on R2D2 and GDR2 priors, but rather focus on the interpretation of the R2D2 and GDR2 prior as variable importance measures by assessing the values of $\boldsymbol{\phi}$.

2.6 The INLA framework

As we have seen, the analytical posterior is possible to obtain for some hierarchical structures (e.g. (2.72)). However, in the case of GLMMs, the posterior distribution is not in general analytically tractable (Fong et al. 2010). This calls for the use of numerical methods, such as Markov Chain Monte Carlo (MCMC) methods, to be able to sample from the posterior distribution. Such methods are computationally expensive, and require careful analysis to justify convergence and proper mixing of the Markov chains to make sure we sample from the steady state posterior distribution. Therefore, it is desirable, under certain conditions, to look at other methods that are more computationally efficient. In this thesis we will consider the Integrated Nested Laplace Approximation (INLA) method (Gómez-Rubio 2020).

2.6.1 Introduction to INLA

The INLA method is an alternative to the classical Marko Chain Monte Carlo methods, that has significant advantages at the cost of some structural assumptions. In order to apply INLA, consider the vector of observations $\mathbf{y} = (y_1, \dots, y_n)$, which may also contain missing values. Given an appropriate link function $g(\mu_i) = \eta_i$, we can model the observations as independent given the linear predictor

$$\eta_i = \alpha + \sum_{j=1}^{n_\beta} \beta_j z_{ji} + \sum_{k=1}^{n_f} f^{(k)}(u_{ki}) + \varepsilon_i , \quad i = 1, \dots, n , \quad (2.79)$$

where α is the intercept, β_j are the regression coefficients for the covariates z_{ji} , $f^{(k)}$ are random effects for the vector of covariates $\{\mathbf{u}_k\}_{k=1}^{n_f}$ and ε_i is the error term. This gives rise to the key assumption that the INLA method needs in order to be applicable, namely that the latent field \mathbf{x} , denoted as

$$\mathbf{x} = (\eta_1, \dots, \eta_n, \alpha, \beta_1, \dots, \beta_n) , \quad (2.80)$$

is a Gaussian Markov Random Field (GMRF). Further, it is assumed that observations are independent given this latent field and the latent field is distributed according to some hyperparameters $\boldsymbol{\theta}$. The structure of the GMRF is given by a precision matrix $\mathbf{Q}(\boldsymbol{\theta})$, which is sparse and can be represented by a graph $\mathcal{G} = (\mathcal{V}, \mathcal{E})$ (see Section 2.8 for more details). This along with the assumed conditional independence makes computations very fast and is why INLA is effective. Now, the posterior distribution of the latent field \mathbf{x} is given by

$$\pi(\mathbf{x}, \boldsymbol{\theta} | \mathbf{y}) = \frac{\pi(\mathbf{y} | \mathbf{x}, \boldsymbol{\theta}) \pi(\mathbf{x} | \boldsymbol{\theta}) \pi(\boldsymbol{\theta})}{\pi(\mathbf{y})} \propto \pi(\mathbf{y} | \mathbf{x}, \boldsymbol{\theta}) \pi(\mathbf{x} | \boldsymbol{\theta}) \pi(\boldsymbol{\theta}) , \quad (2.81)$$

where $\pi(\mathbf{y} | \mathbf{x}, \boldsymbol{\theta})$ is the likelihood, $\pi(\mathbf{x} | \boldsymbol{\theta})$ is the posterior of the latent field and $\pi(\boldsymbol{\theta})$ is the prior. Since it is assumed that observations are independent given the latent field, we can further express

$$\pi(\mathbf{y} | \mathbf{x}, \boldsymbol{\theta}) = \prod_{i \in \mathcal{I}} \pi(y_i | x_i, \boldsymbol{\theta}) , \quad (2.82)$$

where the index set $\mathcal{I} \subset \{1, 2, 3, \dots, n\}$ only includes actual observed data. The INLA method now attempts to estimate the marginals of the latent effects and

the hyperparameters. These marginals are given by

$$\pi(x_l|\mathbf{y}) = \int \pi(x_l|\boldsymbol{\theta}, \mathbf{y})\pi(\boldsymbol{\theta}|\mathbf{y})d\boldsymbol{\theta} , \quad (2.83)$$

and

$$\pi(\theta_k|\mathbf{y}) = \int \pi(\boldsymbol{\theta}|\mathbf{y})d\boldsymbol{\theta}_{-k} , \quad (2.84)$$

respectively, where $\boldsymbol{\theta}_{-k}$ is the vector of hyperparameters excluding element θ_k (Gómez-Rubio 2020).

2.6.2 Approximating the marginals

As previously mentioned the marginals in Equation (2.83) and Equation (2.84) are generally not tractable, but INLA uses this form of the marginals, to construct nested approximations (Rue et al. 2009). Consider, as in (Rue et al. 2009), the approximation of the marginals in Equation (2.83) and Equation (2.84) as

$$\tilde{\pi}(x_l|\mathbf{y}) = \int \tilde{\pi}(x_l|\boldsymbol{\theta}, \mathbf{y})\tilde{\pi}(\boldsymbol{\theta}|\mathbf{y})d\boldsymbol{\theta} , \quad (2.85)$$

and

$$\tilde{\pi}(\theta_k|\mathbf{y}) = \int \tilde{\pi}(\boldsymbol{\theta}|\mathbf{y})d\boldsymbol{\theta}_{-k} , \quad (2.86)$$

where $\tilde{\pi}(\cdot, \cdot)$ is an approximation of the density $\pi(\cdot, \cdot)$. To be able to compute the above approximations, we need to first specify the approximations under the integral sign. The first one to consider is $\tilde{\pi}(\boldsymbol{\theta}|\mathbf{y})$, which Rue et al. (2009) approximates by

$$\tilde{\pi}(\boldsymbol{\theta}|\mathbf{y}) \propto \frac{\tilde{\pi}(\mathbf{x}, \boldsymbol{\theta}, \mathbf{y})}{\tilde{\pi}_G(\mathbf{x}|\boldsymbol{\theta}, \mathbf{y})} \Big|_{\mathbf{x}=\mathbf{x}^*(\boldsymbol{\theta})} , \quad (2.87)$$

where $\tilde{\pi}_G(\mathbf{x}|\boldsymbol{\theta}, \mathbf{y})$ is the Gaussian approximation of the full conditional of \mathbf{x} evaluated at the mode $\mathbf{x}^*(\boldsymbol{\theta})$ of the full conditional for given $\boldsymbol{\theta}$ (Rue et al. 2009). From Equation (2.87) the posterior marginals of hyperparameter k , $\tilde{\pi}(\theta_k|\mathbf{y})$, can be approximated by integrating out $\boldsymbol{\theta}_{-k}$ using numerical integration. However, an approximation for $\tilde{\pi}(x_l|\boldsymbol{\theta}, \mathbf{y})$ must be chosen to obtain the posterior marginals of the latent effects. To approximate $\tilde{\pi}(x_l|\boldsymbol{\theta}, \mathbf{y})$, (Rue et al. 2009) describe three strategies of varying computational complexity. The cheapest approximation (Gómez-Rubio 2020) is to derive the Gaussian marginals of $\tilde{\pi}_G(\mathbf{x}|\boldsymbol{\theta}, \mathbf{y})$ as

$$\tilde{\pi}_G(x_l|\boldsymbol{\theta}, \mathbf{y}) = \mathcal{N}(\mu_l(\boldsymbol{\theta}), \sigma_l^2(\boldsymbol{\theta})) , \quad (2.88)$$

where $\mu_l(\boldsymbol{\theta})$ is the mean vector and $\sigma_l^2(\boldsymbol{\theta})$ the corresponding vector with marginal variances of the Gaussian approximation (Rue et al. 2009). The second, more costly, approach is to use a Laplace approximation so that

$$\tilde{\pi}_{LA}(x_l|\boldsymbol{\theta}, \mathbf{y}) \propto \frac{\tilde{\pi}(\mathbf{x}, \boldsymbol{\theta}, \mathbf{y})}{\tilde{\pi}_{GG}(\mathbf{x}_{-l}|x_l, \boldsymbol{\theta}, \mathbf{y})} \Big|_{\mathbf{x}_{-l}=\mathbf{x}_{-l}^*(x_l, \boldsymbol{\theta})} \quad (2.89)$$

where $\tilde{\pi}_{GG}(\mathbf{x}_{-l}|x_l, \boldsymbol{\theta}, \mathbf{y})$ is the Gaussian approximation to the density of $\mathbf{x}_{-l}|x_l, \boldsymbol{\theta}, \mathbf{y}$ evaluated at the mode $\mathbf{x}_{-l}^*(x_l, \boldsymbol{\theta})$ (Gómez-Rubio 2020). This approximation requires computations for each value x_l , and so a simplified modification

$$\tilde{\pi}_{LA}(x_l|\boldsymbol{\theta}, \mathbf{y}) \propto \mathcal{N}(\mu_l(\boldsymbol{\theta}), \sigma_l^2(\boldsymbol{\theta})) \exp(\text{cubic spline}(x_l)) \quad (2.90)$$

with a cubic spline fitted to the difference of $\tilde{\pi}_{LA}(x_l|\boldsymbol{\theta}, \mathbf{y})$ and $\tilde{\pi}_G(x_l|\boldsymbol{\theta}, \mathbf{y})$ can be used (Rue et al. 2009). The third method, which is implemented as the default strategy in the INLA framework, is named the *simplified* Laplace approximation (Rue et al. 2009). This method uses a series expansion of $\tilde{\pi}_{LA}(x_l|\boldsymbol{\theta}, \mathbf{y})$ about the mean $x_l = \mu_l(\boldsymbol{\theta})$ to obtain the approximated density $\tilde{\pi}_{SLA}(x_l|\boldsymbol{\theta}, \mathbf{y})$ (Gómez-Rubio 2020). With this expansion, one can correct for skewness and location in the Gaussian approximation, while at the same time maintaining the computational advantages (Gómez-Rubio 2020). For the full derivations of the series expansion and the simplified Laplace approximation, see Rue et al. (2009, chapter 3.2.3).

2.6.3 Parameter estimation and sampling procedure

The parameter estimation procedure in INLA is composed of a number of steps. The mode of the log-likelihood $\ln(\tilde{\pi}(\boldsymbol{\theta}|\mathbf{y}))$ of the hyperparameters are obtained by maximizing with a quasi-Newton method. Then, to obtain the negative Hessian, \mathbf{H} , at the modal configuration $\boldsymbol{\theta}^*$, finite differences are applied (Gómez-Rubio 2020). The negative Hessian is then decomposed by its eigenvalues by $\mathbf{H}^{-1} = \mathbf{V}\Lambda\mathbf{V}^T$ and the hyperparameters are rescaled using \mathbf{z} such that

$$\boldsymbol{\theta}(\mathbf{z}) = \boldsymbol{\theta}^* + \mathbf{V}\Lambda^{1/2}\mathbf{z}, \quad (2.91)$$

to more effectively explore the hyperparameter space (Gómez-Rubio 2020). Then, the hyperparameter space is explored using either a regular grid with some step-size h or a central composite design (CCD) (Gómez-Rubio 2020, and references therein). The exploration is done to obtain a set $\{\boldsymbol{\theta}^1, \dots, \boldsymbol{\theta}^K\}$ that captures principal portion of the mass in the probability distribution (Martino & Riebler 2019). Once a set of hyperparameters is obtained, $\tilde{\pi}(x_l|\boldsymbol{\theta}, \mathbf{y})$ is approximated by $\tilde{\pi}_G(x_l|\boldsymbol{\theta}, \mathbf{y})$, $\tilde{\pi}_{LA}(x_l|\boldsymbol{\theta}, \mathbf{y})$ or $\tilde{\pi}_{SLA}(x_l|\boldsymbol{\theta}, \mathbf{y})$ and finally one can compute the desired marginal $\pi(x_l|\mathbf{y})$ using a numerical integration scheme on the form

$$\pi(x_l|\mathbf{y}) \simeq \sum_{k=1}^K \tilde{\pi}(x_l|\boldsymbol{\theta}^{(k)}, \mathbf{y}) \tilde{\pi}(\boldsymbol{\theta}^{(k)}|\mathbf{y}) \Delta_k. \quad (2.92)$$

A similar scheme for numerical integration can be used to obtain the marginals $\pi(\theta_k|\mathbf{y})$. Lastly, the joint posterior distribution can be approximated from the so-called Skew Gaussian Copula class, as specified in Chiuchiolo et al. (2021), and allows for sampling from the joint distribution.

The INLA method is implemented in the R-package **R-INLA** (Gómez-Rubio 2020) and is used in this thesis to fit the models and draw from the obtained posteriors. We note that for the random effects it is common to work with the precision matrix, which is defined as the inverse covariance matrix, rather than the covariance matrix directly. Therefore, all estimates on random effects will be given as precision rather than variance. Throughout the thesis, when INLA is applied, we will place PC priors on the model parameters, as recommended by Simpson et al. (2017) and Gómez-Rubio (2020).

2.7 Quantitative genetics and relative variable importance

An important application of GLMMs, which we will later analyse, is in the context of evolutionary biology and quantitative genetics. More specifically, one wishes to estimate the variance of the random effect which contributes to direct heritage of traits between relatives. Further, with this estimate, one uses its proportion of total model variance to evaluate the interaction between inheritance and environmental factors in developing distinct traits. We will now describe how this can be seen as a special case of wanting to estimate relative variable importance of random effects.

The Animal Model

To introduce the animal model and biological terminology, the section will rely heavily on the work of Kruuk (2004) and Conner & Hartl (2004). The animal model is a mathematical model, used as a tool for quantitative genetic analysis in evolutionary biology where the aim is to explain the phenotypic variation in a population. A phenotype is defined as "*the outward appearance of an organism for a given characteristic*" (Conner & Hartl 2004), such as eye color, height or behavior. In an organism, the observed phenotypic trait in an individual is a result of the complex combination of environmental effects and genotype. The genotype of a trait can be defined as "*the diploid pair of alleles present at a given locus*", and is the outcome of genetic inheritance (Conner & Hartl 2004). As evolutionary biology seeks to explain diversity among individuals in a population (Kruuk 2004), a decomposition of the phenotypic variance is of great interest. The simplest partition is to define the phenotypic variance as the sum of the genetic variance and environmental variance (Conner & Hartl 2004). However, for species that mate with other individuals in the population rather than self-fertilize, it is common to further decompose the genetic variance into three parts. The **total phenotypic variance** can therefore be partitioned as

$$\sigma_P^2 = \sigma_G^2 + \sigma_E^2 = \sigma_A^2 + \sigma_D^2 + \sigma_I^2 + \sigma_E^2 , \quad (2.93)$$

where σ_P^2 is the total phenotypic variance, σ_G^2 is the **genetic variance**, σ_E^2 is the **environmental variance**, σ_A^2 is the **additive genetic variance**, σ_D^2 is the **dominance genetic variance** and σ_I^2 is the **interaction genetic variance** (Conner & Hartl 2004). The parameter σ_A^2 , the variance of the additive genetic effect, is of particular interest, as the additive genetic effects are the only effects directly transferred to the offspring from its parents (Conner & Hartl 2004). Thus, the animal model aims to estimate σ_A^2 to gain inference on how changes in phenotypic values across generations occur, which is defined as phenotypic evolution (Conner & Hartl 2004). The animal model can be stated as a generalized linear mixed model, by letting

$$\boldsymbol{\eta} = g(\boldsymbol{\mu}) = \mathbf{X}\boldsymbol{\beta} + \mathbf{U}\boldsymbol{\alpha} , \quad (2.94)$$

where $\boldsymbol{\mu}$ is the mean of the observations \mathbf{y} of the phenotypic trait(s), $\boldsymbol{\eta}$ is the linear predictor, \mathbf{X} the design matrix of the fixed effects, $\boldsymbol{\beta}$ the population coefficients, \mathbf{U} the design matrix of the random effects and $\boldsymbol{\alpha}$ the vector of random effects.

One of the random effects in the animal model, $\boldsymbol{\alpha}_A \sim \mathcal{N}(0, \mathbf{G})$, accounts for the additive genetic effect. The values of the vector $\boldsymbol{\alpha}_A$ contains the so-called breeding values Wilson et al. (2010), which are defined as the effect of an individuals genes on the value of the phenotypic trait in its offspring (Conner & Hartl 2004) As in Section 2.3.2, we let \mathbf{G} denote the covariance matrix of the random effect $\boldsymbol{\alpha}_A$, which in the animal model can be derived from the expected covariance between relatives (Kruuk 2004). This derivation can be done by considering the coefficient of coancestry, $\Theta_{i,j}$, defined as "*the probability that an allele drawn at random from an individual i will be identical by descent to an allele drawn at random from individual j* " (Kruuk 2004). We use the coefficient of coancestry to define the expected covariance between relatives, or additive relationship matrix, as $\mathbf{A}_{i,j} = 2\Theta_{i,j}$ and consequently $\mathbf{G} = \sigma_A^2 \mathbf{A}$ (Kruuk 2004).

Heritability

As mentioned, we are particularly concerned in the additive genetic variance σ_A^2 and functions of it, such as the **heritability**. Heritability in the narrow sense, is defined as (Wilson 2008) the proportion of the total phenotypic variance that is present due to the additive genetic variance,

$$\frac{\sigma_A^2}{\sigma_P^2} = \frac{\sigma_A^2}{\sigma_A^2 + \sigma_D^2 + \sigma_I^2 + \sigma_E^2}. \quad (2.95)$$

The narrow sense heritability is what one considers for outbreeding species, and therefore, when we refer to heritability, we refer to the narrow sense heritability. In quantitative genetics, heritability is perhaps the most frequently estimated and discussed measure (Conner & Hartl 2004). Heritability has this role, as it can be used to partly explain how quickly the mean phenotypic values evolve, when populations are subject to artificial or natural selection (Conner & Hartl 2004). This is directly linked to the aim of quantitative genetics, which is to explain diversity and the cause of diversity (Kruuk 2004). As a subject of much misinterpretation, it is important to note that the definition of heritability is based purely on variance, and consequently heritability refers only to variation within a population. Further, as heritability is calculated for a specific population, environment and over time, it is not to be viewed a fixed value (Conner & Hartl 2004). Nonetheless, heritability is a widely used quantity to compare populations, species and traits, and is an important tool for understanding the evolutionary forces that drive genetic diversity and thereby evolution (Conner & Hartl 2004). The estimation process is often carried out using the animal model, and as heritability is the result of a variance decomposition of the model fit, we can connect it to variable importance. Recalling our preferred definition of the R^2 in Equation (2.59), (2.60), (2.61) and (2.62), one can quickly notice that the definition of heritability is very similar. In fact, generalizing the definition of variable importance from Grömping (2007) to also yield for random intercepts, one can define the heritability as the relative variable importance of the additive genetic effect. Therefore, estimating heritability can be seen as a special case of estimating relative variable importance, and serves as a suitable application for variable importance measures.

2.8 The Animal Model as a Gaussian Markov Random Field

INLA is a powerful tool for fitting latent Gaussian models (LGMs) as it provides a computationally efficient alternative to the traditional MCMC methods (Rue et al. 2009). To be applicable it relies heavily on the latent field, which is Gaussian, to possess the Markov property and thereby have the structure of a Gaussian Markov Random Field (GMRF). If a Gaussian random variable $\mathbf{X} = (X_1, \dots, X_n)$ possesses the Markov property it means that for some $i \neq j$, X_i is independent of X_j conditioned $X_{-i,j}$, where $X_{-i,j}$ denotes all other elements of \mathbf{X} except X_i and X_j (Rue et al. 2009). This property is readily visualized in a conditional independence graph (Figure 2.1, right), and in the animal model the pedigree structure (Figure 2.1, left) derived from the family relation can be used as the conditional independence graph (Wermuth & Lauritzen 1983, as cited in Steinsland & Jensen (2010)). The pedigree of a population is a directed acyclic graph (DAG) where each node represents an individual and the directed edges represent the parent-offspring relationship. This gives rise to the conditional independence graph, which can be found by inserting edges between parents that share offspring and removing the directions in the pedigree (Wermuth & Lauritzen 1983). An individual(node) in this graph will therefore only have edges, meaning it is conditionally dependent on, its parents, the parent(s) of its offspring, and its offspring. For example, in Figure 2.1 bird U_4 is conditionally dependent on birds U_1 and U_2 as they are its parents, bird U_6 as it is the offspring of U_4 and on bird U_5 as it is the other parent of the offspring of U_4 . U_3 and U_7 therefore does not provide additional information on U_4 (Steinsland & Jensen 2010). Therefore, this GMRF structure from the pedigree can be used to effectively sample and obtain parameter estimates from the animal model (Steinsland & Jensen 2010), and allows us to use the INLA framework for model fitting. The pedigree can also be used to construct the relatedness matrix \mathbf{A} , previously defined as the expected covariance between relatives, and the gives rise to the sparse precision matrix $\mathbf{Q} := \mathbf{A}^{-1}$ which is needed for calculations. As we consider each node an individual, the corresponding value of that node is its breeding value $\boldsymbol{\alpha}_\mathbf{A}$ (Steinsland & Jensen 2010).

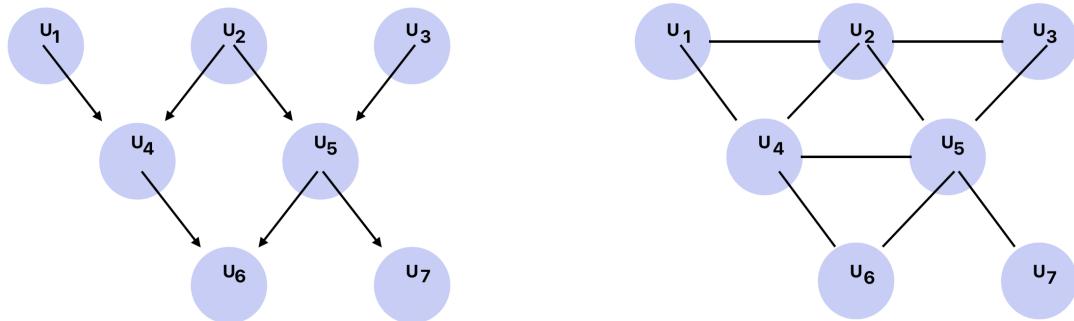


Figure 2.1: Illustration of a pedigree as a GMRF, figure and figure text inspired by Figure 1 in Steinsland & Jensen (2010). On the left, a pedigree structure is depicted as a directed acyclic graph (DAG), where birds U_1 and U_2 are the parents of bird U_4 , birds U_2 and U_3 the parents of bird U_5 , and birds U_4 and U_5 the parents of bird U_6 . Bird U_7 has one known parent in U_5 , and one unknown. On the right, the conditional independence graph of the pedigree structure is given, where the parents sharing offspring is assigned an edge and the direction is removed.

CHAPTER
THREE

METHODS

Based on the presented background theory, we now present our novel method for combining this into a relative variable importance tool for Bayesian GLMMs called Bayesian Variable importance (BVI). The proposed method is an extension of the method presented in Arnstad (2024) so that it now applies to GLMMs modelled with Binomial, Poisson in addition to Gaussian responses. The BVI method assumes the distinct random effects to be independent and does not include variable importance for random slopes.

For the complete model formulation of all methods used in this thesis, all files are uploaded to GitHub, with a link in Appendix A. If categorical covariates with more than two levels are contained in the fixed effects, they should be encoded using distinct names in order to make sure the method can handle them correctly.

3.1 Variable importance in the Bayesian framework

There are a few considerations necessary in order to calculate variable importance on GLMMs in a Bayesian framework. Firstly, the characteristics of the Bayesian framework must be considered. When fitting a GLMM in the frequentist framework, point estimates of the fixed regression coefficients as well as point estimates of the variance of the random effects are obtained. These estimates are then used to calculate relative variable importance measures. In contrast, a Bayesian GLMM tries to estimate the joint posterior distribution of parameters. From the posterior distribution, one can obtain samples of all parameters, that can be used to approximate a posterior distribution for each parameter. It is these samples that we will use for further calculations.

Secondly, we argue that the most intuitive way to calculate variable importance is on the link (or latent) scale. The reasoning behind this is the definition of residual variance for models with additive overdispersion in Nakagawa & Schielzeth (2013). This definition makes variable importance calculations on GLMMs analogous to that of LMMs, thus supporting a unified approach to both types of models. Therefore, we consider only GLMMs modeled with additive overdispersion, although we

believe the method could be extended to handle multiplicative overdispersion as well. These considerations are the basis of our proposed method for calculating relative variable importance in Bayesian GLMMs. The presented method can handle categorical variables with more than two categories as long as they are dummy encoded. Random slopes are excluded from our method due to the added computational complexity and the debatable improvement of GLMMs and R^2 values with random slopes as mentioned in Johnson (2014). We now go in to detail on how the different components of the GLMM model are handled in our method, to finally develop a relative importance measure for GLMMs.

3.2 Extending the R^2 to Bayesian GLMMs

The core of our Bayesian variable importance measures is a decomposition of the R^2 value so that each covariate is assigned a share of relative variable importance. We now combine the definition of the R^2 for GLMMs presented Section 2.4 and the R^2 for the Bayesian linear regression from Section 2.5.5 to yield our proposed distribution of the R^2 for Bayesian GLMMs. Consider the linear predictor

$$g(\mathbb{E}[\mathbf{y}]) = g(\boldsymbol{\mu}) = \boldsymbol{\eta} = \mathbf{X}\boldsymbol{\beta} + \mathbf{U}\boldsymbol{\alpha}, \quad (3.1)$$

for some response \mathbf{y} and monotonic and differentiable link function $g(\cdot)$. The variance components of the linear predictor can be decomposed into variance from the fixed effects and the random effects. Define the variance of the fixed effects as

$$\sigma_f^2 = \text{Var}(\mathbf{X}\boldsymbol{\beta}), \quad (3.2)$$

and let $\sigma_{\alpha_i}^2$ denote the variance of the i -th random effect, with random effects assumed independent. For Gaussian responses corresponding to an LMM, the residual variance σ_ε^2 is explicitly modelled as a parameter. The residual variance coincides with the overdispersion in the model, and the distributional variance with the identity link is zero (Nakagawa & Schielzeth 2013). However, for non-Gaussian responses, the residual variance of the model when considering additive overdispersion is defined as

$$\sigma_\varepsilon^2 = \sigma_e^2 + \sigma_d^2, \quad (3.3)$$

where σ_e^2 is the additive dispersion and σ_d^2 is the distributional variance (Nakagawa & Schielzeth 2013). A table containing the distributional variances for the link functions used in this thesis can be found in Table 3.1. Given that we can obtain samples for the variance components, we define for a sample s the marginal and conditional R^2 for the Bayesian GLMM as

$$R_{s,m}^2 = \frac{\sigma_{f,s}^2}{\sigma_{f,s}^2 + \sum_{i=1}^q \sigma_{\alpha_i,s}^2 + \sigma_{\varepsilon,s}^2} \quad \text{and} \quad R_{s,c}^2 = \frac{\sigma_{f,s}^2 + \sum_{i=1}^q \sigma_{\alpha_i,s}^2}{\sigma_{f,s}^2 + \sum_{i=1}^q \sigma_{\alpha_i,s}^2 + \sigma_{\varepsilon,s}^2}, \quad (3.4)$$

respectively, where $\sigma_{\varepsilon,s}^2 = \sigma_{e,s}^2 + \sigma_d^2$ is the sampled residual variance and σ_d^2 is distribution specific and the same for all samples. The posterior distribution of the marginal and conditional R^2 will then be approximated by the distribution of the samples of $R_{s,m}^2$ and $R_{s,c}^2$ for $s = 1, \dots, S$ respectively.

Distribution	Link Function	Parameter	σ_d^2
Normal	Identity	$\mu, \sigma^2 > 0$	0
Binomial	Logit	$0 < p < 1$	$\pi^2/3$
Poisson	Log	$\lambda > 0$	$\ln(1 + 1/\mathbb{E}[\lambda])$

Table 3.1: Distribution-specific variance σ_d^2 for the Gaussian, Binomial and Poisson distributions with link functions. The full expression $\mathbb{E}[\lambda]$ is given in (3.18). Distributional variances correspond to the variances in Nakagawa & Schielzeth (2013) and the calculation for the log-link Poisson follow the recommendations of Nakagawa et al. (2017).

3.3 Decomposing the R^2 value

We now seek to decompose the proposed R^2 value and assign each covariate with a proportion of the variance explained, i.e. assign each covariate with a *relative variable importance*. Recall that the fixed and random effects are assumed to be independent, so that one can consider the variances of the fixed and random effects separately. Further, the residual variance, if present, is also considered as independent of both fixed and random effects.

3.3.1 Applying the relative weights method in the Bayesian framework

To remedy the problems of calculating importance of correlated covariates, we will apply the relative weights method to the fixed effects before fitting the model. Following Section 2.2.4, we project the design matrix \mathbf{X} of the fixed effects to obtain the matrix \mathbf{Z} . The model is fit using \mathbf{Z} as an approximated design matrix of fixed effects, and from the joint posterior distribution samples of the coefficients $\boldsymbol{\beta}_{\mathbf{Z}}$ can be drawn. Each sample $\boldsymbol{\beta}_{\mathbf{Z},s}, s = 1, \dots, S$ and the matrix Λ can be used to approximate a sample of the importance of the columns \mathbf{X} , with the matrix Λ acting as weights from the projected space to the original covariate space. Using equations (2.19) and (2.20), we calculate this sample as

$$\text{IMP}(\mathbf{X})_s = \Lambda^{[2]} \boldsymbol{\beta}_{\mathbf{Z},s}^{[2]}, \quad (3.5)$$

where $\text{IMP}(\mathbf{X})_s$ is a column vector containing the approximated importance of column k of \mathbf{X} on the k -th entry for $k = 1, \dots, p$ and $\boldsymbol{\xi}^{[2]}$ again denotes the Schur product of $\boldsymbol{\xi}$ with itself. To calculate the relative variable importance, note that we estimate $\sigma_{f,s}^2$ in (3.4) by

$$\sigma_{f,s}^2 \simeq \sum_{k=1}^p \text{IMP}(\mathbf{X})_{s,k}. \quad (3.6)$$

Therefore, we define the relative importance of column k of \mathbf{X} in our method as

$$\text{RI}(\mathbf{X})_{s,k} = \frac{\text{IMP}(\mathbf{X})_{s,k}}{\sum_{j=1}^p \text{IMP}(\mathbf{X})_{s,j} + \sum_{i=1}^q \sigma_{\alpha_i,s}^2 + \sigma_{\varepsilon,s}^2}, \quad (3.7)$$

where $\sigma_{\alpha_i,s}^2$ and $\sigma_{\varepsilon,s}^2$ are defined as in Section 3.2. For sufficiently large S , we believe these samples can be used to construct an approximation of the posterior distribution of the relative importance for each fixed effect.

3.3.2 Random effects

The presented background theory on relative variable importance has mostly been developed for linear regression models. As long as the random effects are assumed not to be correlated, introducing random effects does not change the general idea. For each random effect, an approximation of the posterior distribution is constructed from the samples of the joint posterior distribution. Then, the proportion of variance explained by random effect i is calculated as

$$\text{RI}(\alpha_i)_s = \frac{\sigma_{\alpha_i,s}^2}{\sum_{k=1}^p \text{IMP}(\mathbf{X})_{s,k} + \sum_{k=1}^q \sigma_{\alpha_k,s}^2 + \sigma_{\varepsilon,s}^2}. \quad (3.8)$$

In addition to the relative importance of the random effects, a quantity of interest is the intraclass correlation, often also called the within cluster correlation or repeatability (Fahrmeir et al. 2013). The ICC represents the correlation between observations within the same cluster, and is defined for a random effect α_i in (Nakagawa et al. 2017) as

$$\text{ICC} = \frac{\sigma_{\alpha_i}^2}{\sum_{k=1}^q \sigma_{\alpha_k}^2 + \sigma_{\varepsilon}^2}. \quad (3.9)$$

Thus, following the same logic as before we can sample the ICC as

$$\text{ICC}_s = \frac{\sigma_{\alpha_i,s}^2}{\sum_{k=1}^q \sigma_{\alpha_k,s}^2 + \sigma_{\varepsilon,s}^2}, \quad (3.10)$$

and obtain an approximate posterior distribution of the ICC. It is also worth noting that the relative importance of the distributional variance are calculated in the same manner. Although not a quantity of interest, it can give an indication of how influential the added variance from the specific distribution is with respect to the covariates.

As previously mentioned, it is common to report the precision of random effects rather than the variance. Since the random effects are assumed to be independent, one can invert the precision estimate to obtain the variance. Another way of estimating the variance is to take the variance of the sampled values for the random vector α . Both methods seem to give very similar results as long as the sample size is large enough, and we therefore see both methods as fit for estimating the variance of random effects.

3.3.3 Drawing samples

A critical part in performing the calculations the BVI method requires, is to obtain samples from the joint posterior distribution. To do this, we utilize the built-in function from the INLA framework called `inla.posterior.sample()`. This function uses the approximation of the posterior distribution fitted with INLA by numerical integration, and therefore the accuracy of the samples is dependent on how well the numerical integration is carried out (Gómez-Rubio 2020). INLA provides several integration options, so one can choose the resolution one desires, but this comes at the cost of computational complexity. The BVI method is

implemented using the default integration strategy in INLA, which is either the grid strategy for a hyperparameter vector of dimension less than or equal to two or the central composite design (CCD) for a larger dimension hyperparameter vector (Martino & Riebler 2019). Further, if the model fit is poor or if the model is misspecified, the samples will suffer from this as well. Recall that INLA assumes a Gaussian latent layer, so this condition is crucial to obtain a representative set of samples. Lastly, INLA is a tool that is continuously in development, and the authors state that a skewness correction is in the works (Gómez-Rubio 2020).

3.4 Gaussian simulation study

To evaluate the performance of our proposed method a simulation study was conducted in Arnstad (2024), which we will reproduce here to provide a comprehensive overview. The study investigates how the BVI compares to the relative importance decomposition(Relaimpo, see package description in Grömping & Lehrkamp (2023)) as presented in Grömping (2007) and the two methods presented in Matre (2022). The Relaimpo method uses the LMG decomposition and considers only fixed effects and can therefore only be compared with the BVI in the fixed effects. The two methods in Matre (2022), ELMG and the ERW, are extensions of the LMG and relative weights methods respectively, to include random intercepts. These extensions allow us to compare the results for the random intercept model to our BVI method.

To simulate the data we consider the model as in (3.1), with the link function $g(\cdot)$ being the identity function. We have a sample size $n = 10^4$, $\boldsymbol{\alpha} = (\alpha_1, \dots, \alpha_m)$ where $\alpha_j \stackrel{iid}{\sim} \mathcal{N}(0, \sigma_\alpha^2 = 1)$ as a single random intercept for $m = 200$ clusters of $n_j = 50$ observations each, $\mathbf{X} \sim \mathcal{N}(\boldsymbol{\mu}, \Sigma) \in \mathbb{R}^{n \times p}$, where $\boldsymbol{\mu} = (1, 2, 3)$, $\Sigma_{ii} = 1$, $\Sigma_{i,k} = \rho_{i,k}$, $k \neq i$ and $p = 3$ consisting of three fixed effects, \mathbf{U} as a design matrix of appropriate dimension and a random error $\boldsymbol{\varepsilon}_i \stackrel{iid}{\sim} \mathcal{N}(0, \sigma^2 = 1)$. Further, the true vector of regression coefficients is set to be $\boldsymbol{\beta}_{\mathbf{X}} = (1, \sqrt{2}, \sqrt{3})^T$ so the total model, including an intercept column of ones, can be written as

$$\mathbf{y} = \mathbf{1} + \mathbf{X}\boldsymbol{\beta}_{\mathbf{X}} + \mathbf{U}\boldsymbol{\alpha} + \boldsymbol{\varepsilon}. \quad (3.11)$$

To investigate how different correlations between the fixed effects are handled by the method, we consider four different correlation levels between the fixed covariates in our data. That is achieved by letting $\rho_{1,2} = \rho_{1,3} = \rho_{2,3}$ take on the values $\{0, 0.1, 0.5, 0.9\}$. For each correlation level, we simulate $N = 1000$ datasets and fit each of the four methods BVI, Relaimpo, ELMG and ERW. To get a comparable measure from the Bayesian framework to the frequentist framework, we use the posterior means of the sampled posterior distribution of $\text{RI}(\mathbf{X})$ by the BVI method when estimating the quantities in (3.7) and (3.8).

From this setup, the theoretical variance of the response is

$$\text{Var}(\mathbf{y}) = \beta_{1,\mathbf{X}}^2 + \beta_{2,\mathbf{X}}^2 + \beta_{3,\mathbf{X}}^2 + 2 \sum_{i=1}^3 \sum_{k=i+1}^3 \beta_{i,\mathbf{X}} \beta_{k,\mathbf{X}} \rho_{ik} + \sigma_\alpha^2 + \sigma_\varepsilon^2, \quad (3.12)$$

and the theoretically correct relative importances for the case $\rho = 0$ are

$$\text{RI}(\mathbf{x}_1) = \beta_{1,\mathbf{x}}^2 = \text{RI}(\alpha) = \sigma_\alpha^2 = \frac{1}{8}, \quad \text{RI}(\mathbf{x}_2) = \beta_{2,\mathbf{x}}^2 = \frac{2}{8}, \quad \text{RI}(\mathbf{x}_3) = \beta_{3,\mathbf{x}}^2 = \frac{3}{8}, \quad (3.13)$$

as in Grömping (2007).

Further, the theoretically expected marginal R^2 values can be calculated as the variance of the fixed effects divided by the total variance given in 3.12. Similarly, the expected conditional R^2 are given by the sum of variance of the fixed effects and random intercepts divided by the total variance. The R^2 values are listed in Table 3.2.

ρ	R^2_{marg}	R^2_{cond}
0	0.750	0.875
0.1	0.781	0.890
0.5	0.852	0.926
0.9	0.889	0.945

Table 3.2: The theoretically expected marginal variance explained (left column) and conditional variance explained (right column) for different correlation levels between the fixed effects.

These values provide an empiric way of checking if our method fulfills the proper decomposition criteria listed in Section 2.2.1, by seeing if the relative importances for each effect sum to the model R^2 .

3.5 Heritability of phenotypic traits

As we have seen in Section 2.7, the concept of variance decomposition in GLMMs is not new and has been used in quantitative genetics with the animal model for many years (e.g. Kruuk (2004)). The main quantity of interest in such studies has been the heritability of phenotypic traits, which is defined as the ratio of additive genetic variance to total phenotypic variance (Wilson 2008). We now aim to illustrate how we calculate the heritability of phenotypic traits using the BVI method, and hence illustrating why heritability is a special case of variable importance. This involves modeling a pedigree covariance structure in random effects, which is a pivotal feature of the BVI method.

To demonstrate the broader inference possible with the BVI method, we have added the estimated posterior distributions of relative importance for all covariates in the supplementary material Appendix C. As heritability is a well-known quantity in quantitative genetics, there exists many other studies to compare our results with. Therefore, we investigate only the heritability estimates, but also want to emphasize that the BVI method is capable of providing more extensive inference on all covariates.

3.5.1 Heritability as relative variable importance

By comparing (2.95) with (3.8), it is clear that the way we have defined relative variable importance of a random effect coincides with the definition of heritability, if the random effect is the additive genetic effect and one assumes the total phenotypic variance σ_P^2 to be captured by the other fixed and random effects present. Therefore, when applying the BVI method to model a phenotypic trait, the relative variable importance of the random effect accounting for additive genetic variance can be interpreted as the heritability of the phenotypic trait. This is a highly relevant and useful application of our method and has been a major motivation for the development of the BVI method. It should be mentioned here that in the frequentist framework, fixed effects are assumed to not have an associated variance. Therefore, fixed effects are commonly not featured in formulae for the variance decomposition when estimating heritability (see Kruuk (2004) and Wilson et al. (2010)). Further, the discrimination between fixed and random effects are not always clear in biology. Often, no variance component of fixed effects is calculated. This means that they do not go into the calculation of the total phenotypic variance. However, there may be effects that are modelled as fixed, but still contribute to the phenotypic variance. To avoid confusion on this topic, we have implemented our method such that any covariate that contributes with variance in the model, is included in the calculation of total phenotypic variance. We see this to be the most clear and general way to handle the problem.

3.5.2 House sparrow study

We now apply the BVI method to a dataset gathered on house sparrows (*Passer domesticus*) from a study on the coast of Helgeland, Norway (Steinsland & Jensen 2010). The entire bird population on five islands have been surveyed since 1993 and several morphological traits have been measured. Blood samples were drawn to determine the relatedness between birds, and we therefore have a pedigree structure for the birds (Steinsland & Jensen 2010, citing Jensen et al., 2003, 2004, 2008). In the dataset we use we have $N = 3116$ birds with one or more observations on the traits and covariates. For a more thorough description of the house sparrow study, see Steinsland & Jensen (2010, and references therein). We model three phenotypic traits using a Gaussian LMM, namely the body mass, wing length and tarsus length. The fixed effects in the model consist of observations of *sex*, a standardized inbreeding coefficient denoted *FGRM*, the standardized *month* of the year (measurements were made during May-August), the *age* of each bird, and dummy variables encoding the location of the *native island* group of the bird (three levels, outer islands, inner islands or other islands). In addition, we model the *hatchyear* as an independent and identically distributed (i.i.d.) random intercept. To account for the correlation between relatives, we include a random effect for the additive genetic variance. It is the sampled variances of the additive genetic random effect that will determine the heritability of each trait. We derive the relatedness matrix of the birds from our pedigree, and specify this as the covariance matrix for the additive genetic variance term. Lastly, to account for individual differences we add an i.i.d. random intercept for the individual bird. We prefer to use the INLA framework, described in Section 2.6, to fit our LMM as it is computationally efficient and easy to use. Each prior is internally parametrized

in INLA by $\theta = \ln(\tau)$ with τ being the precision of the prior. This means when placing priors, they are always placed on the scale of the internal parameter θ , and if we want to place a prior on the external scale we must take this into account. For the fixed effects, we place penalising complexity (PC) priors with the initial value being $\ln(0.5)$ on the external scale and parameters $U = \sqrt{2}$ and $a = 0.05$ as the input parameters discussed in section Section 2.5.3. Similarly, we place PC priors on each random effect, with the effects *hatchyear* and *individual differences* having $U = 1$ and $a = 0.05$. The initial value of the priors for *hatchyear* and *individual differences* is set to be $\tau_0 = \ln(1)$ to correspond to 1 on the external scale. The additive genetic effect is assigned $U = \sqrt{2}$ and $a = 0.05$, with $\tau_0 = \ln(0.5)$. These priors have been chosen through discussion with the supervisor of the thesis and researchers with domain knowledge in biology. The approximation of the posterior marginals $\tilde{\pi}(x_l|\mathbf{y}\boldsymbol{\theta})$ will be made using the simplified Laplace approximation for all components, as described in Section 2.6.2 and Section 2.6.3. In this case, we also investigate the different integration strategies INLA provides. Two similar models are fit, with the only distinction being that one uses the grid integration strategy and the other applies the CCD strategy. We draw $N_{\text{samp}} = 10^4$ samples from the posterior distribution of the Bayesian GLMM fitted with the BVI method to estimate the posterior relative importances of the covariates.

3.6 Non-Gaussian studies

In this section, we present the methodology used to apply the Bayesian Variable Importance method to non-Gaussian GLMMs. This is a key extension of the method, as it allows the method to handle a wider range of models. We will analyse the Binomial and Poisson GLMMs, both via a simulation study and a case study.

3.6.1 Binomial and Poisson simulation studies

There are three primary reasons why we wish to conduct a simulation study with our method. The first being the ability to evaluate how well our method assigns relative variable importance to all covariates in the model. The real life case studies available mostly have the heritability, or some other function of the additive genetic variance, as the objective of analysis (Steinsland & Jensen 2010). We aim to provide the heritability, but at the same time provide information on the relative variable importance of all covariates present in the model. The second motivation is that the Bayesian framework is stochastic, and so is our method. We wish to assess the variability of this stochasticity by simulating different datasets with the same underlying structure, and see the spread of the estimates. We hope that this can provide signs that any fitted model can be seen as a random sample of a distribution centered around the true value. Lastly, the fundamental challenge that variable importance measures face, is the task of allocating the part of the variance explained due to correlation between covariates. Therefore, we wish to evaluate how our model performs for different correlation levels. This will give insight into how robust it is, and if the method handles correlated covariates in a sensible manner.

We simulate $N = 10^4$ responses from a Binomial distribution with a logit-link and from a Poisson distribution with log-link. The linear predictor contains three fixed effects and one random intercept. The fixed effects are, for simplicity but without loss of generalization, $\mathbf{X} \sim \mathcal{N}(\boldsymbol{\mu}, \Sigma)$ with $\boldsymbol{\mu} = (0, 0, 0)^T$, $\Sigma_{i,i} = 1$ and $\Sigma_{i,k} = \rho, k \neq i$. The true regression coefficient for the Binomial model are set to be $\boldsymbol{\beta} = (1, \sqrt{2}, \sqrt{3})^T$. In the Binomial model, the random effect $\boldsymbol{\alpha} = (\alpha_1, \dots, \alpha_m)$ comes from $m = 100$ clusters, each with $n_j = 100$ observations for $j = 1, \dots, m$. Further, we draw the random effect from a normal distribution such that $\alpha_j \stackrel{iid}{\sim} \mathcal{N}(0, \sigma_\alpha^2 = 1)$. This means that the total variance of the linear predictor $\boldsymbol{\eta}$ is $\sigma_\eta^2 = 7$. For the Poisson model, to avoid numerical instabilities, it was necessary to standardize the linear predictor used in the simulation study. Thus, the true regression coefficients were set to be $\boldsymbol{\beta} = \frac{1}{\sqrt{7}}(1, \sqrt{2}, \sqrt{3})^T$ and the random effect $\alpha_j \stackrel{iid}{\sim} \mathcal{N}(0, \sigma_\alpha^2 = 1/7)$. To investigate the impact of correlated fixed effects, we fit five different models letting ρ vary for each model by taking on the values $\rho \in \{-0.4, -0.1, 0, 0.1, 0.4\}$. The INLA framework is used to fit the GLMMs and the methodology described used to calculate the relative importance. All fixed and random effects receive the same PC prior as used in the comparison with the `rptR` package, that is with initial values of $\tau_0 = \ln(1)$ and parameters $U = 1$ and $a = 0.01$. As has been done throughout the thesis, the simplified Laplace approximation is used to approximate the posterior marginals of the latent field conditioned on the observations and hyperparameters. We fit $N_{\text{sim}} = 500$ Binomial and Poisson models with different datasets for each correlation level. For each fitted model, a sample of the posterior distribution is made to calculate the relative importance measures of all covariates. A single sample is drawn to analyse the variability of our method.

In the simulation study, when parameters are simulated so that we know their true value, we can empirically calculate the relative importance of the parameters when they are not correlated. When uncorrelated, the proportion of variance explained by each covariate in the linear predictor is equal to the square of the true coefficient divided by the total model variance. By defining $\sigma_{x_k}^2$ as the variance contribution to the linear predictor (the latent scale) for fixed effect k and σ_α^2 as the variance contribution of the random effect, we then have for the Binomial model with logit-link

$$\sigma_{x_1}^2 = \sigma_\alpha^2 = 1 \quad \text{and} \quad \sigma_{x_2}^2 = 2 \quad \text{and} \quad \sigma_{x_3}^2 = 3 , \quad (3.14)$$

and for the Poisson model with log-link

$$\sigma_{x_1}^2 = \sigma_\alpha^2 = 1/7 \quad \text{and} \quad \sigma_{x_2}^2 = 2/7 \quad \text{and} \quad \sigma_{x_3}^2 = 3/7 . \quad (3.15)$$

Then, the relative importance of the covariates can be calculated as

$$\begin{aligned} \text{RI}(\mathbf{X}_1) &= \text{RI}(\alpha_1) = \frac{\sigma_{x_1}^2}{\sum_{i=1}^3 \sigma_{x_i}^2 + \sigma_{\alpha_1}^2 + \sigma_d^2}, \\ \text{RI}(\mathbf{X}_2) &= \frac{\sigma_{x_2}^2}{\sum_{i=1}^3 \sigma_{x_i}^2 + \sigma_{\alpha_1}^2 + \sigma_d^2}, \\ \text{RI}(\mathbf{X}_3) &= \frac{\sigma_{x_3}^2}{\sum_{i=1}^3 \sigma_{x_i}^2 + \sigma_{\alpha_1}^2 + \sigma_d^2} . \end{aligned} \quad (3.16)$$

In our simulation study, the Binomial model with logit-link is assigned $\sigma_d^2 = \pi^2/3$. The distributional variance of the Poisson model with log-link is given by

$$\sigma_d^2 = \ln(1 + 1/\mathbb{E}[\mathbf{y}]) = \ln(1 + 1/\mathbb{E}[\lambda]) , \quad (3.17)$$

where

$$\mathbb{E}[\lambda] = \exp\left(\beta_0 + 0.5\left(\sum_{k=1}^q \sigma_{\alpha_k}^2 + \sigma_e^2\right)\right) , \quad (3.18)$$

is the quantity used in Table 3.1 (Nakagawa et al. 2017). So we obtain, using a single random intercept, $\sigma_d^2 = 0.6581$ with $\beta_0 = 0$, $\sigma_\alpha^2 = 1/7$ and $\sigma_e^2 = 0$. Therefore, we can summarize the expected relative importance of our three models as in Table 3.3. For the case of correlated covariates, no expected results for the relative importance of covariates are available, as there is no consensus on how to properly distribute the variance explained shared by covariates (Grömping 2007).

Model	$\mathbb{E}[\text{RI}(\boldsymbol{\alpha})]$	$\mathbb{E}[\text{RI}(\mathbf{X}_1)]$	$\mathbb{E}[\text{RI}(\mathbf{X}_2)]$	$\mathbb{E}[\text{RI}(\mathbf{X}_3)]$
Binomial, logit	0.0972	0.0972	0.1944	0.2915
Poisson, log	0.0861	0.0861	0.1723	0.2585

Table 3.3: The expected relative importance of the covariates in the different models when they are uncorrelated.

In practice, the distributional variance of the Poisson model should be calculated using the estimated values, and the distributional variance will therefore be dependent on the fitted model (Nakagawa et al. 2017).

In addition to the expected importance of covariates in the uncorrelated case, we can calculate the expected marginal and conditional R^2 values for all correlation levels on the latent scale. Recalling that each of the $p = 3$ columns of \mathbf{X} is initialized to have variance equal to 1, the expected marginal R^2 can be calculated as

$$\mathbb{E}[R_{\text{marg}}^2] = \frac{\sum_{i=1}^3 \beta_i^2 + 2 \sum_{i=1}^2 \sum_{k=i+1}^3 \beta_i \beta_k \rho}{\sum_{i=1}^3 \beta_i^2 + 2 \sum_{i=1}^2 \sum_{k=i+1}^3 \beta_i \beta_k \rho + \sigma_\alpha^2 + \sigma_d^2} , \quad (3.19)$$

and similarly for the expected conditional R^2 as

$$\mathbb{E}[R_{\text{cond}}^2] = \frac{\sum_{i=1}^3 \beta_i^2 + 2 \sum_{i=1}^2 \sum_{k=i+1}^3 \beta_i \beta_k \rho + \sigma_\alpha^2}{\sum_{i=1}^3 \beta_i^2 + 2 \sum_{i=1}^2 \sum_{k=i+1}^3 \beta_i \beta_k \rho + \sigma_\alpha^2 + \sigma_d^2} . \quad (3.20)$$

Model Type	Correlation (ρ)	$\mathbb{E}[R^2_{\text{marg}}]$	$\mathbb{E}[R^2_{\text{cond}}]$
Binomial Logit	-0.4	0.262	0.434
Binomial Logit	-0.1	0.532	0.641
Binomial Logit	0	0.583	0.680
Binomial Logit	0.1	0.624	0.712
Binomial Logit	0.4	0.709	0.777
Poisson Log	-0.4	0.214	0.354
Poisson Log	-0.1	0.465	0.561
Poisson Log	0	0.517	0.603
Poisson Log	0.1	0.559	0.638
Poisson Log	0.4	0.651	0.714

Table 3.4: Expected marginal and conditional R^2 values for the Binomial regression with logit-link (top) and Poisson regression with log-link (bottom) for different correlation levels ρ .

3.6.2 Binomial and Poisson case studies

To investigate how well the BVI method generalizes to non-Gaussian responses, we perform a case study using the setup described in the vignette of the R-package `rptR`, found at <https://cran.r-project.org/web/packages/rptR/vignettes/rptR.html> (Stoffel et al. 2017). This package estimates the repeatability of phenotypic traits, which is closely related to heritability. An important clarification for this case study, is that there are multiple formulations of repeatability. Two of the most common ways of looking at repeatability are

$$\begin{aligned} R_1 &= \frac{\text{Additive genetic variance}}{\text{Total variance of covariates}} \\ R_2 &= \frac{\text{Additive genetic variance}}{\text{Total variance of random covariates}} , \end{aligned} \quad (3.21)$$

where the former corresponds to our notion of heritability (Stoffel et al. 2017) and the latter to the ICC (Fahrmeir et al. 2013). We choose to look at the notion corresponding to heritability, and to obtain the result from `rptR` so that they match this, each model must be fit with the argument `adjusted=FALSE` (Stoffel et al. 2017). The dataset used in the `rptR` package vignette, introduced for a different purpose, is simulated to replicate a study on twelve different beetle larvae populations (Stoffel et al. 2017). It contains $N = 960$ observations of the covariates *population*, the discrete *habitat* of the larvae, the dietary *treatment* of the larvae, the *sex* and *container* of which the larvae were contained in. The phenotypes to be modeled by the Binomial and Poisson distributions are the two distinct male color morph and the number of eggs laid by female larvae respectively. Both models use *treatment* as the only fixed effect and place i.i.d. random intercepts on the *population* and *container* covariates. Note that a more complex covariance structure could be modelled by the BVI method, but the `rptR` package does not allow for this, so for comparing the methods we see it as suitable with i.i.d. random intercepts. As before, our modelling is carried out using INLA, whereas the models in the vignette are calculated from functions in the `rptR` package. The priors

placed on the fixed effect *treatment* and random effects *population* and *container* are PC priors with initial values $\tau_0 = \ln(1)$ on external scale and parameters $U = 1$ and $a = 0.01$ for all effects. As before, our preferred approximation of the posterior marginals of the latent field conditioned on the observations and set of hyperparameters is the simplified Laplace approximation. Furthermore, we also here draw $N_{\text{samp}} = 10^4$ samples from the posterior distribution of the Bayesian GLMM fitted with the BVI method to estimate the posterior distributions of the repeatability.

3.7 Simulation study with R^2 -induced Dirichlet decomposition priors and Generalized Decomposition Priors on R^2

As mentioned, the Bayesian variable importance field is not very large. However, as discussed in Section 2.5.6, the R2D2 priors decompose the R^2 value and can therefore be interpreted as a variable importance measure. We find it sensible to try and compare the resulting variable importance distributions, even though the R2D2 priors have not been developed for this goal specifically. To be able to evaluate the two measures, we simulate a linear regression model with $p = 3$ covariates and $n = 1000$ observations. The covariates are for simplicity simulated as $\mathbf{X} \sim \mathcal{N}(\boldsymbol{\mu}, \boldsymbol{\Sigma})$ with $\boldsymbol{\mu} = (0, 0, 0)^T$, $\boldsymbol{\Sigma}_{i,i} = 1$ and $\boldsymbol{\Sigma}_{i,j} = \rho$ for $i \neq j$. As before, we vary the correlation by letting $\rho \in \{-0.4, -0.1, 0, 0.1, 0.4\}$. The true regression coefficients are initialized as $\boldsymbol{\beta} = (1, \sqrt{2}, \sqrt{3})$, and we simulate a random error by $\varepsilon \sim \mathcal{N}(0, \sigma^2 = 1)$. By noting that $\text{Var}(\mathbf{y}) = 7$, it is clear that in the uncorrelated case, the relative importance of the covariates can be calculated as

$$\text{RI}(\mathbf{X})_1 = \frac{1}{7} \quad \text{RI}(\mathbf{X})_2 = \frac{2}{7} \quad \text{RI}(\mathbf{X})_3 = \frac{3}{7}. \quad (3.22)$$

Further, the R^2 for this model is by the definition in (2.8)

$$R^2 = \frac{\text{Var}(\mathbf{y}) - \sigma^2}{\text{Var}(\mathbf{y})}, \quad (3.23)$$

which gives values that are summarized in Table 3.5.

ρ	R^2
-0.4	0.604
-0.1	0.830
0	0.857
0.1	0.877
0.4	0.913

Table 3.5: Expected R^2 values for the correlation levels ρ used in the linear regression for analyzing the BVI method in comparison to the R2D2 priors.

To fit the linear regression using R2D2 priors for the marginal R^2 we use functions from the GitHub repository Zhang (2024) by the author of Zhang et al. (2020).

For the GDR2 priors, we utilize the Stan code available on Romero & Bürkner (2024) by the authors of Aguilar & Bürkner (2024). The hyperparameters for the marginal $R^2 \sim \text{Beta}(a, b)$ are set so that $\mathbb{E}[R^2] \simeq 0.857$ which is approximately the theoretical R^2 of 6/7. This is done for the R2D2 priors by using the default value $b = 0.5$ from Zhang (2024) and noting that the expected value of the $\text{Beta}(a, b)$ distribution is $a/(a+b)$ (Tjelmeland et al. 2000). We follow the Gibbs sampler for the marginal R2D2 prior as described in (Zhang et al. 2020, section 5.3) and run the MCMC iteration $N = 10^4$ times, with a burn in of 9000 samples. This gives 1000 samples from the posterior distribution of the marginal R^2 as well as 1000 samples of each ϕ_j for $j = 1, 2, 3$. The BVI draws the same amount of samples from the posterior distribution of the fitted Bayesian GLMM. For the GDR2 priors, the implementation requires a prior on the expected value and the precision of the R^2 value directly. These are calculated by letting $a_\pi = 0.7$, the reference unit α_K be zero, the expected R^2 equal 6/7 and then solving for the precision τ according to the properties of the Beta distribution given in Aguilar & Bürkner (2024). The choice of a_π corresponds to a scenario in which one assumes the covariates to be approximately equally important and the difference between the GDR2 prior and R2D2 prior is substantial (Aguilar & Bürkner 2024). Similarly, as for the R2D2 case, we run the MCMC iteration $N = 10^4$ times, with a burn in of 9000 samples, but we also fit four Markov chains this time to ensure proper mixing. This means we obtain 4000 samples for the GDR2 priors. The samples of ϕ_j from the linear regression using R2D2 and GDR2 priors are then seen as the posterior distribution of relative variable importance of \mathbf{x}_j , and compared to that of the BVI method. We will refer to the results by using R2D2 and GDR2 priors as the R2D2 method and the GDR2 method respectively. To evaluate all methods, we fit a frequentist linear regression model and evaluate the importance metrics according to the Relaimpo method by using the package `Relaimpo` (Grömping & Lehrkamp 2023) in R as described in Grömping (2007). As the Relaimpo method implements the LMG, which is feasible in this context, it poses perhaps the most robust and reliable benchmark available. We draw 1000 bootstrap samples of the LMG metrics and use this to create confidence intervals for the Relaimpo method.

RESULTS

4.1 Gaussian simulation study

As the simulation study from Arnstad (2024) for Gaussian LMMs was included in the previous chapter, this section presenting the results from it also follow from that Arnstad (2024). To present the results of the simulation study we consider each effect separately in different plots, that is we show results for the importance of variables X_1, X_2, X_3 and the random effect α , in distinct plots. We used violin plots to visualize the estimated quantities, as they contain much information in a compact way. The violin plot is analogous to a density plot, but the density is shown along the y -axis and mirrored about the y -axis to form a symmetrical shape. Each violin therefore displays the distribution of our simulated estimates. Lastly, we consider how our BVI model estimates the conditional and marginal R^2 values from a Bayesian perspective, comparing these results to the R^2 value obtained from the decompositions by the Relaimpo, ELMG, and ERW methods, where the first method only considers models without random effects.

4.1.1 Relative importance of the fixed effects

Here, the distribution of the relative importance allocated to each fixed effect from the simulations are shown. There are four different distributions for each method, which corresponds to the four different correlation levels. The horizontal dashed line displays the theoretically correct relative importance from (3.13) when the covariates are pairwise independent.

In general, it can be seen (Figure 4.1) that the distributions in the case of uncorrelated data are unbiased with some variation around the theoretically correct relative importance. For a correlation of $\rho = 0.1$ the distributions of the estimates are shifted marginally compared to the uncorrelated case for all methods. The importance attributed to X_1 and X_2 , in Figure 4.1, is larger when compared to the uncorrelated case, whereas the importance attributed to X_3 is smaller. All methods seem to shift the relative importance estimate for the covariate with the same amount in the same direction. This shift is both expected and desirable, when considering the values found in Table 3.2 for the theoretically correct variance explained. Therefore, we should expect our method to assign different shares when

we have various levels of covariate correlation, which it does. This trend continues for the correlation level $\rho = 0.5$, where the distributions are shifted further in the same directions as for $\rho = 0.1$. Lastly, for $\rho = 0.9$ we see the largest reallocation of the distributions, which follows the same trend as for the other correlation levels.

The rise in importance for X_1 and X_2 for increasing correlation can be understood by the relation $\mathbf{Z}\Lambda = \mathbf{X}$ in the relative weights method. When the matrix \mathbf{X} is not correlated, Λ is close to the identity matrix, but with an increase in correlation the diagonal elements grows smaller and off diagonal elements grow larger. An increase in off diagonal values would for X_1 and X_2 imply that a larger value is multiplied with β_3^2 , which is larger than β_1^2 and β_2^2 . Therefore, it is expected to see a rise in importance as correlation increases for X_1 and X_2 , and the opposite for X_3 . In all figures, the BVI method is in agreement with the other methods when allocating importance for different correlation levels. The width of the distributions seem to become lower as the correlation increases, most notably for $\rho = 0.9$, where the distributions exhibit significantly smaller dispersion than for $\rho = 0$. Generally all methods seem to follow the same trends and produce similar results for all three fixed effects. As correlation increases the trend is that the relative importance assigned to X_1 and X_2 increases, in contrast to the decrease in relative importance assigned to X_3 .

In general, the BVI method is in agreement with the theoretical results for uncorrelated data derived in Chapter 3 and is consistent with the other three methods for correlated fixed effects.

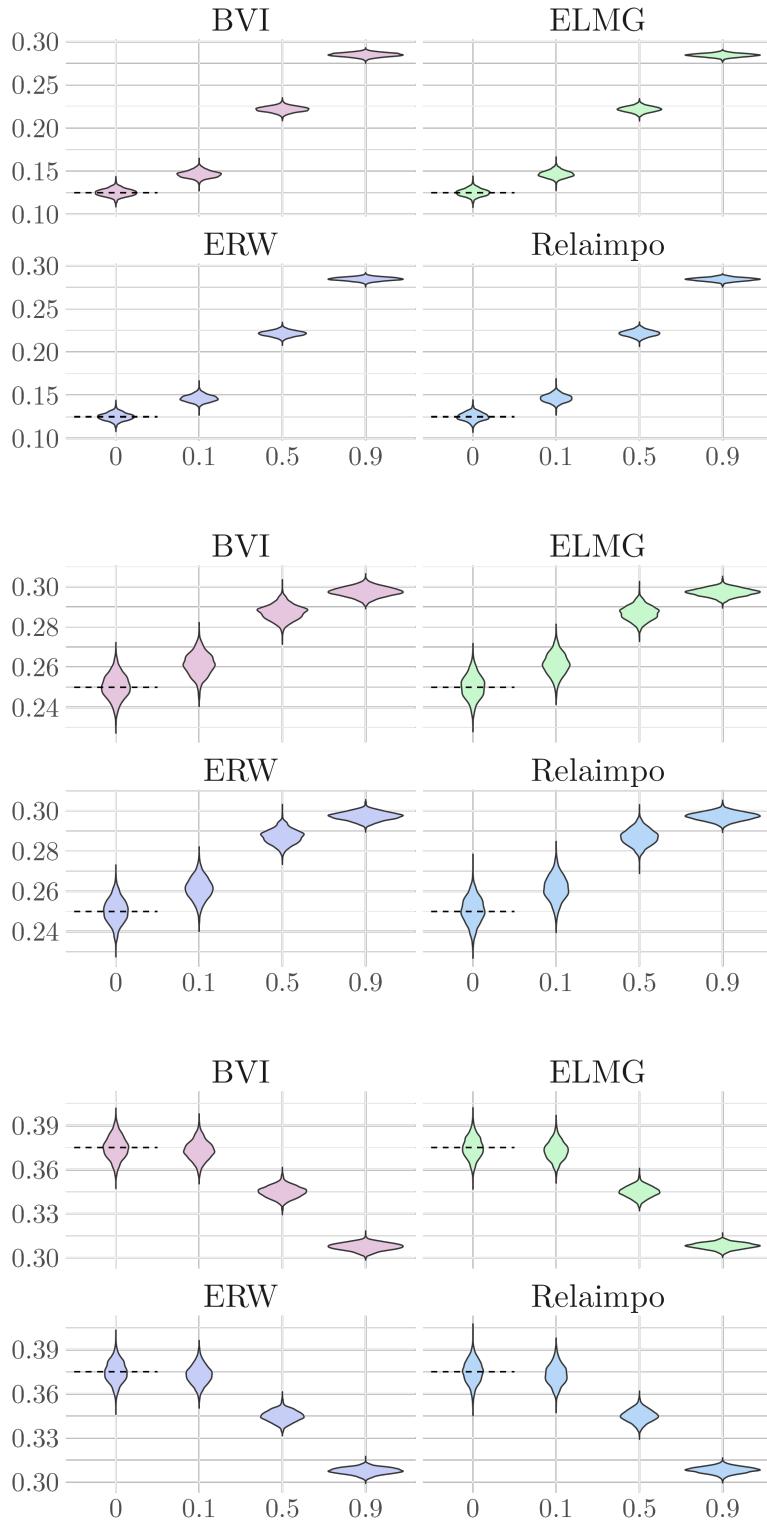


Figure 4.1: Violin plots for the relative importance of the fixed effects X_1 (top), X_2 (middle) and X_3 (bottom) for different correlation levels displayed along the x-axis, calculated from the ensemble of simulated datasets by the BVI, ELMG, ERW, and the Relaimpo methods. The horizontal line displays the theoretically correct importance of each fixed effect in the case of uncorrelated data. For the BVI method, the distributions of posterior means are shown to compare to the distribution of point estimates from the other three methods.

4.1.2 Relative importance of the random effects

Considering a model with one random intercept, we can no longer compare our model with the Relaimpo method, which is only implemented for the linear regression in the Relaimpo R package (Grömping 2007). Therefore, we now compare the BVI method only with the ELMG and ERW methods, which have been extended from the Relaimpo method (Matre 2022). We display (Figure 4.2) the distribution of the relative importance, or variance, assigned to the random intercept α for different correlation levels. The random intercept α follows a univariate normal distribution with mean zero and variance equal to 1. As before the horizontal line shows the theoretical relative importance from (3.13) that α has in the model when the fixed effects are uncorrelated.

It is apparent (Figure 4.2) that both the location and width of the relative importance distribution of all methods are largely indistinguishable. The distributions take on a moderately smaller value when $\rho = 0.1$ and the location of the estimates is further decreased for $\rho = 0.5$ and $\rho = 0.9$. For the latter correlation level, the distributions are located around a value that is less than half of the value of the centering when the fixed effects are uncorrelated. To re-emphasize, this is both expected and desirable since the increase in response variance comes solely from the correlation of fixed effects, so the random effects now contribute to explain a smaller proportion of the variance, *i.e.* the importance is lower.

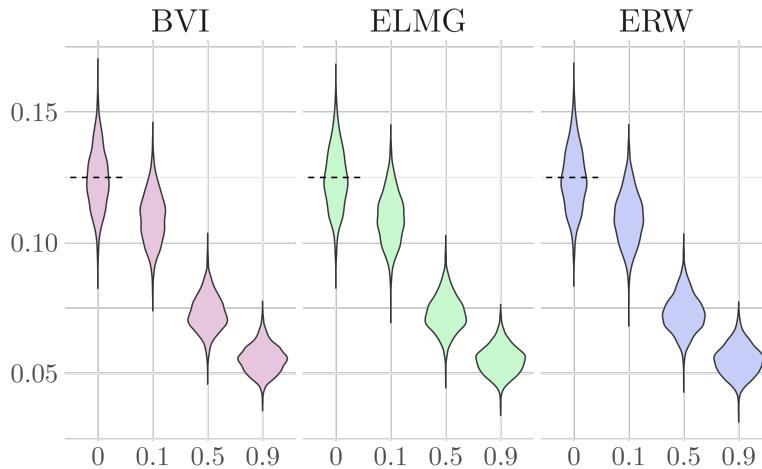


Figure 4.2: Violin plots for the relative importance of the random effect α , that is, $\hat{\sigma}_\alpha^2$ for different correlation levels displayed along the x-axis, calculated from the ensemble of simulated datasets by the BVI, ELMG and the ERW method. For the BVI method the distributions of posterior means of the marginal distribution of $\hat{\sigma}_\alpha^2$ are shown to compare to the point estimates of the other two methods. The horizontal line displays the theoretically correct importance σ_α^2 of the random effect in the case of uncorrelated data.

4.1.3 Total variance explained - R^2 estimates

As a useful by-product from the previous results we can get the total variances explained by our model (Figure 4.3). The marginal variance explained, R_{marg}^2 , is

the variance explained by the fixed effects, and we get results for all four methods, including Relaimpo. Below, the marginal variance explained and the total conditional variance explained, R^2_{cond} , is displayed. This is the variance given all the fixed effects and the random effect. To complement the conditional and marginal variances explained, a horizontal line is drawn for each correlation level corresponding to the theoretically correct variance explained, found in Table 3.2, for the correlation level.

We see that the four methods produce very similar results of R^2_{marg} for the fixed effects across all correlation values, albeit a slightly larger width for the BVI method can be seen. When considering the conditional variance R^2_{cond} , the dispersion of the BVI method is strikingly larger compared to the other methods. It is not immediately clear why, but it could be a result of the relatively large dispersion of the estimated posterior marginal variance of α . Both the marginal and the conditional variance are centered around the theoretically correct value with some variability, particularly visible for conditional variance of the BVI method. The centering of the distributions for both the marginal and conditional variances resemble each other for all methods, regardless of correlation level.

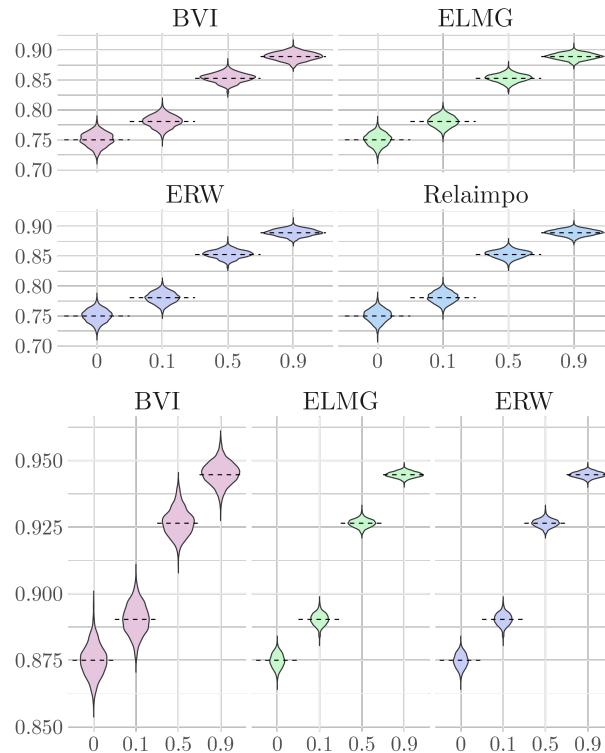


Figure 4.3: Violin plots for the total marginal (top) and conditional (bottom) variance explained (R^2) for different correlation levels displayed along the x-axis, calculated from the ensemble of simulated datasets by the BVI, ELMG, the ERW and the Relaimpo method(only marginal variance explained can be computed). For the BVI method the posterior means of the sampled posterior distributions of β and the marginal distribution of $\hat{\sigma}_\alpha^2$ in each simulation are used to compare to the point estimates of the other two methods. The horizontal lines display the theoretical explained variance for each correlation level ρ as in Table 3.2.

4.2 Heritability of house sparrow traits

We now investigate the results of applying our method to the house sparrow dataset. As previously discussed (Section 2.7), estimating the heritability of phenotypic traits can be seen as a special case of relative variable importance and so the findings we present are directly obtained by our method. The samples of relative variable importance, are sampled from the variance component that captures additive genetic variance, and we use the results from Silva et al. (2017) and Muff et al. (2019) for comparison. For this analysis, the covariance structure of the pedigree required us to model more complex random effects than i.i.d. random intercepts, and so the `rptR` package could not be used for comparisons. In Table 4.1 the mean of sampled heritability along with quantile values is presented, as well as the corresponding measures from the comparable studies.

	h_{mass}^2		h_{wing}^2		h_{tarsus}^2	
	Est.	CI	Est.	CI	Est.	CI
Silva et al. (2017)	0.300	[0.231, 0.369]	0.388	[0.353, 0.461]	0.415	[0.333, 0.497]
Muff et al. (2019)	0.288	[0.219, 0.371]	0.344	[0.294, 0.409]	-	-
BVI grid	0.282	[0.231, 0.342]	0.356	[0.322, 0.393]	0.401	[0.329, 0.468]
BVI CCD	0.284	[0.255, 0.315]	0.356	[0.334, 0.380]	0.401	[0.363, 0.438]

Table 4.1: Heritability estimates and confidence intervals from Silva et al. (2017), posterior means of additive genetic variance divided by the posterior means of total phenotypic variance in Muff et al. (2019) with corresponding confidence intervals, and the mean and quantile values of the heritability samples obtained from the BVI method with grid and CCD integration strategies for the phenotypic traits: body mass, wing length, and tarsus length. Note that in Muff et al. (2019) no estimate of tarsus length heritability was provided.

For the sampled heritability of body mass (Figure 4.4), we have a mean of 0.282 (Table 4.1) for the grid strategy and 0.284 for the CCD strategy. Both distributions are centered around the mean, with the CCD strategy having a smaller spread. This is also seen from the 95th percentile, which is approximately the interval [0.231, 0.342] for the grid strategy compared to [0.255, 0.315] for the CCD strategy. The grid strategy provides a distribution that does not seem to be fully symmetric, having a longer tail to the right than on the left. The CCD strategy, on the other hand, gives a more symmetric distribution with lighter tails. Perhaps a longer tail to the right is observable also here. To fit both the grid and CCD model and obtain the samples, the BVI method took about 156 seconds.

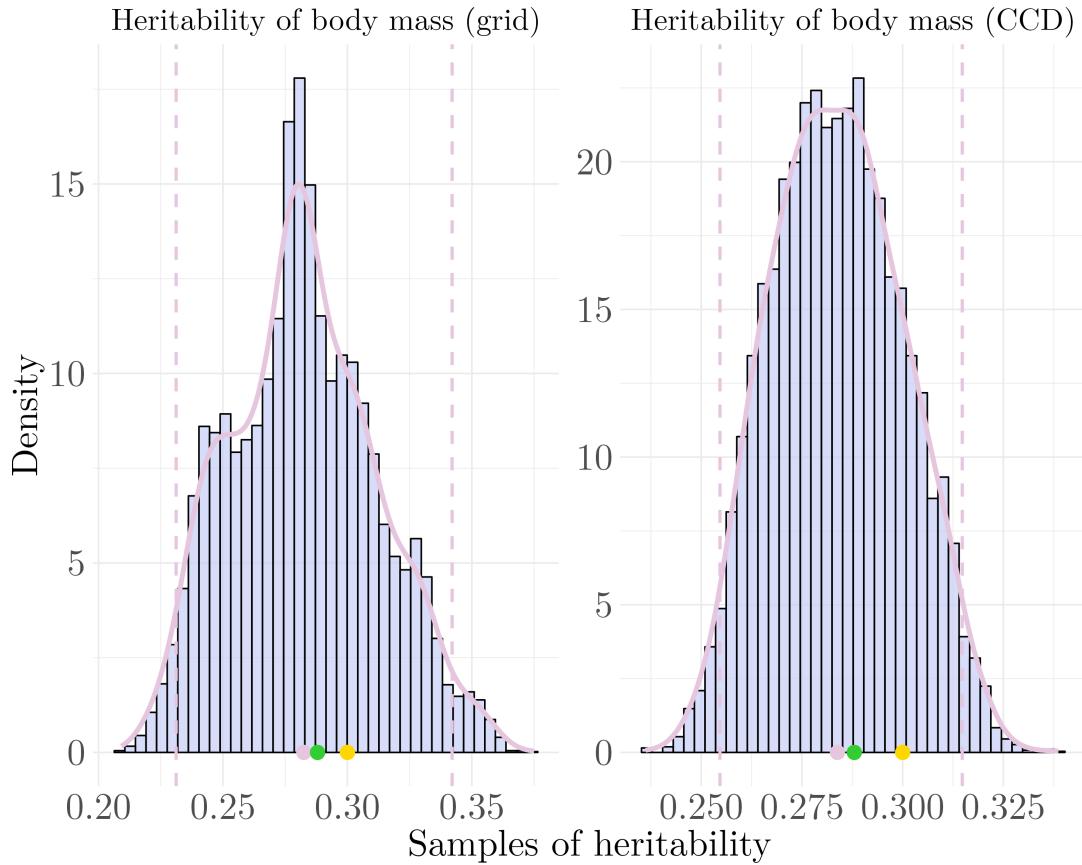


Figure 4.4: Histogram depicting the estimated heritability values of body mass by the BVI method for the grid integration strategy(left) and CCD integration strategy (right) for the house sparrow dataset. The mean of the samples is marked as a pink circle at the bottom of the histogram, with the lower and upper value for the 95% percentile marked as dashed lines. The heritability estimate from Silva et al. (2017) and Muff et al. (2019) are marked as gold and green dots respectively at the bottom of the histogram.

The samples of wing length heritability both form a roughly symmetric curve, with the grid and CCD samples centered around the same mean of 0.356. The grid strategy has again a slightly larger spread, with the 95th percentile being approximately the interval [0.322, 0.393], compared to the CCD strategy with the interval [0.334, 0.380]. One could argue that for the grid strategy, the right tail is a bit longer also for the heritability of wing length. The samples of heritability for wing length therefore exhibits less dispersion than those for heritability of body mass. In this case, the model fit and sampling procedure took 162 seconds for both integration strategies in total.

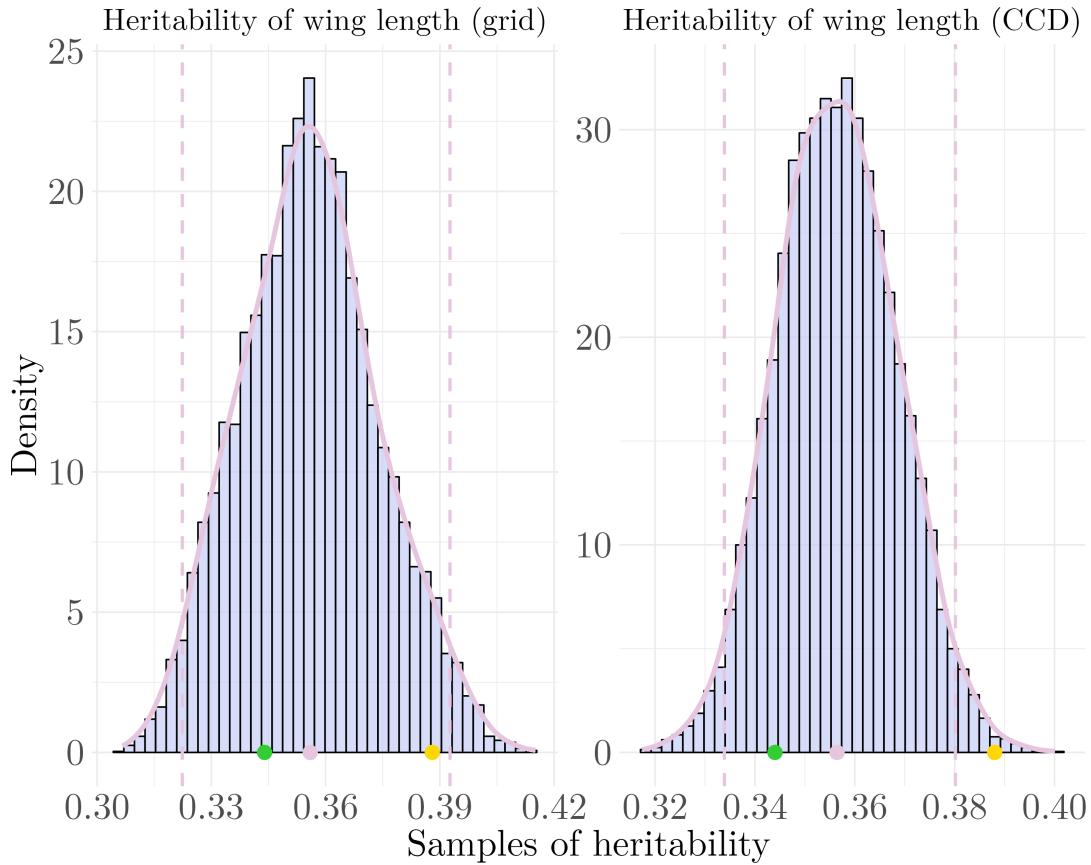


Figure 4.5: Histogram of heritability values for wing length of the house sparrows estimated by the BVI method for the grid integration strategy (left) and the CCD integration strategy (right). The mean of the samples is marked as a pink circle at the bottom of the histogram, and the lower and upper value for the 95% percentile are featured as dashed lines. The heritability estimate from Silva et al. (2017) and Muff et al. (2019) are marked as gold and green dots respectively at the bottom of the histogram.

The heritability samples of tarsus length (Figure 4.6) has a mean of 0.401 (Table 4.1) for both integration strategies. Both are centered at this value, but while the CCD strategy gives of an approximately normal, unimodal distribution, the grid strategy gives a trimodal distribution. The peaks of the trimodal distribution are distinct, with the center peak being the highest and the right and left peaks symmetric about the center peak at equal height. A possible explanation for this pattern is that for this trait, the grid used for numerical integration might force the sampling to occur most frequently at the three modal values. Again, we see that the grid strategy exhibits a larger spread than the CCD strategy. The 95th percentile of the grid strategy is captured by the interval [0.329, 0.468], whereas the percentile of the CCD strategy is covered by the interval [0.363, 0.438]. For the heritability of tarsus length, the quantiles for both strategies are larger than the quantiles for heritability of body mass and wing length. The time spent to fit the model and draw the samples was reported to be 151 seconds.

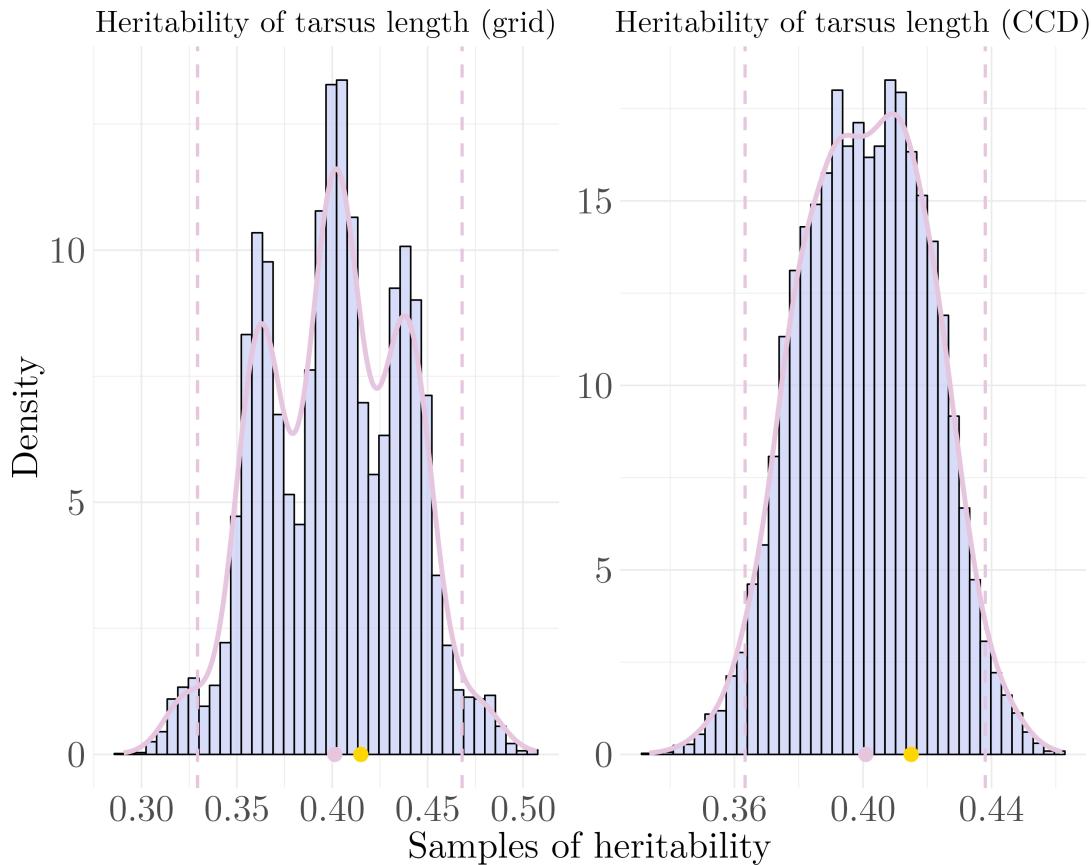


Figure 4.6: Histogram showing estimated heritability values for tarsus length of the house sparrows from the BVI method for the grid integration strategy (left) and the CCD integration strategy (right). The two dots at the bottom represent the mean of the samples (pink) and the estimate from (Silva et al. 2017) (gold). The dashed lines represent the lower and upper value for the 95% percentile.

We see it as natural that some different patterns that are hard to fully interpret occur, as the dataset is from real life and relatively small. Further, the measurements are taken on birds that are quite small, so one should expect measurement error to some degree.

4.3 Non-Gaussian simulation study

In this section, we lay forth the results of our simulation study on a Binomial and a Poisson regression. In this study, we draw one sample of the posterior distribution for each fitted model to construct the estimates presented. We note that it has been difficult to find suitable methods to compare the non-Gaussian models with. In parallel to fitting our model as described in Section 3.6.1, we fit a model using the `rptR` package with 100 bootstrap samples. This allows us to directly compare the importance of the random effect and the marginal and conditional R^2 values. However, it does not compute the importance of each isolated fixed effect.

4.3.1 Binomial simulation

We begin by presenting the results obtained from the simulation study. The first model to be analysed is the Binomial regression on binary response, modelled with the logit-link function. For each correlation level, we draw a sample from the resulting posterior distribution using the Bayesian Variable Importance method. Then, the BVI method calculates the derived measures from Chapter 3 to estimate the relative importance of all covariates in each model. These estimates thereby construct the estimated posterior distributions of the relative importance measures of all covariates. A summary of the 500 estimated importances are shown in the supplementary material (Appendix C, specifically Table C.1), which contains the mean and values for the lower and upper 95% quantile.

4.3.1.1 Fixed effects

The sampled posterior distribution of relative importance allocated to the three fixed effects X_1 , X_2 and X_3 are shown for each correlation level (Figure 4.7). We see that the distributions generally form a normal shape around the mean, with somewhat varying spread. As correlation levels go from negative to positive, meaning that the variance contribution from the fixed effects increase, the importances of the fixed effects also increase. This is expected, as the shared covariance increases and is spread across the correlated fixed effects. The difference is quite substantial, with the average relative importance allocated to X_1 for $\rho = -0.4$ being 0.0200 compared to 0.1724 for $\rho = 0.4$. The same pattern is seen for X_2 and X_3 , with the average relative importance increasing from 0.0769 to 0.2392 for X_2 and from 0.1666 to 0.2984 for X_3 when going from $\rho = -0.4$ to $\rho = 0.4$. For $\rho = 0$ (middle plot of Figure 4.7), it is clear that the average estimate for relative importance of all fixed effects is very similar to the expected importance (Table 3.3) shown as a dashed green line.

We notice that the covariates X_1 and X_2 are allocated a significantly larger share when correlation goes from $\rho = 0$ to $\rho = 0.4$, whereas X_3 is almost unchanged for the same correlation levels. This was also experienced in the simulation study on LMMs in Section 4.1 from Arnstad (2024). It is explained by the fact that off-diagonal elements of Λ increase positively when the fixed effects are positively correlated, while the diagonal elements decrease. In the uncorrelated case, Λ should be equal to the identity matrix. The columns of Λ therefore act as weights and due to this, when $\rho = 0.4$, X_1 will receive an importance estimate where β_2^2 and β_3^2 will have positive weights contrary to $\rho = 0$ where the only weight is put on β_1^2 . Since β_1^2 is smaller than β_2^2 and β_3^2 , the higher positive correlation level yields a higher importance estimate for X_1 . The same pattern is seen for X_2 , where β_1^2 is smaller and β_3^2 is larger than β_2^2 . This means the importance of X_2 is estimated to be higher for $\rho = 0.4$ than for $\rho = 0$, but the increase is smaller than the increase for X_1 (Arnstad 2024). In contrast, the importance of X_3 is then estimated with more weight on β_1^2 and β_2^2 , which are both smaller than β_3^2 , and thus the importance is not notably increased. If one had introduced a larger positive correlation level than $\rho = 0.4$, we would therefore expect the importance of X_3 to even decrease, as was seen in Arnstad (2024). It is hard to say, based on these results, whether the inverse pattern can be seen for negative correlation lev-

els, but it could be noted that the decrease in importance is less for X_3 compared to X_1 and X_2 when ρ changes from 0 to -0.1 .

Generally, it seems that the method is able to capture the expected effects of varying correlation levels, and is in close agreement with the expected theoretical values when the fixed effects are uncorrelated.

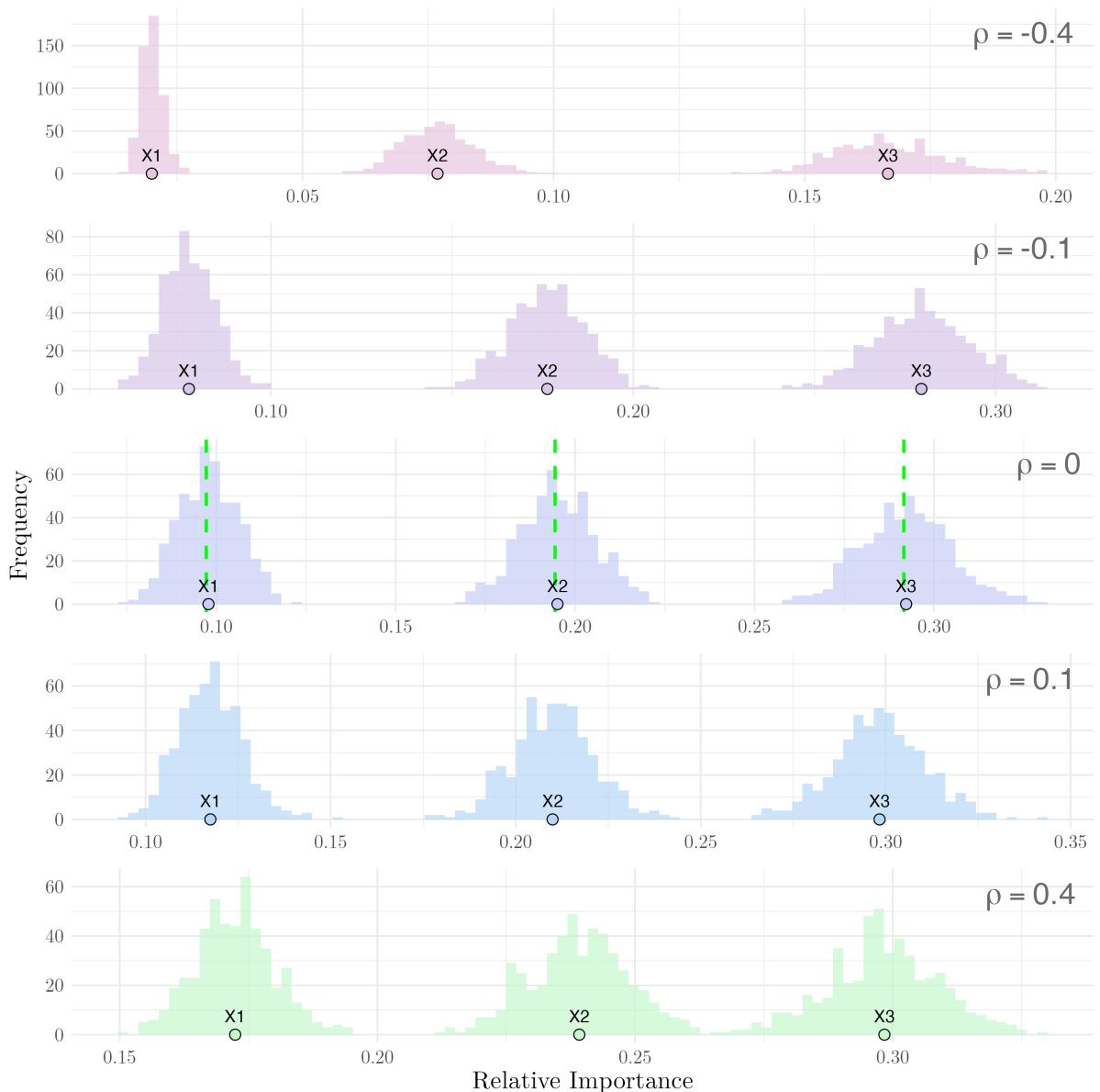


Figure 4.7: Histogram with the distribution of relative importance of the fixed effects present in the Binomial regression for the different correlation levels $\rho = -0.4$ (top), $\rho = -0.1$ (second from the top), $\rho = 0$ (middle), $\rho = 0.1$ (second from bottom) and $\rho = 0.4$ (bottom). The values are calculated by the Bayesian Variable Importance method from the $N_{\text{sim}} = 500$ simulations in the simulation study. The vertical green line for $\rho = 0$ displays the expected relative importance in the case of uncorrelated data. The mean of the relative importance for all simulations is denoted at the bottom of each histogram as a circle.

4.3.1.2 Random effect

The sampled posterior importances for the random effect in the logit model (Figure 4.8) all seem to be roughly normally distributed around the mean. The spread of the random effects seem to be larger than the spread seen in the fixed effects for negative correlations, and more similar for independent and positively correlated covariates. One can see that when the correlation in fixed effects go from negative to positive, the estimated importance of the random effect shrinks. Specifically, when $\rho = -0.4$ the average estimate of relative importance for the random effect is 0.1694 compared to only 0.0667 when $\rho = 0.4$. This naturally occurs as the variance contribution from the random effect should be held constant for the correlation levels, and the variance contribution from the fixed effects rise as the correlation increases. Therefore, the proportion of variance explained, which is our definition of relative variable importance, will decrease for the random effect. For $\rho = 0$ we see that the average relative importance estimate lies very near the expected value of 0.0972 as shown in Table 3.3. The orange dot at the bottom of the histograms in Figure 4.8 displays the estimated relative importance of the fixed effect from the `rptR` package, and we see that the estimates are quite close to the mean of the BVI method. The largest difference from the BVI and the `rptR` package is 0.0267 and are found when $\rho = -0.1$. This difference is 25% of the average estimated relative importance from the BVI method, and is therefore not negligible. However, the methods seem to be in agreement for the overall trends and the methods are closer in accordance with each other for the other correlation levels.

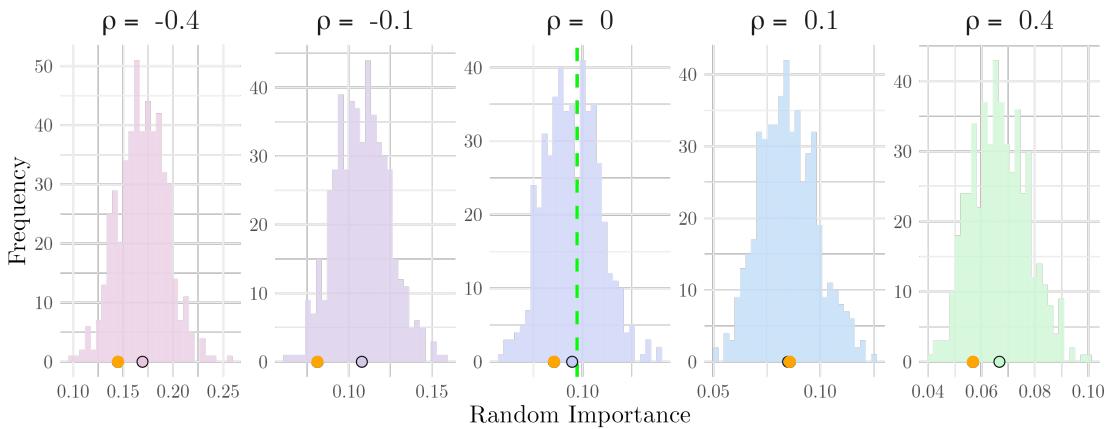


Figure 4.8: Histogram with values from the BVI method for each of the $N_{\text{sim}} = 500$ simulations, estimating relative importance of the random effect α across the different correlation levels ρ . The mean of the estimated relative importance from all simulations is displayed at the bottom as a circle and the orange dot at the bottom displays the estimate from the `rptR` package. The vertical green line for $\rho = 0$ is the expected relative importance as in Table 3.3.

4.3.1.3 R^2 estimates

An important measure in this simulation study is the models sampled posterior distribution of marginal and conditional R^2 (Figure 4.9). The expected values for the marginal and conditional R^2 are shown in Table 3.4, and are displayed as vertical green lines in each plot. It is clear that, regardless of correlation level, the BVI method is able to estimate the marginal and conditional R^2 close to what we expect. The distributions of R^2 values seem to have the shape of a bell curve and are symmetric around the mean value. The spread is naturally larger than for the individual fixed effects and random effects, as the R^2 is constructed from these importances. For the R^2 values, the largest difference between the mean of the simulations from the BVI method and the expected values is 0.002 for the marginal R^2 when $\rho = 0$ and 0.001 for the conditional when $\rho = -0.4$. When comparing to the results from the `rptR` package, it seems that the BVI method consistently estimates larger R^2 values both marginally and conditionally. The largest difference in means between the BVI and the `rptR` estimates for the marginal R^2 is found when $\rho = 0$ and is 0.0448 which makes up 7% of the average estimated value. For $\rho = -0.4$, the mean difference accounts for 14% of the average estimated marginal R^2 value. For the conditional R^2 the largest difference is 0.0617 and found for $\rho = -0.4$, which also makes up 14% of the average estimated value. Generally, the BVI method seems to be in line with our expectation for R^2 values, but deviates somewhat from the `rptR` package estimates.

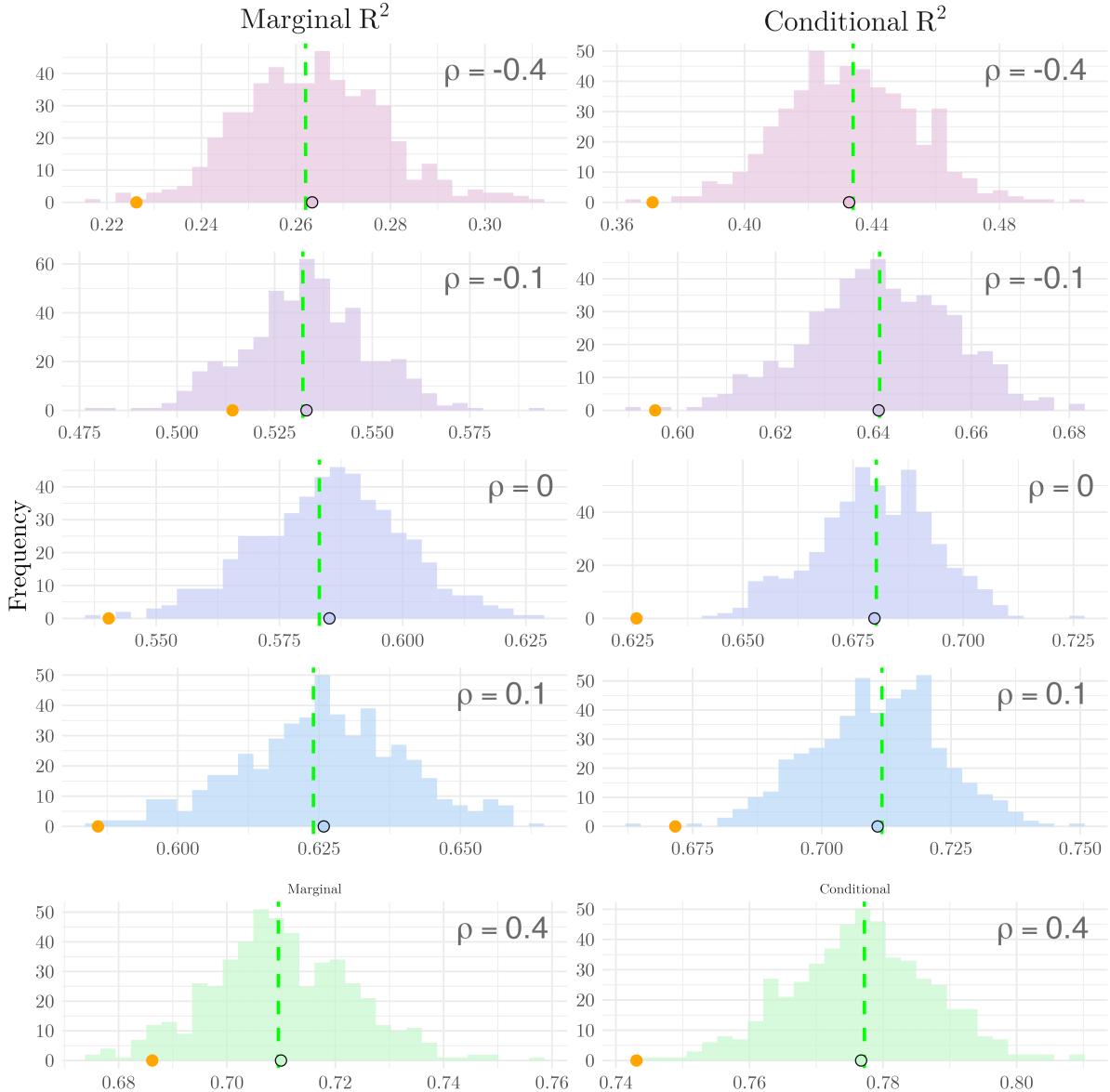


Figure 4.9: Histograms with the estimated marginal R^2 (left) and conditional R^2 (right) from the BVI method for the Binomial regression for the different correlation levels $\rho = -0.4$ (top), $\rho = -0.1$ (second from the top), $\rho = 0$ (middle), $\rho = 0.1$ (second from bottom) and $\rho = 0.4$ (bottom). The values are calculated by the Bayesian Variable Importance method from the $N_{\text{sim}} = 500$ simulations in the simulation study. The expected values are displayed as vertical green lines, and can be found in Table 3.4, while the orange dot denotes the estimate from the `rptR` package. The mean value of the R^2 values for all simulations is marked with a circle at the bottom of each histogram.

4.3.2 Poisson simulation

The second model fit in the simulation study is a Poisson regression with log-link. The Poisson models were fit with the same correlation levels as the Binomial simulation and $N_{\text{sim}} = 500$ models for each level were fitted. For each fitted model, the Bayesian Variable Importance method draws a sample and then calculates the rel-

ative variable importance measures as described in Chapter 3. The results of these calculations construct the estimated posterior distributions of relative importance for all covariates. In the supplementary material (Appendix C, see Table C.2), a summary of the mean and 95% quantile values for different correlation levels is displayed.

4.3.2.1 Fixed effects

As for the Binomial model, we first look at the fixed effects. The estimates of posterior relative importance for the fixed effects (Figure 4.10) are very similar in shape as the Binomial model. Again, the spread is somewhat varying for the different correlation levels. Overall, the overall estimates are smaller for the Poisson model. This is mainly due to the log-link function having a larger associated distributional variance relative to the variance of the covariates, than the logit-link. Therefore, the contribution of fixed effects to the total model variance becomes smaller. Consequently, the Poisson results are less spread and have more narrow quantile values than the Binomial. It seems the estimates form a normal curve about the mean, and for $\rho = 0$ the average estimated importance is close to the expected value. The same influence of varying correlation levels can be seen as in the Binomial model, namely that we obtain larger importance of the fixed effects when the correlation increases. Again the difference is notably large, with the average relative importance of X_1 going from 0.0163 for $\rho = -0.4$ to 0.1579 for $\rho = 0.4$. For X_2 and X_3 the average relative importance for the same correlation levels increases from 0.0627 to 0.2192 and from 0.1357 to 0.2740 respectively.

Further, we see the same pattern of larger increase in importance to X_1 than for X_2 and larger for X_2 than X_3 , as for the Binomial model, when correlation increases from $\rho = 0$ to $\rho = 0.4$. Specifically, X_1 increases with 0.0720, X_2 with 0.0461 and X_3 with 0.0152 when we go from uncorrelated covariates to the highest correlation level. This is in line with our expectations, as the diagonal elements of Λ decrease with increasing correlation levels and the off-diagonal elements increase. The inverse pattern is once again not so clear, as the decrease of X_3 is larger than the decrease of X_1 and X_2 . However, if one looks at the relative decrease in importance, it is clear that the decrease is larger for X_1 and X_2 than for X_3 when going from $\rho = 0$ to $\rho = -0.4$. The inverse pattern may therefore be hard to detect, due to the relatively large difference in magnitude of the coefficients. For the case $\rho = 0$, we see that the mean of our samples and the expected relative importance are very close (Table 3.2), and it seems the model express the expected pattern of relative importance for varying correlation levels.

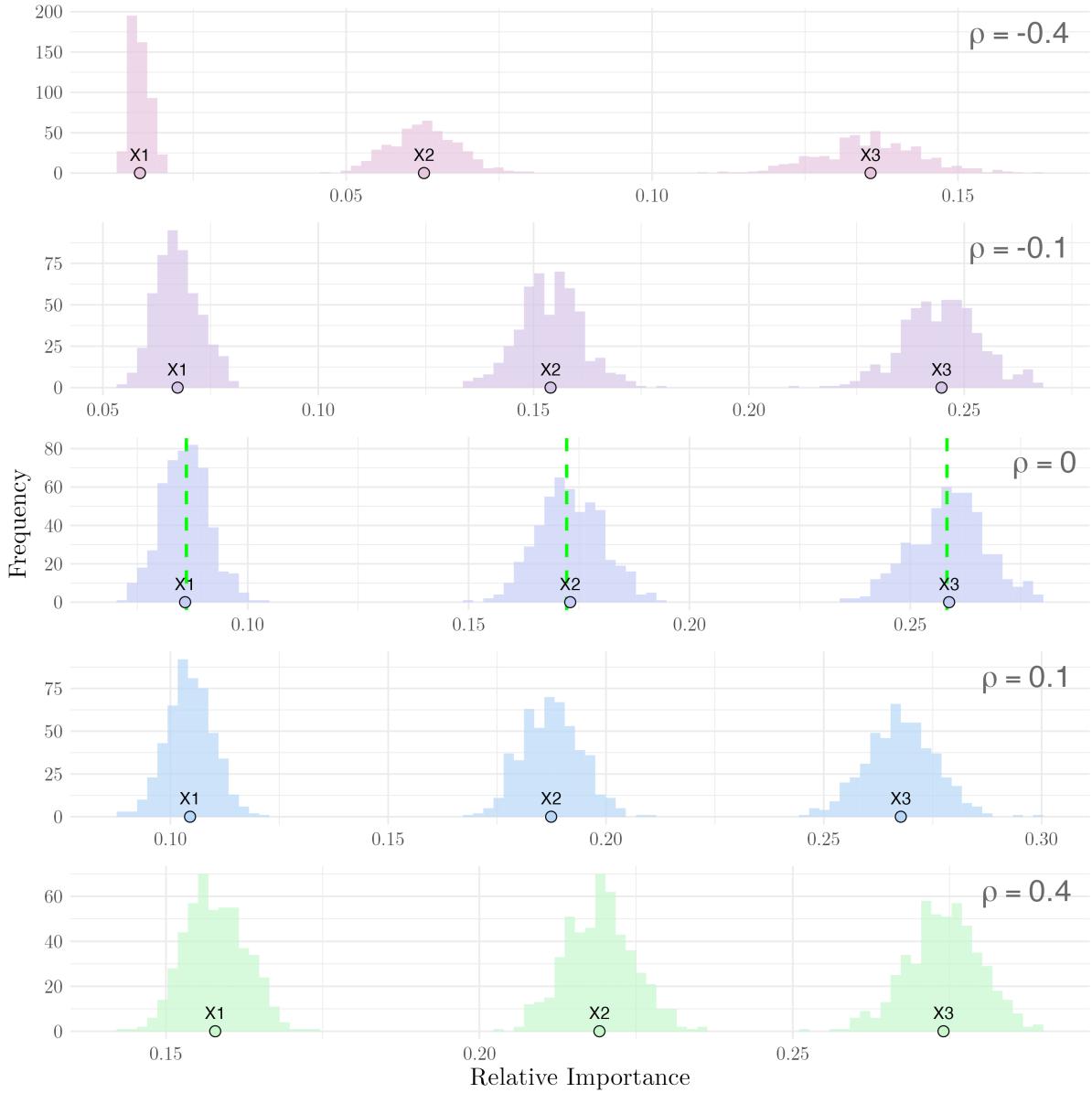


Figure 4.10: Histogram with relative importance of the fixed effects present in the Poisson regression for the different correlation levels $\rho = -0.4$ (top), $\rho = -0.1$ (second from the top), $\rho = 0$ (middle), $\rho = 0.1$ (second from bottom) and $\rho = 0.4$ (bottom). The values are calculated by the Bayesian Variable Importance method from the $N_{\text{sim}} = 500$ simulations in the simulation study. The true regression coefficients are $\beta = (1, \sqrt{2}, \sqrt{3})^T$ and the vertical green line for $\rho = 0$ displays the expected relative importance in the case of uncorrelated data. The mean of the relative importance for all simulations is denoted at the bottom of each histogram as a circle.

4.3.2.2 Random effect

When looking at the sampled posterior distribution of relative importance estimates of the random effect in the Poisson mode (Figure 4.11), we see that they too are in general a bit smaller than the same estimates for the Binomial case. This is again a consequence of the Poisson model having a relatively larger distri-

butional variance, and so we expect these results. Likewise, the distributions can be said to be normal and symmetric around the mean also for the random effect in the Poisson model. We notice a somewhat larger spread in the random effect importances than the fixed effects for the Poisson model. The shrinkage effect of increasing correlation is also here apparent, with the average relative importance of the random effect going from 0.1396 for $\rho = -0.4$ to 0.0627 for $\rho = 0.4$. We again emphasize that this is natural and anticipated, as the random effect variance is constant across correlation levels, while the variance of the fixed effects vary. The expected value when $\rho = 0$ is 0.0861 as shown in Table 3.2, and we see that the average estimate is close to this value. The orange dots, denoting the estimates from the **rptR** package are close to the average estimate from the BVI method, with the largest difference being seen for $\rho = 0.4$ with a difference of 0.0145. This is also the largest relative difference, being 23% of the average estimated relative importance. The method seems to be in agreement with the expected values for the relative importance of the random effect, and the **rptR** estimates are close to the BVI method.

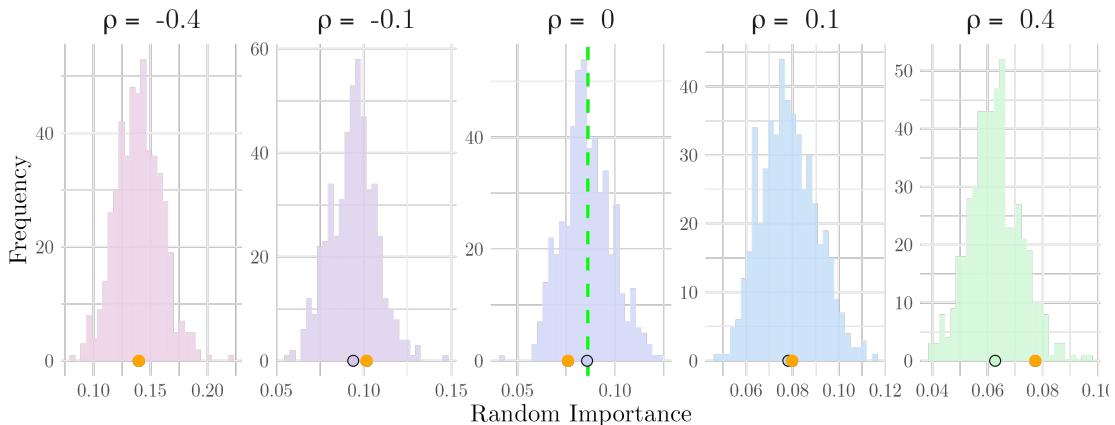


Figure 4.11: Histogram with relative importance estimates for the random effect α for varying values of ρ calculated by the BVI method. The study conducted $N_{\text{sim}} = 500$ simulations and the mean of the relative importance for all simulations is displayed at the bottom of each histogram as a circle. The vertical green line for $\rho = 0$ is the expected relative importance as in Table 3.2.

4.3.2.3 R^2 estimates

Moving on to the estimated posterior R^2 distributions for the Poisson model (Figure 4.12), we see that the expected values from Table 3.4 are in close agreement with the average marginal and conditional R^2 estimated from the BVI method for all correlation levels. The largest difference in expected values and average values from the BVI method is found for $\rho = -0.4$ and is 0.0011 and 0.0004 for the marginal and conditional R^2 respectively. The R^2 distributions seem roughly normal and symmetric around the mean value, with a plausible size of the spread. Contrary to what was the case, it now seems that the BVI method consistently estimates lower values than the **rptR** package. The estimates from the **rptR** package deviate quite a bit from our method, and they seem to deviate more as the correlation increases. For $\rho = -0.4$ the methods agree, but in general the differ-

ences are noticeable. It is hard to say why this occurs, especially since our method coincides very closely with the expected values as well.

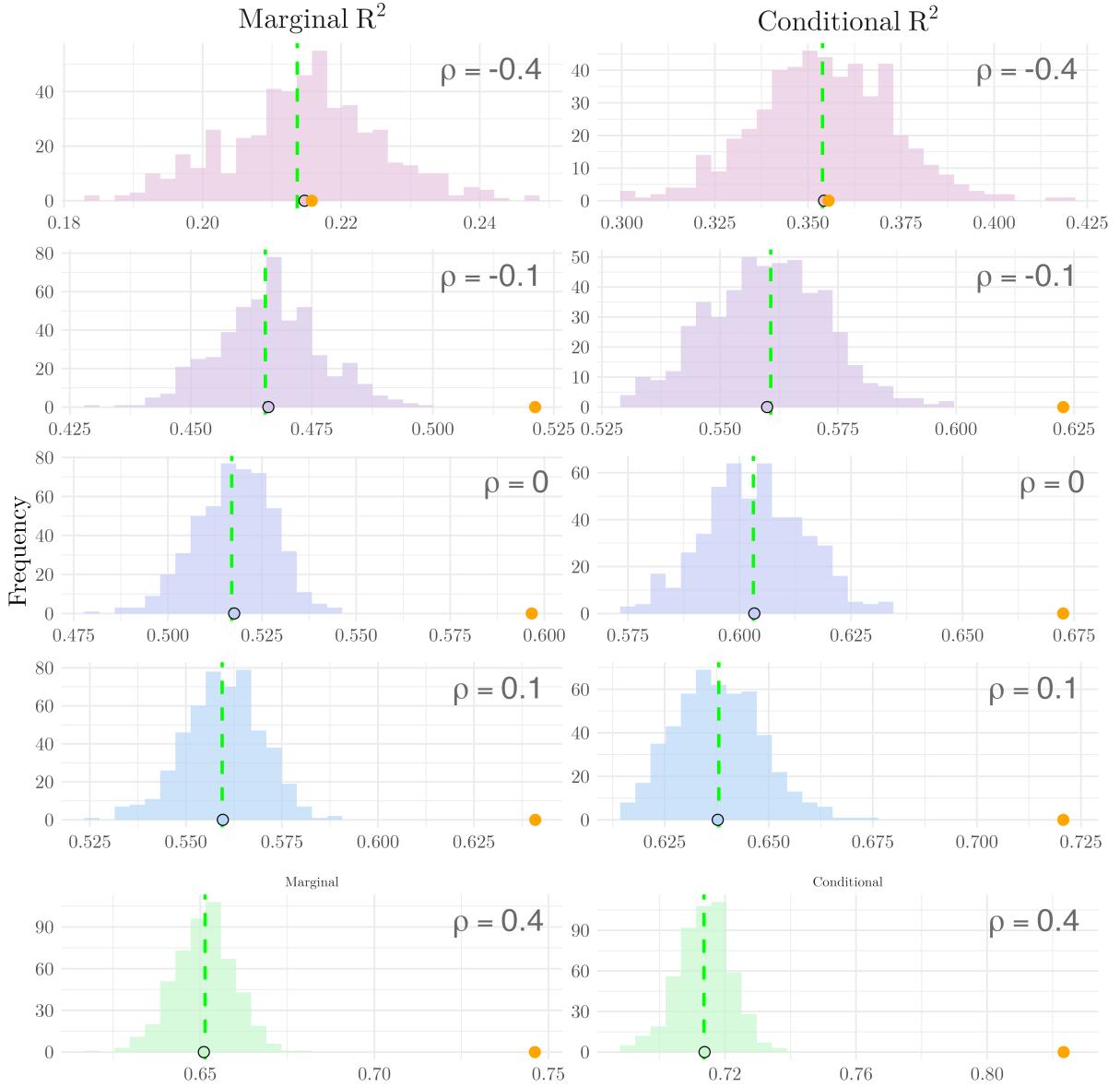


Figure 4.12: Histograms with the estimated marginal R^2 (left) and conditional R^2 (right) from the BVI method for the Binomial regression for the different correlation levels $\rho = -0.4$ (top), $\rho = -0.1$ (second from the top), $\rho = 0$ (middle), $\rho = 0.1$ (second from bottom) and $\rho = 0.4$ (bottom). The values are calculated by the Bayesian Variable Importance method from the $N_{\text{sim}} = 500$ simulations in the simulation study. The expected values are displayed as vertical green lines, and can be found in Table 3.4, while the orange dot denotes the estimate from the `rptR` package. The mean value of the R^2 values for all simulations is marked with a circle at the bottom of each histogram.

4.4 Comparison with rptR package

To further assess our method, a comparison to the vignette for the `rptR` was made. The package described in this vignette estimates the repeatability of phenotypic traits, which for some definitions coincide with heritability and therefore can be seen as a special case of variable importance. Thus, we were able to apply the Bayesian Variable Importance method and compare the results. No expected results were available, and so we can only compare our method to the results made by the authors of the vignette. It should however be noted, that the `rptR` package returns the marginal R^2 as the only measure of importance for the fixed effects, whereas our method directly decomposes this value and assigns a share to each fixed effect. To obtain uncertainty estimates in the likelihood framework, Stoffel, Nakagawa and Schielzeth have built in bootstrap functionality. This is used in our comparison, to evaluate computational complexity and confidence intervals.

The repeatability of the color of male beetles is modelled by a Binomial GLMM with binary outcome and logit-link. We use the same formulation as in the model `rep11` from the vignette (Stoffel et al. 2017), with the parameter `adjusted=FALSE`. We see that the sampled posterior distribution of repeatability (Figure 4.13) from the Bayesian Variable Importance method is centered around a mean of 0.1932, which is very similar to the estimate by Stoffel which is 0.1958. The obtained distribution appears unimodal, with the mode and mean aligning closely. Perhaps a slightly longer tail on the right side can be observed. From 10^3 bootstrap samples, the `rptR` estimates a 95% confidence interval of [0.051, 0.338], which is a bit larger than our estimated 95th percentile of [0.114, 0.280]. In terms of computation time, the Bayesian Variable Importance method used 6 seconds to obtain the model fit and 10^4 samples, whereas the `rptR` package used 66 seconds to obtain the model fit and the same number of bootstrap samples.

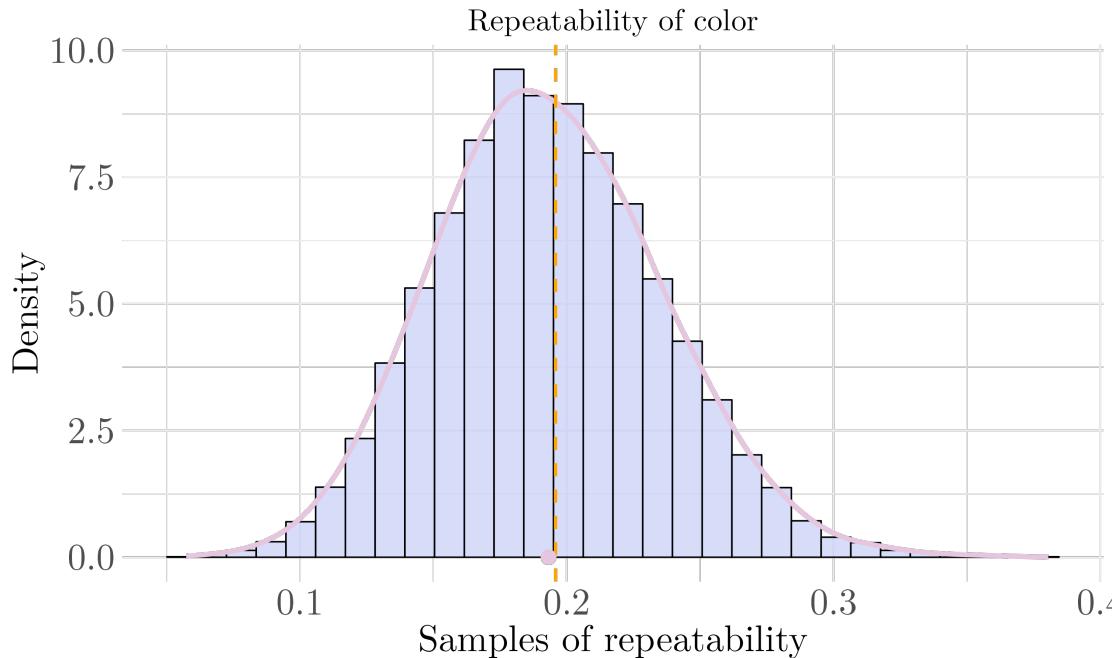


Figure 4.13: Histogram with heritability values for the color of male beetles from the BVI method, with the estimate from the `rptR` package marked as a dashed line with orange color.

To estimate the repeatability of the number of eggs laid by female beetles, we use a Poisson GLMM with log-link. The model used in our method corresponds to `rep9`, but as is described in the vignette after fitting `rep9`, we set the option `expect="latent"` so that the method calculates the distributional variance as in Table 3.1. This corresponds to the recommendations of Nakagawa et al. (2017) as previously mentioned. Likewise, this model is estimated from the `rptR` package with `adjusted=FALSE`. From the plotted samples of posterior repeatability of eggs laid (Figure 4.14), we see a very similar distribution as that of the Binomial color model. The distribution is symmetric and centered around a mean of 0.3585. Further, the estimate from the `rptR` model is 0.3795 with a confidence interval of [0.131, 0.542], compared to our 95th percentile of [0.278, 0.445]. The 10^3 bootstrap samples and model fit for the `rptR` package took 2 minutes and 13 seconds, whereas the BVI method used 8 seconds to obtain the model fit and 10^4 samples.

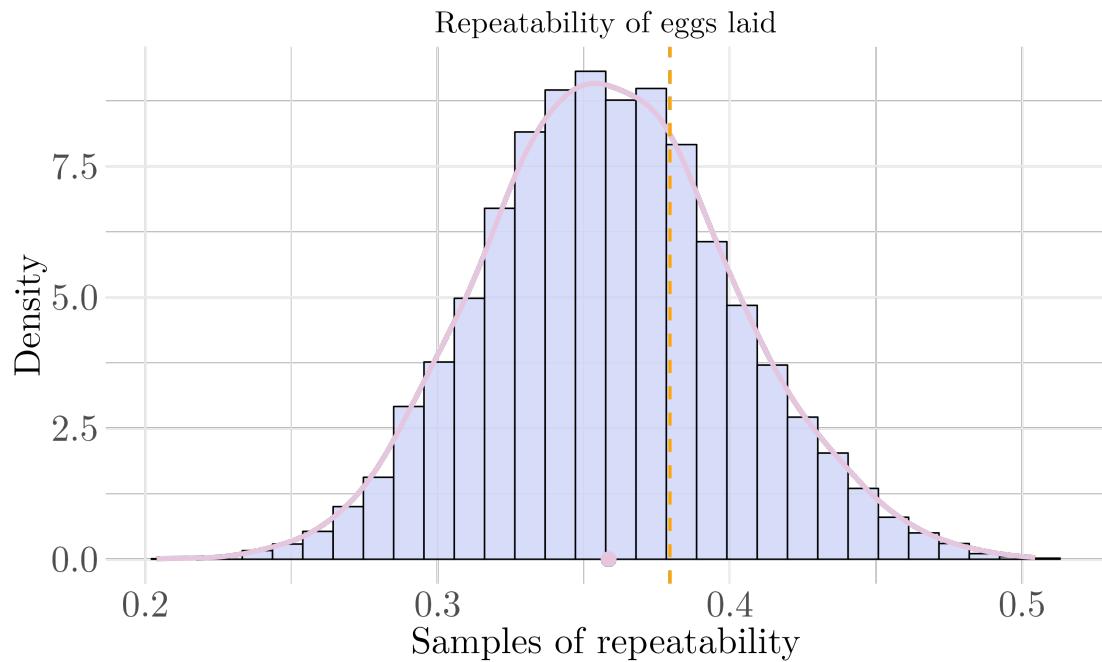


Figure 4.14: Histogram with heritability values for eggs laid by female beetles from BVI method, with the estimate from the `rptR` package marked as a dashed line with orange color.

Importantly, note that the estimates from the BVI method will vary each time a model is fit, as it is stochastic. In this comparison, we only fit a single Bayesian GLMM with the BVI method. Therefore, it could be that another fit from the BVI method might align closer with the results of Stoffel and Nakagawa, but it could also be further off.

4.5 Comparing the BVI method with R^2 -induced Dirichlet decomposition priors and Generalized Decomposition Priors on R^2

To explore other possible relative variable importance tools in the Bayesian framework, we have discussed R2D2 and GDR2 priors in Section 2.5.6 and Section 3.7. We now present the results of applying the R2D2 and GDR2 priors to a linear regression, and see how they can be interpreted as relative importance measures. The results are compared to the BVI method and the LMG method serves as a robust benchmark for all methods. Please note once again that the R2D2 and GDR2 priors are originally developed for prediction models, and that the results presented here are based on our interpretation of how the R2D2 and GDR2 priors can be used for relative importance. The theoretical importances for uncorrelated covariates are found in (3.22) and the theoretical R^2 for all correlation levels are found in Table 3.5.

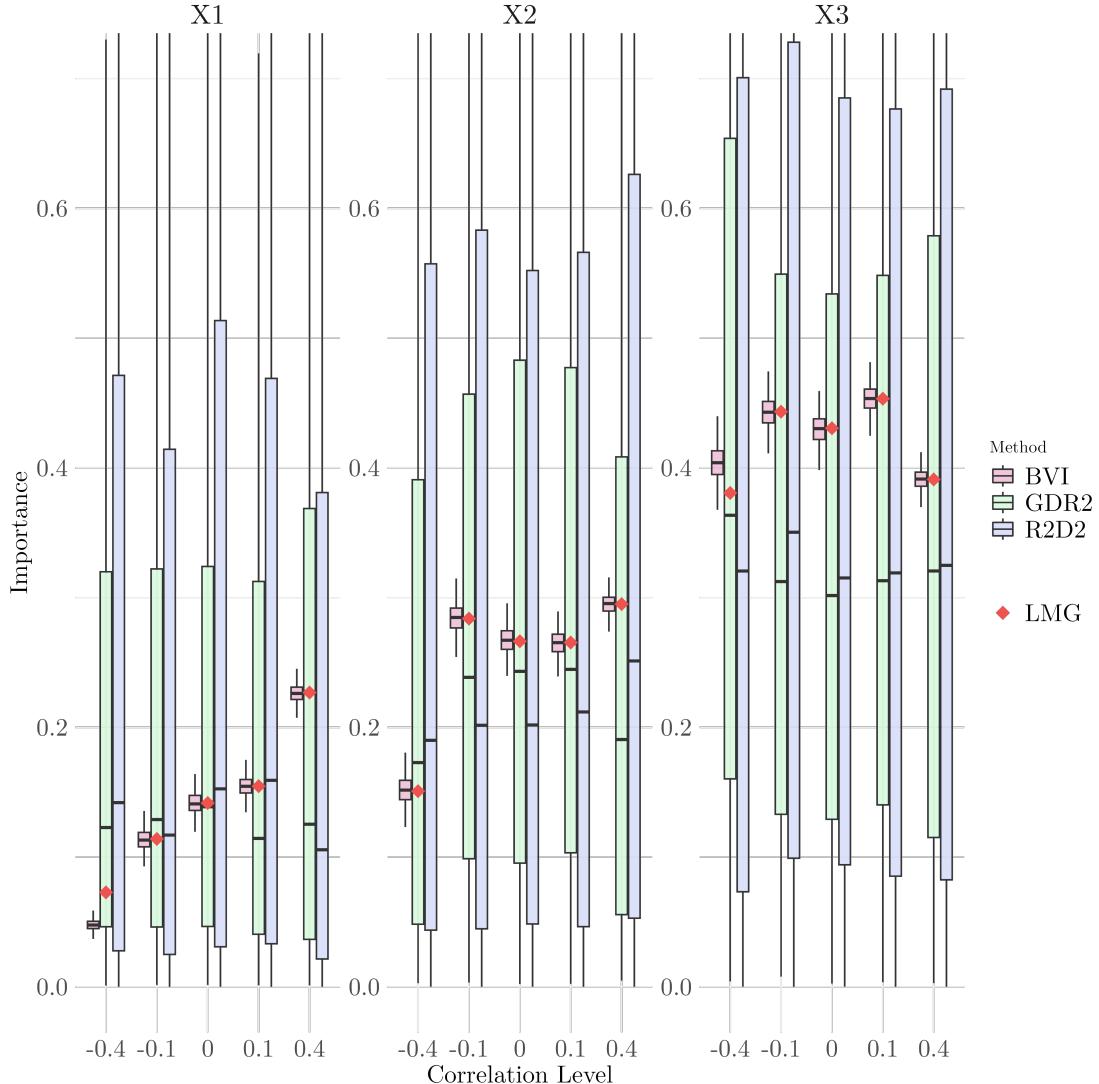


Figure 4.15: Box plots of the relative importance distributions for X_1 (left), X_2 (middle) and X_3 (right) for varying correlation levels by the R2D2, GDR2 and BVI methods. The correlation levels ρ are denoted along the x-axis, and the red diamond represents the relative variable importance calculated from the LMG method.

The first thing one notices from the posterior relative importance distributions (Figure 4.15), is that the spread from the R2D2 method is larger than the spread of the GDR2 method, which again is significantly larger than the spread of the BVI method. In the box plots, the 25th and 75th quantiles define the interval of each box. The mean values of the relative importance distributions for the R2D2 and GDR2 methods do not seem to follow any pattern at all for varying correlation. They are more similar across correlation levels for all covariates, and do not adjust for correlation as the benchmark LMG method. Although it is hard to find patterns and the R2D2 and GDR methods give very flexible results, it becomes clear from they grasp the larger aspects, in that X_3 is estimated to be the covariate with the largest variance contribution, followed by X_2 and lastly X_1 . The BVI method is in close agreement with the LMG method, with deviation for X_1 and X_3 when $\rho = -0.4$. Overall the BVI and LMG methods follow the same

pattern for varying correlation, as we expect and have previously discussed.

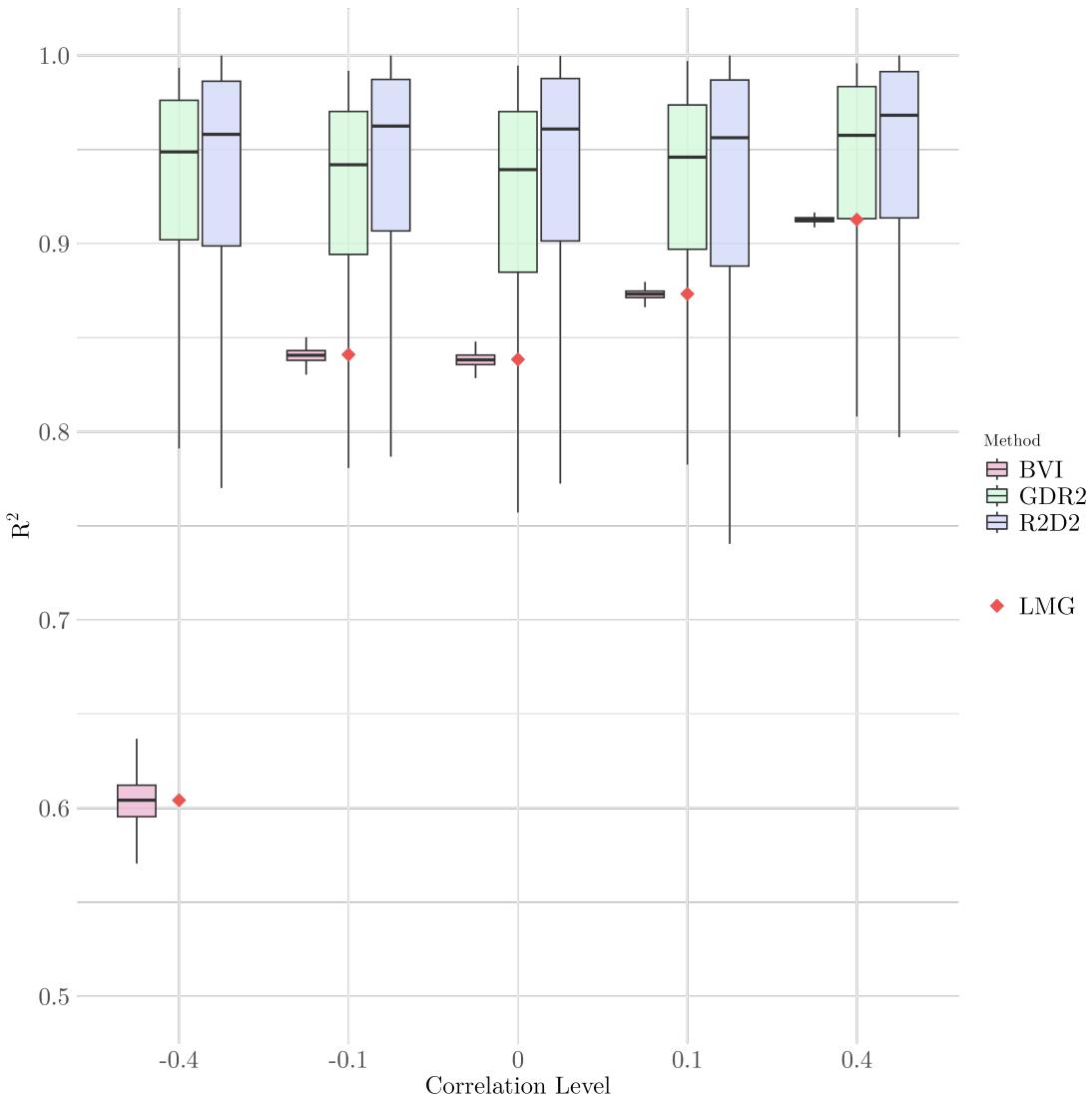


Figure 4.16: Box plots of the estimated R^2 distributions for varying correlation levels ρ for the R2D2, GDR2 and BVI methods. The correlation levels ρ are denoted along the x-axis, and the red diamond represents the relative variable importance calculated from the LMG method.

The estimated posterior marginal R^2 distributions for the R2D2 and GDR2 priors (Figure 4.16) are uniform across the correlation levels, and significantly larger than both the estimates of the BVI and LMG methods. For the R^2 , the spread of the R2D2 and GDR2 methods are quite similar, with both being much larger than that of the BVI method. Overall, the R2D2 and GDR2 estimates a much larger R^2 than the BVI and LMG methods, and the means deviate quite much from the expected values. The BVI method is consistent with the LMG method, and they both follow the expected pattern of the R^2 . For the expected R^2 , the BVI and LMG methods seem to align closely with the expectation, with a small deviation for both methods when $\rho = 0$.

As the R2D2 and GDR2 methods are not specifically relative variable importance

measures, one should also interpret the results with this in mind. The development of the R2D2 and GDR2 priors have been done to produce robust predictions for high dimensional linear regression models, and the results presented here are therefore not an evaluation of the R2D2 and GDR2 priors as relative importance measures. Moreover, the interpretation made to obtain these results was made by the author, and the results should be taken with caution. The main motivation behind this comparison was to explore other possible relative importance measures in the Bayesian framework, as this is a small field with few available methods.

CHAPTER
FIVE

DISCUSSION & FURTHER WORK

The main objectives of this thesis was to develop a general method applicable to a wide range of GLMMs, allowing complex covariance structures in the random effects, and to provide interpretable and trustworthy results. In addition, the method should be easily accessible for researchers across disciplines, and be computationally feasible in most applications. Our attempt to reach these objectives has culminated in the Bayesian Variable Importance method, which puts forth a novel framework for estimating relative variable importance in generalized linear mixed models. The work presented in this thesis is motivated by the increased inference possible in the Bayesian framework and partially builds on the authors previous work in Arnstad (2024).

Summary of contributions

The development of the BVI method has involved utilizing the relative weights method (Johnson 2000) to project the fixed covariates into an orthogonal space. The projection, or approximation, of these covariates are used to fit the model, before a back-transformation is applied to relate the estimated results back to the original covariate space. To obtain inference on the Bayesian GLMMs, we have translated frequentist concepts, such as the R^2 measure, to fit in the Bayesian framework. This translation has been inspired by the work in Gelman et al. (2017), but is also a result of the authors own work. Once the methodology was developed, we revisited the Gaussian simulation study from Arnstad (2024). This simulation study gave us indications that the BVI method was sound for the LMMs, and so it was applied to a real world dataset. The dataset was gathered from house sparrows on Helgelandskysten, Norway, and was used to investigate the heritability properties of the sparrow population. We saw that the BVI method was in close agreement with other heritability studies, which was pleasant. Moving from the LMMs to the GLMMs, we conducted a simulation study for the Binomial and Poisson GLMMs in which the underlying structure was known. The results from the simulation study were promising, and so the method was then compared to a similar relative variable importance measure, `rptR`, in the frequentist framework. This comparison showed that while the BVI method and the `rptR`

method gave quite similar results, the BVI method allows for a more thorough assessment of the covariates and models each of the fixed effects separately. Hopefully, this is advantageous for the researcher using the BVI method, as it allows for a more detailed analysis of the covariates. Lastly, we explored how similar the BVI method was to related Bayesian methods that use shrinkage priors. The results showed that the R2D2 and GDR2 methods are not very precise, and so we argue that the BVI method is a more suitable choice for estimating relative variable importance. This is not surprising, as the shrinkage prior methods were not specifically developed for variable importance. The full methodology has been implemented in an R package, `BayesianVariableImportance`, which is available on the authors GitHub, with a link to the repository provided in Appendix A. In Appendix B a usage example of the package is supplied, which is also available on the authors GitHub along with all code used to obtain the results of this thesis.

Being a general method, our aspirations are that the BVI method will be applied by researchers across disciplines that are interested in the statistical properties of covariates in GLMMs. The BVI method does not aim to give researchers an exact measure of variable importance, but rather provide posterior distributions of relative importance that should be interpreted by the researcher in the field of application. As the distributions will naturally have an uncertainty, it is advantageous if this uncertainty is assessed and understood as a part of the analysis. Hopefully, this can give broader inference on the importance of the covariates, which will in turn lead to more informed conclusions on the effect of covariates on a response. In itself, the BVI poses an analogue to the frequentist relative variable importance measure `rptR` for non-Gaussian responses, but with the added benefit of directly estimating the relative importance distributions of fixed effects. Further, for Gaussian data, it also poses an analogue to more established methods such as the LMG method (Grömping 2007), the extended LMG method (Matre 2022) and the extended relative weights method (Matre 2022) as discussed in Arnstad (2024). Lastly, the BVI method allows one to specify covariance structures in the random effects, which can be beneficial when modeling complex data structures.

Assessment and validation

For relative variable importance measures, some criteria are found in Section 2.2.1 that it is desirable to fulfill. The simulation study on Gaussian LMMs (Section 3.4) shows that the BVI method compares very nicely to the robust LMG method, as well as its extension and the extension of the relative weights method. We argue, as was done in Arnstad (2024), that this is a promising result. Although no theoretical results were derived, one could argue that the simulation study implies that the BVI method gives a proper decomposition, at least in expectation. This is perhaps the most fundamental criteria, as decomposing the R^2 is the main objective of the BVI method. When assessing how the BVI method performs on GLMMs, in which the response variance is not on the same scale as the covariates, this criterion is hard to assess. Instead of aiming to decompose the total model variance, we find

it natural to rather aim for a proper decomposition of the models R^2 on the latent scale. From the definition of R^2 for GLMMs in Nakagawa & Schielzeth (2013), the simulation study shows that the posterior distributions of the marginal and conditional R^2 are generally symmetrically distributed around the expected R^2 value. As the R^2 values in our thesis are constructed from the relative importance assigned to covariates, this indicates that the allocation of relative importance is sensible. Based on these observations, we argue that the BVI method, in posterior expectation, is capable of providing a satisfactory decomposition of the R^2 in GLMMs. Further, the results from the simulation studies for the isolated covariates and the R^2 strengthen our belief that the BVI method correctly captures the expected patterns for different correlation levels. Consequently, we argue that the BVI method allocates the covariates with a plausible relative importance, both for Gaussian and non-Gaussian models. The non-negativity criteria is fulfilled by recalling that the relative importance estimates of fixed effects are squared, and no variance estimate for random effects can be negative. Consequently, the posterior relative importance distributions will not contain negative values. As discussed in Arnstad (2024), the exclusion criteria will not be used in our assessment, as Grömping (2007) argues this is not in general reasonable. Lastly, violating the inclusion criteria is seen as unlikely to occur in practice, although it is mentioned in Matre (2022) that the extensions of the LMG method and the relative weights method can violate this criterion. It has not yet been properly assessed how the inclusion criteria applies to the BVI method. A suggestion that was debated in (Arnstad 2024) is whether one should directly translate the desirable criteria for relative importance measures in the frequentist framework to the Bayesian framework. In the case of the inclusion criteria, we interpret this to mean that if the posterior relative importances of a non-zero regressor contains zero, this is a violation the criteria. The Bayesian framework is designed to provide uncertainty, and therefore subjecting its result against a rigid threshold of containing or not containing zero is not necessarily reasonable. By not considering the inclusion criteria, zero values in the relative importance distributions of a non-zero regressor would require the researcher to carefully assess the covariate. A careful evaluation of the results is in line with what we intend the BVI method to invoke, and therefore the violation of the inclusion criteria might not pose a problem at all. With this in mind, the results both the Gaussian and non-Gaussian simulation studies show that the BVI method produces results that align well with what we expect, and that the results are plausible. Therefore, we believe that for most practical applications, the general idea behind the criteria of variable importance measures are fulfilled by the BVI method.

Another part of validating the BVI method, is to assess how well the methodology performs on real data. To investigate this, we first applied the BVI method to construct an LMM to estimate the heritability of three phenotypic traits of a population of house sparrows. In this analysis, both the grid and CCD integration strategies were applied in the model fitting procedure. The model formulation and pedigree structure was constructed with the help of domain experts, and we compare the results to Silva et al. (2017) and Muff et al. (2019). For all traits and both integration strategies, the posterior distribution of the heritability from the BVI method covers the estimates from the domain experts, and places them

close to the mean. We observed that the average heritability estimate for body mass from the BVI method was narrowly smaller than both estimates from Silva et al. (2017) and Muff et al. (2019). Investigating the wing length, the average heritability estimated by the BVI method was marginally larger than that of Muff et al. (2019) and a bit smaller than the estimate from Silva et al. (2017). The average heritability estimate for tarsus length was very close to the estimates from Silva et al. (2017), and in this case Muff et al. (2019) had no estimate. All posterior distributions of heritability were seemingly normally distributed, except for that of tarsus length, and perhaps body mass, using grid strategy. That the BVI method is in such close agreement with estimates from published papers by domain experts is very promising, and strengthens our belief that the methodology and implementation can be used in practice.

It was difficult to find a real world example to compare the Binomial and Poisson GLMMs to. The solution was to compare the BVI method to the `rptR` method in a case study on repeatability as well as in the non-Gaussian simulation study where the package was applicable. The case study was created by the authors of the `rptR` package, which suggests a repeatability measure for GLMMs. As repeatability is closely linked to heritability, it was possible to use the package in such a way that it could be compared to the BVI method. In the case study, the BVI method and the `rptR` package closely agreed for both the Binomial and Poisson models. The spread of the posterior repeatability from the BVI method was more narrow than the confidence interval from the `rptR`. In terms of computational efficiency, the BVI method was significantly faster than all the models from `rptR` as it does not need to bootstrap to quantify the uncertainty. When using the `rptR` package for comparisons in the simulation study, the results were moderately similar for the relative importance of random effects. However, for the R^2 estimates, the BVI method and the `rptR` package seems to systematically differ. In the Binomial model, the BVI method almost consistently estimated larger R^2 values than that of the `rptR` package. For the Poisson model, the opposite occurred as the BVI method now produced consistently smaller R^2 estimates than the `rptR` package, except when $\rho = -0.4$ in which case they were very similar. Overall, the BVI method was much closer to the expected R^2 values in the non-Gaussian simulation study than the `rptR`. It is hard to say why the BVI differs systematically from both the `rptR` method, but it is pleasing to see that the BVI method aligns well with the expected values.

The field of Bayesian variable importance measures for regression models is not very large, but there has been some research on the topic. Specifically, the use of continuous shrinkage priors for linear models of high dimension has attracted attention (Aguilar & Bürkner 2024). Two priors that can be applied as continuous shrinkage priors and that have favorable properties for variable importance are the R^2 -induced Dirichlet Decomposition (R2D2) priors (Zhang et al. 2020) and its generalization to Generalized Decomposition R^2 (GDR2) priors. Through a simulation study on a linear regression model, the use of R2D2 and GDR2 priors were compared to the BVI method with the LMG method as a benchmark. The results show that the R2D2 and GDR2 priors are not very rigid, by estimating almost uniform distributions of relative posterior variable importance. The almost

uniform distribution may not be reasonable for relative variable importance, but it is sensible for cases where there is little or no prior information available. The shrinkage prior methods generally do not follow the patterns we see from the BVI and the Relaimpo methods, and yield very uncertain estimates. We argue that for the specific task of assigning relative variable importance, the BVI method is more suitable and more reliable than the R2D2 and GDR2 methods. However, the use of these shrinkage priors are primarily not focused on calculating the specific variable importance. Shrinkage prior methods could perhaps be developed further, with an emphasis on variable importance, to yield more suitable estimates for posterior relative variable importance distributions. As the BVI method and the shrinkage prior methods have been developed for different purposes, the R2D2 and GDR2 methods differ from the BVI method in some fundamental ways. Firstly, to our knowledge, the shrinkage prior methods have yet to be applied for GLMMs and so direct comparison for the most complex models is not possible. Further, we sample values of coefficients and random effects *a posteriori* and then estimate the relative importances based on the samples. This means that the estimates from the model are used, which in most cases do not vary greatly for different model fits to the data. On the other hand, the R2D2 and GDR2 priors consider the relative variable importance as a parameter in the model, and therefore places prior values on the relative variable importance directly. When placing the priors directly on the importance, one must keep in mind that the choice of priors are the most criticized topic in Bayesian statistics (Robert 2007). Taking into account that the user is often not informed about the underlying mechanism in the relative variable importance parameter, we assume that they will parameterize the priors in such a way that they reflect the lack of information. Therefore, we see it as sensible that the users initial lack of precision propagates into the final posterior distribution of relative variable importance. This could be a reason why the estimates for the shrinkage priors are more spread out than estimates for the BVI method. Moreover, the priors themselves differ, as the R2D2 and GDR2 priors are continuous shrinkage priors, which are designed to shrink small effects towards zero. We use penalising complexity priors, which puts the emphasis on the complexity of the model. This means the general idea for the priors used is the same, but the implementation and interpretation of their results are different. Lastly, it should be mentioned that the results in this thesis are based on the authors interpretation of how the shrinkage priors can be used for relative variable importance. The author gained this knowledge by reading the papers Zhang et al. (2020) and Aguilar & Bürkner (2024) and by discussing the topic with the authors of Aguilar & Bürkner (2024). Therefore, we believe that one could further optimize the use of shrinkage priors for variable importance by further studying the topic. MORE?

Limitations

It could be questioned if our investigations of the Bayesian Variable Importance method has been sufficient. For example, in the simulation study we do not allow for more extreme correlation levels than -0.4 and 0.4 . This is not a very large value, and therefore some analysis on the method for higher correlation lev-

els could be of interest. The reasoning behind using such moderate correlation levels, is that the model fitting procedure was often compromised for more correlated covariates. As mentioned in the results, we experienced model crashes for a correlation of 0.4, and the frequency of crashes rose when testing with higher correlation. It is therefore not clear how well the model will perform if covariates share much information, but results for an LMM with correlation levels of 0.9 can be found in Arnstad (2024). Further, our initial idea was to also include a Binomial model with the probit-link function in the simulation thesis. This idea was abandoned due to severe model fitting problems, in which INLA would not converge. This was of course unfortunate, but as the result could not be trusted, we chose to omit them from the thesis. We do however believe that if the probit model had been a good fit, the method would be able to calculate the relative importances similarly as for the logit model.

Another topic of discussion, regards choice of priors. It would be natural, given more time, to investigate how different priors would perform and also if one could tune the hyperparameters of the priors applied. As priors are a large subject in themselves, a thorough analysis of prior effects on the BVI method was not performed. We chose to follow the recommendations of Simpson et al. (2017) to use penalising complexity priors, as these had desirable properties and are designed to nicely fit INLA models (Simpson et al. 2017). The parameters of our PC priors follow the default values in the R-INLA package, and we have not investigated how these could be tuned to better fit the data. This could be done to further solidify the results of the BVI method, but would also require more time and resources than what was available in the scope of this thesis.

Many of the foundational calculations made in the BVI method relies on approximations and sampling. The relative weights method can be viewed an approximation of the Lindemann, Merenda and Gold (LMG) method, and the accuracy of INLA is dependent on how well the marginals are approximated. It is to be expected that the errors made in these approximations are propagated to the outputted results of the BVI method. Further, the integration strategy used to compute the marginal posterior distributions of covariates, which again is used to approximate the joint posterior, can affect the sampled values. The samples may either be compromised by poor numerical integration due to a high dimensional hyperparameter vector, or perhaps the assumption of the latent layer being Gaussian is not met, causing the samples to not be representative of the true posterior. An example of this was seen in Section 4.2, where the grid integration strategy gave a posterior distribution for the heritability of tarsus length a trimodal shape. As the grid strategy considers only points that are closer to the log-density than a certain threshold (Gómez-Rubio 2020, See Figure 2.1 for Illustration), we have a slight suspicion that the points considered do not allow for sufficient exploration of the hyperparameter space. This could force the sampling to occur only in a small region of the sampling space, causing the oscillating pattern observed in the posterior heritability of tarsus length. However, this hypothesis is merely speculative, and requires further investigation. Nonetheless, this exemplifies the sensitivity of the BVI method to the chosen strategy, as the interpretations derived from using the grid strategy in this scenario may differ significantly from those obtained us-

ing the CCD strategy. However, we argue that the results obtained from the BVI method are satisfactory. For the simulation studies, the results align well with our expectation in cases where we can give an expectation. Further, the patterns for varying correlation levels seem to be logical, and the results plausible, even though a true value is hard to obtain. Based on this, we believe that the BVI method can pose a useful tool, which is accurate enough for moderate correlation levels and does not require a very informative prior.

Further work

We are not at the time aware of a similar variable importance tool for Bayesian GLMMs as the BVI method. Therefore, there is still much work to be done in this field, and many opportunities for expanding the BVI method. Currently, we have implemented the BVI method to handle Gaussian, Binomial and Poisson distributed responses, but there are a number of other distributions that could be of interest. In Nakagawa et al. (2017), the quasi-Poisson, negative Binomial and Gamma distributions are analysed, so these would be natural extensions. Further, extending the BVI to also handle multiplicative overdispersion would allow the user to specify if the overdispersion should be modelled as additive or multiplicative and would be a valuable addition.

Although not developed for relative variable importance, the shrinkage priors R2D2 and GDR2 could be further explored to see if they can pose as viable variable importance measures. Recently, the author was also introduced to the article Fuglstad et al. (2020). In the article, a framework for selecting priors based on a hierarchical decomposition of total model variance is proposed (Fuglstad et al. 2020). When prior knowledge is not available, the authors use the Dirichlet decomposition as the R2D2 method, however penalising complexity (PC) priors are used if the user has a logical idea of how to decompose the variance. In addition to using PC priors, the method proposed is designed for latent Gaussian models, which are both features of the BVI method when using INLA to fit the Bayesian GLMM. Therefore, this method could be of interest as a possible bridge between the BVI method and the discussed shrinkage prior methods. Due to time constraints, this was not investigated further in this thesis, but could be valuable to explore in the future.

It was also desirable in Arnstad (2024), to go deeper in to the theoretical properties that the BVI method possesses. As the BVI method is first and foremost a tool for researchers, the main focus of this thesis was put on developing a credible variable importance measure and wrap this in an R package so that it could be applied. Due to the complexity of this, the time and resources did not allow for a full theoretical breakdown of the BVI method. Such analysis would be of very high interest, in particular some proofs in expectation for the variable importance estimates would be helpful, to further solidify the credibility of the method.

We did not consider random slopes when developing the BVI method, but this

could also be a possibility for further work. As the random slopes are often associated with a fixed effect, the correlation structure one obtains with random slopes is much more complex than that of random intercepts. This could be a difficult challenge to implement, but as discussed in Section 2.4.2, the proposal by Johnson (2014) could be a good starting point. One should however be careful to make sure that the random slope improves the model significantly, and that this improvement outweighs the potential computational burden including random slopes could have.

Conceptually, variable importance is in itself a debated topic. The first question one can ask is what the definition of relative importance is. In Grömping (2007), relative importance is based on variance decomposition, and we have chosen to follow this notion. However, this definition has the disadvantage that an agreement of allocation of importances for correlated covariates seems impossible (Grömping 2015). This problematic issue is present in our results when the fixed effects were correlated, making evaluation of the method difficult. For our method, the pattern observed was a consequence of the relative weights method, rather than a general method for distributing the shared variance between covariates. The search for a unified variable importance framework has given us methods such as the LMG (Grömping 2007), Proportional marginal variance decomposition (PMVD) (Grömping 2007), the relative weights method (Johnson 2000) and dominance analysis methods (Budescu 1993). Yet, no one has been able to provide a method that is completely accepted by the field of mathematics. For these reasons, variable importance as a subject, and its methods, have received criticism (Grömping 2007). However, we believe that variable importance methods can give researchers valuable information and spark ideas, and that they therefore should have a place in the statistical toolbox. That being the case, we wish to emphasize that all statistical methods are limited by the assumptions they rely on and the data they are applied to. As Chevan & Sutherland (1991) put it; "*Statistical techniques do not build theory - theoreticians do*".

CHAPTER
SIX

CONCLUSIONS

The goal of this thesis was to provide a novel variable importance measure in the Bayesian framework for generalized linear mixed models. To do so, we applied the relative weights method and fit a Bayesian GLMM. Then, we extended a simple definition of the R^2 for GLMMs into the Bayesian framework to obtain our proposed definition. The posterior distribution of the Bayesian GLMM is sampled, before the R^2 is decomposed and distributed to the covariates, to allocate them a relative importance. The methodology is named the Bayesian Variable Importance (BVI) method and wrapped in an R package.

From simulation studies, case studies and real world applications, it has been shown that the BVI method is capable of providing plausible and robust estimates. The uncertainty in estimates is quantified, and the method allows researchers to carry out comprehensive inference. Being a relatively general method, the BVI method can be applied to a wide range of regression models, and has proven to be computationally efficient. It is available to any reader with access to the statistical software R, and has many areas of applications across sciences. There is much potential for further augmentation of the method, both theoretically and practically. It is our aspiration, that the BVI method provides a useful tool, and that it can drive further research in the field of variable importance measures.

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APPENDIX
A

GITHUB REPOSITORY

All code and data used to produce results and all latex files used to produce this document are included in the Github repositories linked below. Please note that the package developed for the masters thesis encapsulates the package developed for the project thesis. Further explanations are given in the readme-files.

GitHub repository link

- Package developed for masters thesis: <https://github.com/AugustArnstad/BayesianVariableImportance>
- Package developed for project thesis: <https://github.com/AugustArnstad/BayesianImportance>
- Full masters thesis: <https://github.com/AugustArnstad/TMA4900-Master-Thesis>
- Full project thesis: <https://github.com/AugustArnstad/TMA4500-Specialization-Pro>

APPENDIX

B

BAYESIAN VARIABLE IMPORTANCE USAGE

```
1 ## GENERAL SETUP
2 First, we set up the necessary libraries and configure the
3   environment for our analysis. This includes loading
4   essential packages and setting options for chunk output
5   and plot dimensions."
6
7 ## GENERAL SETUP
8 First, we set up the necessary libraries and configure the
9   environment for our analysis. This includes loading
10  essential packages and setting options for chunk output
11  and plot dimensions."
12
13 ``{r setup, input=FALSE, echo=FALSE}
14 library(formatR)
15 showsol <- FALSE
16 library(knitr)
17 library(devtools)
18 knitr::opts_chunk$set(tidy.opts = list(width.cutoff = 68),
19                       tidy = TRUE,
20                       warning = FALSE,
21                       error = FALSE,
22                       message = FALSE,
23                       echo = TRUE,
24                       fig.width=7,
25                       fig.height=5,
26                       fig.align="center")
27
28
29 ## INSTALLING THE PACKAGE
30 This section ensures the devtools package is installed, which
31   is required for installing packages from GitHub. We then
32   install the BayesianVariableImportance package directly
33   from GitHub using devtools::install_github(). In the
34   package under the Hello.R file, all functions are defined
```

```

    with corresponding documentation.

27  '''{r}
28 # If not already installed, install the 'devtools' package
29 if(!require(devtools)) install.packages("devtools")
30 devtools::install_github("AugustArnstad/
  BayesianVariableImportance")
31 library(BayesianVariableImportance)
32 '''

33
34 ## SIMULATE DATA
35 In this part, we simulate data to demonstrate the
  functionality of the BayesianVariableImportance package.
  We generate random variables used as fixed effects with
  different correlation structures and random effects. Note
  that the coefficients used here are a bit large for the
  Poisson model, consider lowering them. The data is then
  structured into data frames for further analysis. If you
  have a suitable dataset you can use this instead.

36
37  '''{r}
38 library(remote)
39 library(INLA)
40 library(mnormt)
41 library(ggplot2)
42 library(reshape2)
43 library(RColorBrewer)
44 library(tidyr)
45 library(dplyr)

46
47 set.seed(1)

48
49 simulate_data <- function(n = 10000, n_groups = 100,
  covariance_level=0) {
  # Simulate fixed effects

51
52   sigma <- matrix(c(1, covariance_level, covariance_level,
  covariance_level, 1, covariance_level,
  covariance_level, covariance_level, 1),
  3, 3)

53
54   X <- MASS::mvrnorm(n = n, mu = c(0, 0, 0), Sigma = sigma)
55   X1 <- X[, 1]
56   X2 <- X[, 2]
57   X3 <- X[, 3]

58
59   # Simulate random effects groups
60   Z1 <- sample(1:n_groups, n, replace = TRUE)
61   random_effect_contributions_z1 <- rnorm(n_groups, mean = 0,
  sd = 1)[Z1]
62   # Z2 <- sample(1:(n_groups/10), n, replace = TRUE)

```

```

65 # random_effect_contributions_z2 <- rnorm(n_groups/10, mean
66   = 0, sd = 1)[Z2]
67
68 # Coefficients for fixed effects
69 beta1 <- 1
70 beta2 <- sqrt(2)
71 beta3 <- sqrt(3)
72
73 # Linear predictor
74 eta <- beta1*X1 + beta2*X2 + beta3*X3 + random_effect_
75   contributions_z1 #+ random_effect_contributions_z2
76
77 # Binomial with logit link
78 p_logit <- exp(eta) / (1 + exp(eta))
79 y_logit_bin <- rbinom(n, size = 1, prob = p_logit)
80 data_logit <- data.frame(y_logit_bin, X1, X2, X3, Z1)
81
82 # Binomial with probit link
83 p_probit <- pnorm(eta)
84 y_probit_bin <- rbinom(n, size = 1, prob = p_probit)
85 data_probit <- data.frame(y_probit_bin, X1, X2, X3, Z1)
86
87 # Poisson with log link
88 lambda <- exp(eta)
89 y_pois <- rpois(n, lambda = lambda)
90 data_poisson <- data.frame(y_pois, X1, X2, X3, Z1)
91
92 epsilon = rnorm(n, mean=0, sd=sqrt(1))
93 y_normal <- beta1*X[, 1] + beta2*X[, 2] + beta3*X[, 3] +
94   random_effect_contributions_z1 + epsilon
95 data_normal <- data.frame(y_normal, X1, X2, X3, Z1)
96
97
98 list(binomial_logit = data_logit,
99       binomial_probit = data_probit,
100      poisson = data_poisson,
101      normal = data_normal)
102 }
103
104
105
106 ## USAGE
107 Here we demonstrate the usage of the
108 BayesianVariableImportance package. We fit Bayesian
109 binomial, Poisson and Gaussian models and sample posterior
110 distributions for different simulated datasets using
111 functions from the package. Then, plots are made to
112 display the results.

```

```

108  ' ' '{r}
109  set.seed(1234)
110
111 datasets <- simulate_data()
112
113 glmm_logit <- y_logit_bin ~ X1 + X2 + X3 + f(Z1, model="iid",
114   hyper=list(prec = list(
115     prior = "pc.prec",
116     param = c(1, 0.01),
117     initial = log(1)
118   )))
119
120 glmm_pois <- y_pois ~ X1 + X2 + X3 + f(Z1, model="iid", hyper
121   =list(prec = list(
122     prior = "pc.prec",
123     param = c(1, 0.01),
124     initial = log(1)
125   )))
126
127 lmm <- y_normal ~ X1 + X2 + X3 + f(Z1, model="iid", hyper=
128   list(prec = list(
129     prior = "pc.prec",
130     param = c(1, 0.01),
131     initial = log(1)
132   )))
133
134 model_logit <- BayesianVariableImportance::perform_inla_
135   analysis(datasets$binomial_logit, glmm_logit, family = "
136     binomial", link_func = "logit")
135 model_pois <- BayesianVariableImportance::perform_inla_
136   analysis(datasets$poisson, glmm_pois, family = "poisson",
137     link_func = "log")
136 model_normal <- BayesianVariableImportance::perform_inla_
138   analysis(datasets$normal, lmm, family = "gaussian", link_
139     func = "identity")
140
141 imp_logit <- BayesianVariableImportance::extract_importances(
142   model = model_logit,
143   data =
144     datasets
145     $
146     binomial
147     _logit,
148     random_
149     names=c(
150       "Z1"),
151

```

```
142 fixed_names  
=c("X1",  
  "X2", "  
X3"))  
143  
144  
145 imp_pois <- BayesianVariableImportance::extract_importances(  
  model_pois,  
  datasets$poisson,  
  random_names=c("Z1"),  
  fixed_names=c("X1", "X2", "X3  
    "))  
146  
147  
148  
149  
150 #One can also use the dist_factor argument to specify the  
  distribution factor one wishes to use  
151 imp_pois_2 <- BayesianVariableImportance::extract_importances  
(model_pois,  
  datasets$poisson,  
  random_names=c("Z1"),  
  fixed_names=c("X1", "X2", "X3  
    "),  
  dist_factor = log(1 + 1/exp(  
    summary(model_pois)$fixed  
    [1] + 0.5)))  
152  
153  
154  
155  
156  
157  
158 imp_lmm <- BayesianVariableImportance::extract_importances(  
  model_normal, datasets$normal,  
  random_names=c("Z1"),  
  fixed_names=c("X1", "X2", "X3  
    "))  
159  
160  
161  
162  
163 samples_logit <- BayesianVariableImportance::sample_posterior  
  _count(model_logit, glmm_logit, datasets$binomial_logit, n  
  _samp=5000, additive_param = "Z1")  
164 samples_pois <- BayesianVariableImportance::sample_posterior_  
  count(model_pois, glmm_pois, datasets$poisson, n_samp  
  =5000, additive_param = "Z1")  
165 samples_lmm <- BayesianVariableImportance::sample_posterior_  
  gaussian(model_normal, lmm, datasets$normal, n_samp=5000,  
  additive_param = "Z1")  
166  
167 plots_logit <- BayesianVariableImportance::plot_samples(  
  samples_logit)  
168 plots_pois <- BayesianVariableImportance::plot_samples(  
  samples_pois)  
169 plots_lmm <- BayesianVariableImportance::plot_samples(samples  
  _lmm)  
170  
171
```

```
172
173  ' '
174
175 ## IMPORTANCES
176 The simplest way of obtaining the importances can be done by
177 looking at these objects. Note that these are sampled, so
178 they do not represent the mean of the samples used for
179 plotting further down.
180
181   '''{r}
182 imp_logit
183
184 imp_pois
185
186 imp_lmm
187
188
189
190
191
192
193
194
195
196
197
198
199
200
201
202
203
204
205
206 ## CUSTOM PLOT
207 Cutsomizing plots is often very nice to display information
208 in the way you want it. Therefore, we show how one can
209 customize the plots using ggplot2 based on the samples
210 drawn.
211   '''{r}
212 random <- "Z1"
213
214 random_plot <- ggplot(samples_pois$scaled_random_samples, aes
215   (x = !!sym(random))) +
216     geom_histogram(aes(y = ..density..), fill = "#C6CDF7",
217                   alpha = 0.7, bins = 40, color = "black") +
```

```

213 geom_density(color = "#E6C6DF", adjust = 1.5, linewidth
214   = 1.5) +
215   geom_point(aes(x = mean(samples_pois$scaled_random_samples$  

216     Z1), y = 0), color = "#E6C6DF", size = 4) +
217   labs(#title = paste("Heritability of mass"),
218     x = "Samples of relative importance of random effect",
219     y = "Frequency") +
220   theme_minimal() +
221   theme(legend.position = "none",
222     axis.title.x = element_text(size = 24),
223     axis.title.y = element_text(size = 24),
224     axis.text.x = element_text(size = 24),
225     axis.text.y = element_text(size = 24)
226   )
227
228 random_plot
229
230 str(samples_pois)
231
232 # Assuming 'samples_pois$scaled_importance_samples' is your  

233 # data frame
234 data_long <- samples_pois$scaled_importance_samples %>%
235   pivot_longer(cols = c(X1, X2, X3), names_to = "Variable",
236     values_to = "Value")
237
238 # Updated plot code
239 fixed_plot <- ggplot(data_long, aes(x = Value)) +
240   geom_histogram(aes(y = ..density..), fill = "#C6CDF7",
241     alpha = 0.7, bins = 40, color = "black") +
242   geom_density(color = "#E6C6DF", adjust = 1.5, linewidth
243     = 1.5) +
244   facet_wrap(~ Variable, scales = "free_x") +
245   labs(x = "Samples of relative importance of random effect",
246     y = "Frequency") +
247   theme_minimal() +
248   theme(legend.position = "none",
249     axis.title.x = element_text(size = 24),
250     axis.title.y = element_text(size = 24),
251     axis.text.x = element_text(size = 24),
252     axis.text.y = element_text(size = 24))
253
254 # Print the plot
255 fixed_plot
256
257 r2_data <- data.frame(
258   Marginal_R2 = samples_pois$R2_marginal$`Marginal R2`,
259   Conditional_R2 = samples_pois$R2_conditional$`Conditional
260     R2`
261 )
262
263 # Reshape the data from wide to long format

```

```

257 r2_long <- pivot_longer(r2_data, cols = c(Marginal_R2,
258   Conditional_R2),
259   names_to = "R2_Type", values_to =
260   "Value")
261
262 # Create the plot
263 r2_plot <- ggplot(r2_long, aes(x = Value, fill = R2_Type)) +
264   geom_histogram(aes(y = ..density..), alpha = 0.7, bins =
265     40, color = "black") +
266   geom_density(adjust = 1.5, color = "black", alpha = 0.7) +
267   labs(x = "R2 Values", y = "Density") +
268   scale_fill_manual(values = c("Marginal_R2" = "#C6CDF7",
269     "Conditional_R2" = "#E6C6DF")) +
270   theme_minimal() +
271   theme(legend.title = element_blank(),
272     legend.position = "top",
273     axis.title.x = element_text(size = 14),
274     axis.title.y = element_text(size = 14),
275     axis.text.x = element_text(size = 12),
276     axis.text.y = element_text(size = 12))
277
278 # Print the plot
279 r2_plot
280
281 ```

```

Listing B.1: Usage of the BayesianImpGLMM package with plots and examples.

APPENDIX

C

SUPPLEMENTARY MATERIAL

Supplementary figures for the house sparrow study

We include figures containing the estimated relative importance of all covariates, and R^2 estimates, from both integration strategies used in the house sparrow study (Section 3.5). The figures are presented in the same order as in the main text, starting with the body mass model, followed by the wing length model, and finally the tarsus length model.

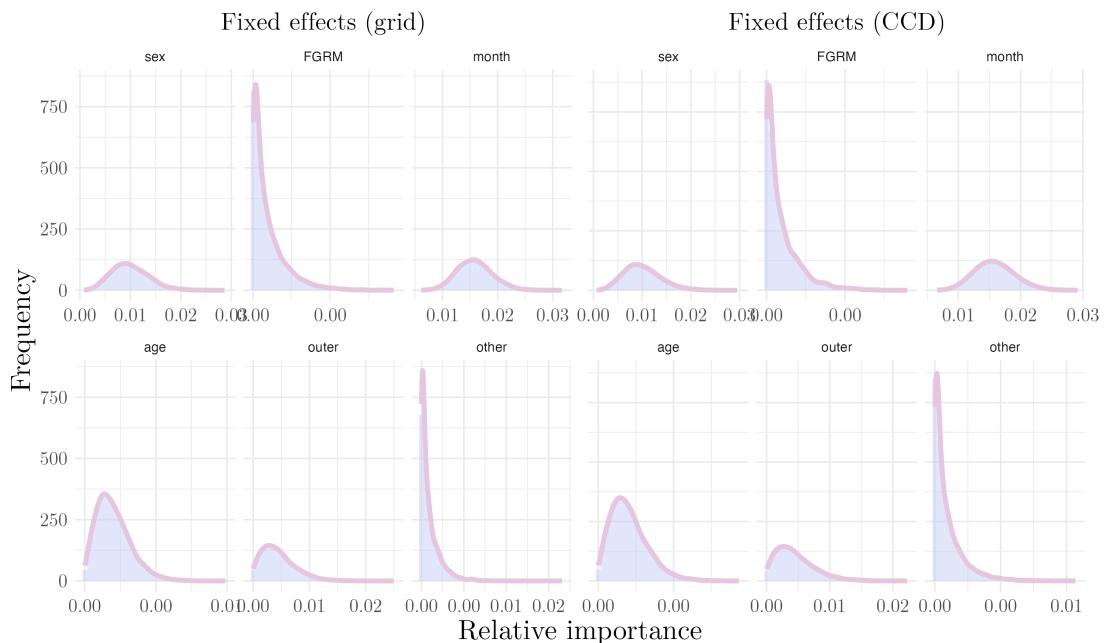


Figure C.1: Posterior relative importance distributions of all fixed effects in heritability of body mass model for house sparrow study. The grid integration is displayed on the left, and CCD on the right.

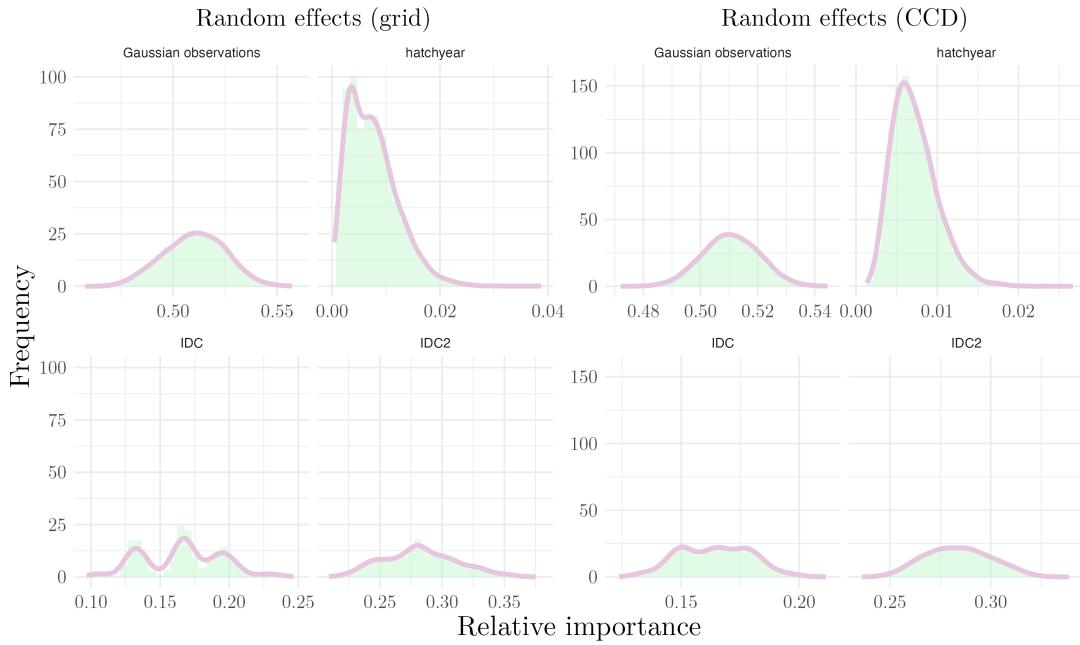


Figure C.2: Posterior relative importance distributions of all random effects in heritability of body mass model for house sparrow study. The grid integration is displayed on the left, and CCD on the right.

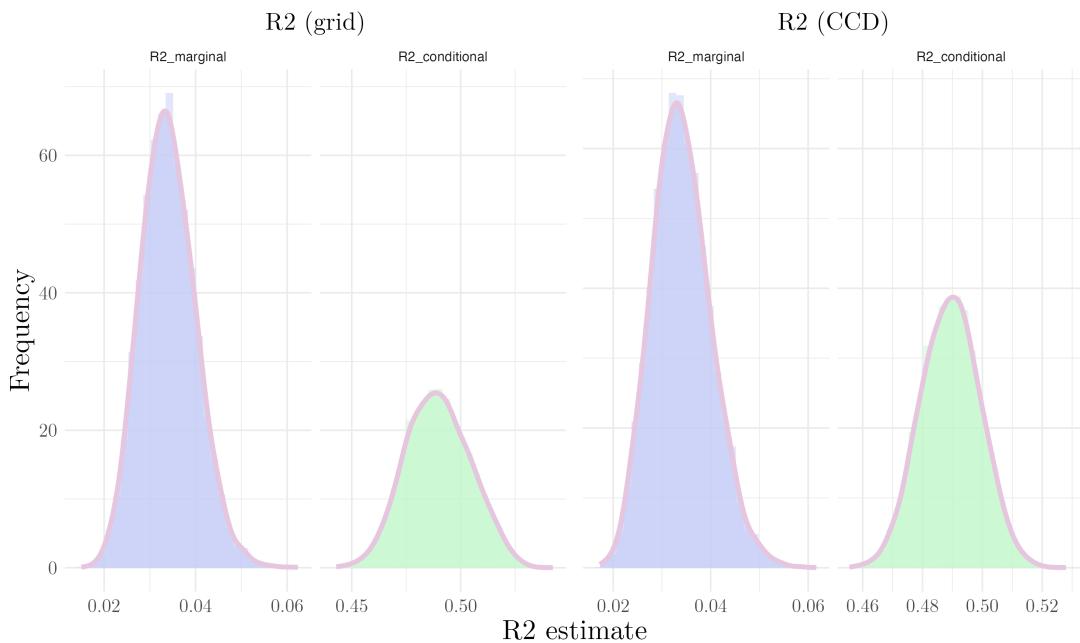


Figure C.3: Posterior distributions of R^2 values in heritability of body mass model for house sparrow study. The grid integration is displayed on the left, and CCD on the right.

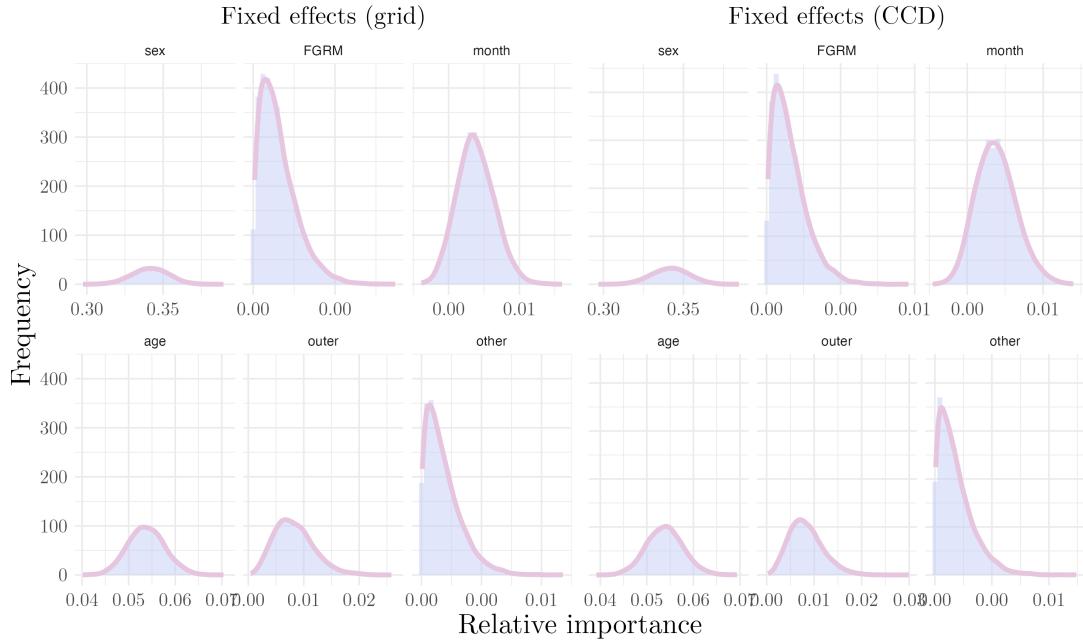


Figure C.4: Posterior relative importance distributions of all fixed effects in wing length model for house sparrow study. The grid integration is displayed on the left, and CCD on the right.

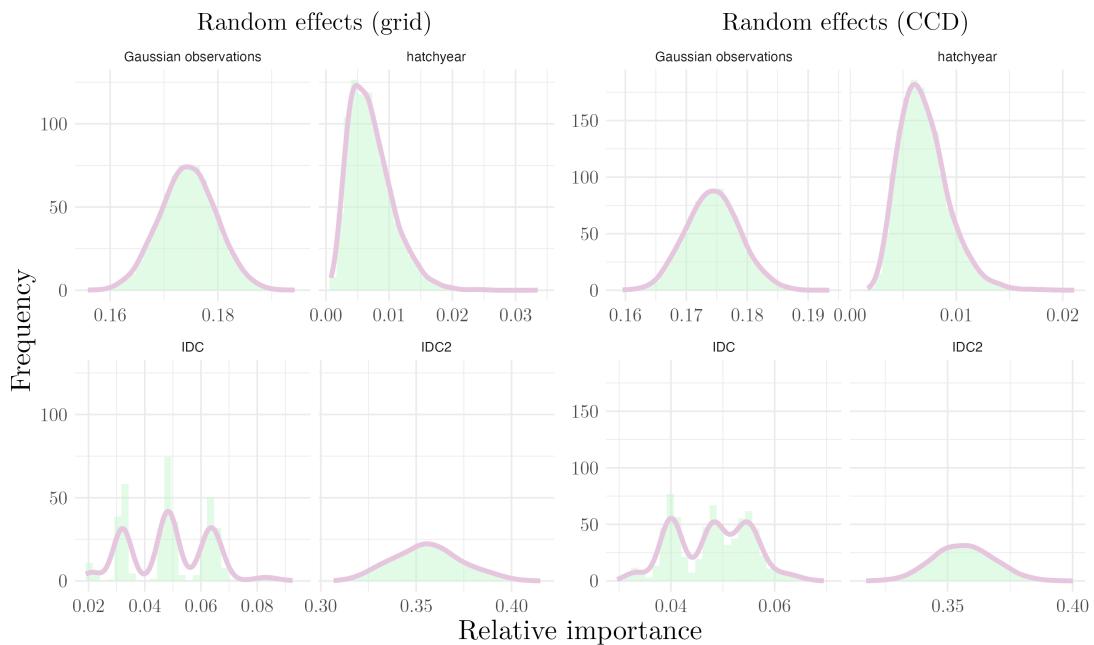


Figure C.5: Posterior relative importance distributions of all random effects in heritability of wing length model for house sparrow study. The grid integration is displayed on the left, and CCD on the right.

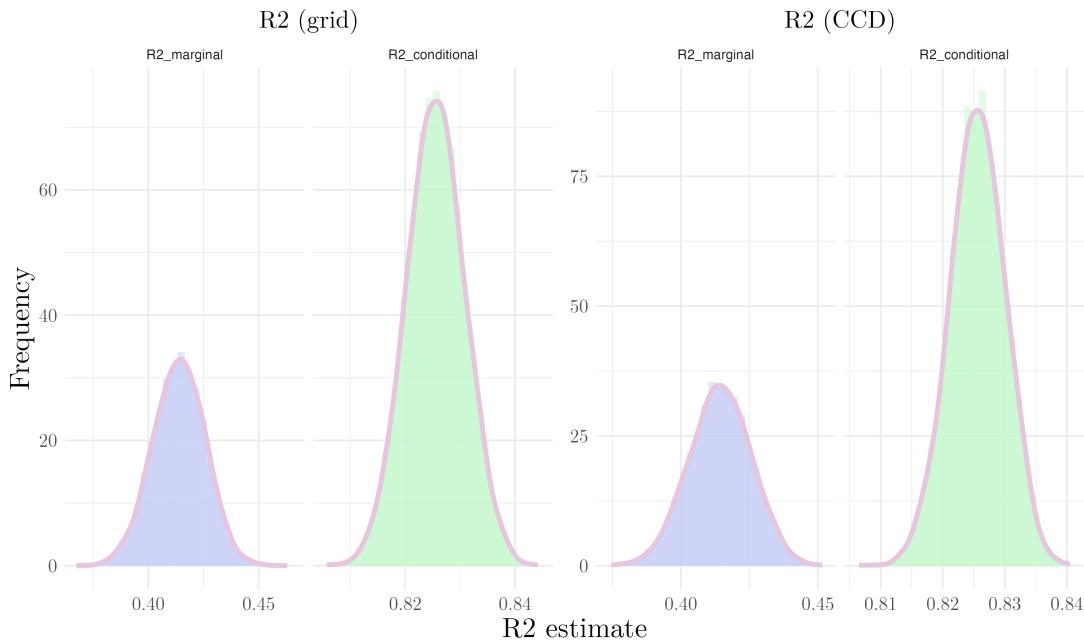


Figure C.6: Posterior distributions of R^2 values in heritability of wing length model for house sparrow study. The grid integration is displayed on the left, and CCD on the right.

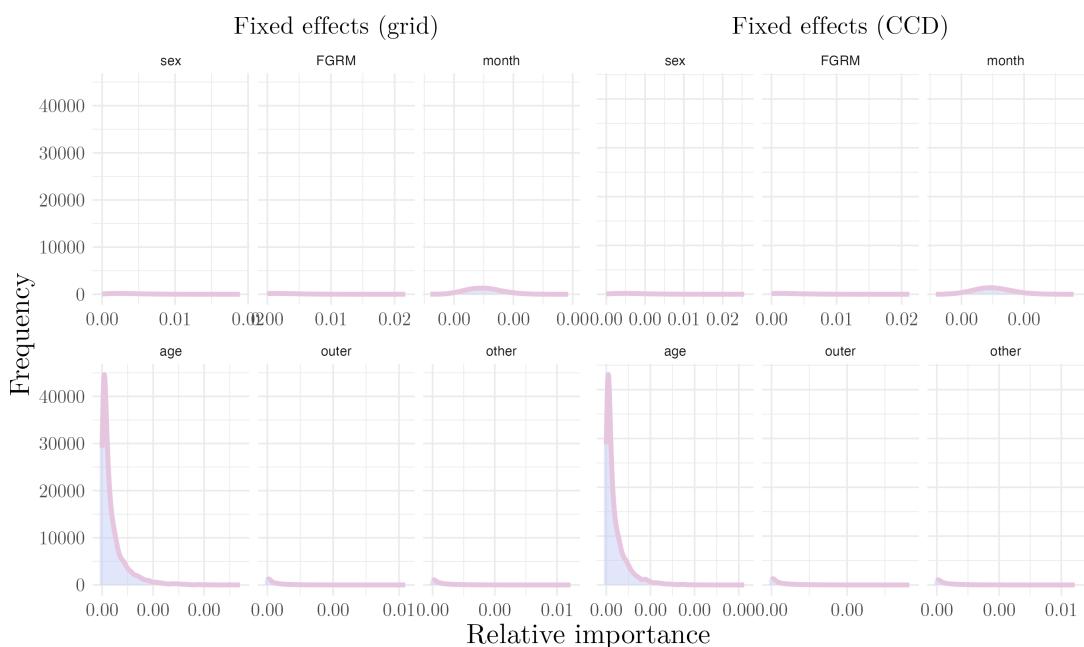


Figure C.7: Posterior relative importance distributions of all fixed effects in heritability of tarsus length model for house sparrow study. The grid integration is displayed on the left, and CCD on the right.

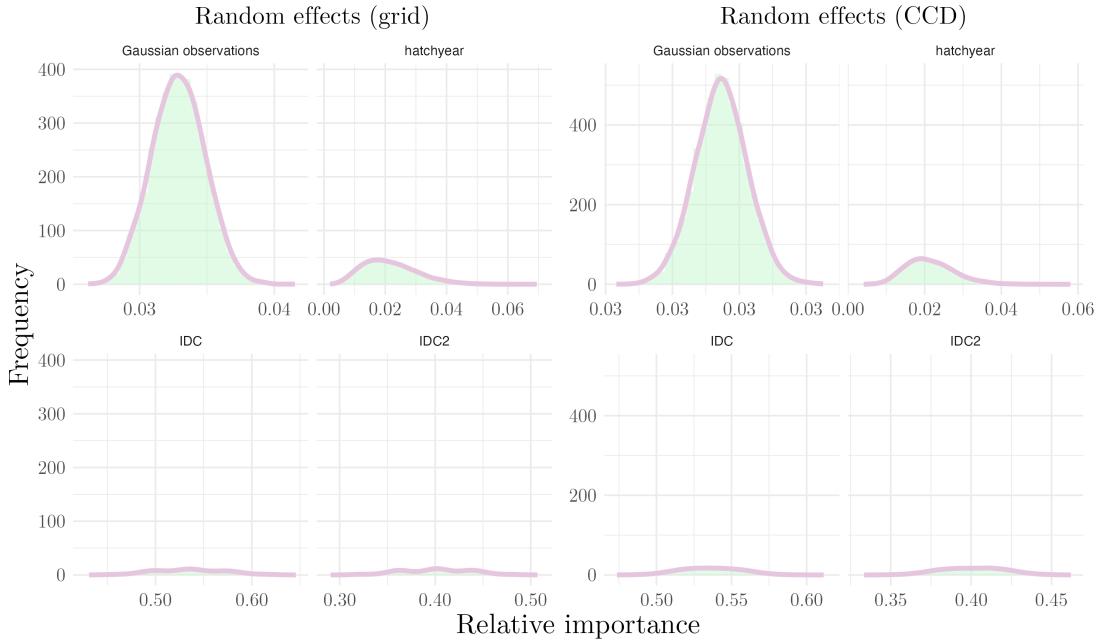


Figure C.8: Posterior relative importance distributions of all random effects in heritability of tarsus length model for house sparrow study. The grid integration is displayed on the left, and CCD on the right.

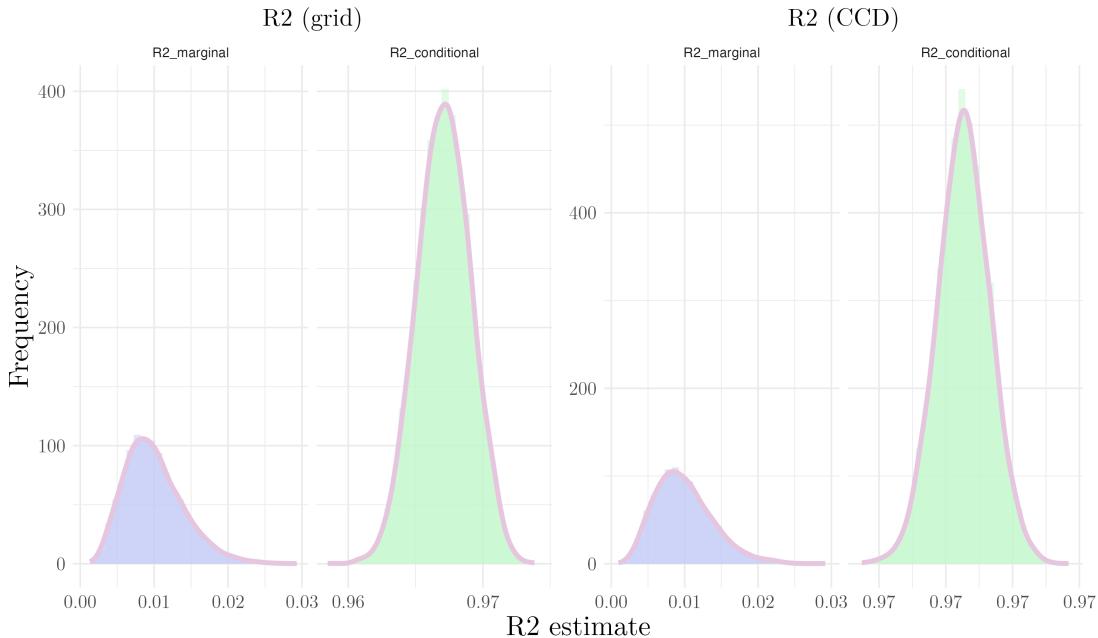


Figure C.9: Posterior distributions of R^2 values in heritability tarsus length model for house sparrow study. The grid integration is displayed on the left, and CCD on the right.

Supplementary tables for the non-Gaussian simulation study

We attach the summarizing tables for the Binomial and Poisson simulation studies here, as they were referred to throughout the thesis. Firstly, Table C.1 contains

summary statistics of the distribution obtained from the Binomial model, while Table C.2 contains the same for the Poisson model.

Measure		$\rho = \mathbf{0}$	$\rho = \mathbf{0.1}$	$\rho = -\mathbf{0.1}$	$\rho = \mathbf{0.4}$	$\rho = -\mathbf{0.4}$
Relative Importance of Random effect	Average	0.0947	0.0850	0.1079	0.0667	0.1694
	2.5%	0.0697	0.0609	0.0771	0.0488	0.1246
	97.5%	0.1223	0.1128	0.1420	0.0889	0.2153
Relative Importance of Fixed effect X1	Average	0.0978	0.1175	0.0774	0.1724	0.0200
	2.5%	0.0817	0.1025	0.0637	0.1580	0.0162
	97.5%	0.1134	0.1350	0.0917	0.1885	0.0245
Relative Importance of Fixed effect X2	Average	0.1951	0.2100	0.1763	0.2392	0.0769
	2.5%	0.1735	0.1903	0.1566	0.2206	0.0643
	97.5%	0.2155	0.2324	0.1956	0.2584	0.0911
Relative Importance of Fixed effect X3	Average	0.2922	0.2984	0.2795	0.2984	0.1666
	2.5%	0.2675	0.2742	0.2539	0.2770	0.1466
	97.5%	0.3187	0.3224	0.3047	0.3201	0.1909
R_m^2	Average	0.5851	0.6258	0.5331	0.7100	0.2634
	2.5%	0.5549	0.5965	0.5023	0.6847	0.2357
	97.5%	0.6146	0.6552	0.5632	0.7360	0.2960
R_c^2	Average	0.6799	0.7108	0.6411	0.7767	0.4328
	2.5%	0.6529	0.6865	0.6118	0.7555	0.3934
	97.5%	0.7046	0.7352	0.6677	0.7967	0.4729

Table C.1: Summary of simulation study results for the quantiles of relative importance estimates of the Logit model across different correlation levels. For $\rho = 0$ the expected values are given in Table 3.2.

APPENDIX
D

MISCELLANEOUS PROOFS

We present a joint proof of the expectation and variance of a random variable belonging to the univariate exponential family. For a random variable Y with a normalized probability density function $f(y|\theta, \phi)$ on the form

$$f(y|\theta, \phi) = \exp\left(\frac{y\theta - b(\theta)}{a(\phi)} + c(y, \phi)\right), \quad (\text{D.1})$$

where θ is the natural parameter and ϕ is the dispersion parameter, the expectation and variance of Y can be expressed as

$$\begin{aligned} \mathbb{E}(Y|\theta) &= b'(\theta) \\ \text{Var}(Y|\theta) &= b''(\theta) \end{aligned} \quad (\text{D.2})$$

This can be proven by considering the following:

$$\frac{df(y)}{d\theta} = \frac{1}{a(\phi)} f(y|\theta, \phi)(y - b'(\theta)), \quad (\text{D.3})$$

and

$$\frac{d^2f(y)}{d\theta^2} = \frac{1}{a(\phi)} f(y|\theta, \phi) \left(\frac{1}{a(\phi)} (y - b'(\theta))^2 - b''(\theta) \right). \quad (\text{D.4})$$

Now, assuming mild regularity to interchange derivation and integration and noting that $\int_{\mathbb{R}} f(y|\theta) dy = 1$, we have

$$\frac{d}{d\theta} \int_{\mathbb{R}} f(y) dy = \int_{\mathbb{R}} \frac{df}{d\theta} dy = 0, \quad (\text{D.5})$$

and

$$\frac{d^2}{d\theta^2} \int_{\mathbb{R}} f(y) dy = \int_{\mathbb{R}} \frac{d^2 f}{d\theta^2} dy = 0. \quad (\text{D.6})$$

Equations (D.5) and (D.6) can be used to derive the relation

$$\begin{aligned} 0 &= \int_{\mathbb{R}} \frac{df(y)}{d\theta} dy = \frac{1}{a(\phi)} \int_{\mathbb{R}} f(y)(y - b'(\theta)) dy \\ &= \frac{1}{a(\phi)} \left(\mathbb{E}(Y|\theta) - b'(\theta) \int_{\mathbb{R}} f(y) dy \right) \\ &= \frac{1}{a(\phi)} (\mathbb{E}(Y|\theta) - b'(\theta)) \\ \implies \mathbb{E}(Y|\theta) &= b'(\theta), \end{aligned} \quad (\text{D.7})$$

and

$$\begin{aligned}
0 &= \int_{\mathbb{R}} \frac{d^2 f(y)}{d\theta^2} dy = \frac{1}{a(\phi)} \int_{\mathbb{R}} f(y) \left(\frac{1}{a(\phi)} (y - b'(\theta))^2 - b''(\theta) \right) dy \\
&= \frac{1}{a(\phi)} \int_{\mathbb{R}} f(y) \left(\frac{1}{a(\phi)} (y - \mathbb{E}(Y))^2 - b''(\theta) \right) dy \\
&= \frac{1}{a(\phi)} \left(\mathbb{E}[(y - \mathbb{E}(Y))^2] - b''(\theta) \int_{\mathbb{R}} f(y) dy \right) \\
&= \frac{1}{a(\phi)} \text{Var}(Y) - b''(\theta) \\
\implies \text{Var}(Y|\theta) &= a(\phi)b''(\theta) \quad \square
\end{aligned} \tag{D.8}$$