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**ManyBabies 5: A large-scale investigation of the proposed shift from familiarity preference to novelty preference in infant looking time**

Pre-data collection manuscript for peer-review

The ManyBabies 5 Team

## Abstract

Much of our basic understanding of cognitive and social processes in infancy relies on measures of looking time, and specifically on infants' visual preference for a novel or familiar stimulus. However, despite being the foundation of many behavioral tasks in infant research, the determinants of infants' visual preferences are poorly understood, and differences in the expression of preferences can be difficult to interpret. In this large-scale study, we test predictions from the Hunter and Ames model of infants' visual preferences.<sup>1</sup> We investigate the effects of three factors predicted by this model to determine infants' preference for novel versus familiar stimuli: age, stimulus familiarity, and stimulus complexity. Drawing from a large and diverse sample of infant participants ( $N = XX$ ), this study will provide crucial empirical evidence for a robust and generalizable model of infant visual preferences, leading to a more solid theoretical foundation for understanding the mechanisms that underlie infants' responses in common behavioral paradigms. Moreover, our findings will guide future studies that rely on infants' visual preferences to measure cognitive and social processes.

**Keywords:** looking time methods; familiarity preference; novelty preference

## Introduction

Scholars and parents alike have long been fascinated by questions concerning the infant mind. What are infants thinking, seeing, hearing, and feeling, and from where and when do these abilities arise? Because it is not possible to query infants directly, scientists must rely on indirect techniques to study early development. However, many of the measures and procedures used with older children and adults, such as button presses or responding to questions, are difficult or impossible to use with infants. This has led researchers to adopt an indirect behavioral measure – infants' looking toward visual stimuli – as one of the primary measures for uncovering the development of psychological processes in infancy.<sup>2</sup>

Much of the prior research using infants' looking time is based on the seminal finding that infants, following repeated exposure to a particular stimulus, typically demonstrate a preference to look at a relatively new stimulus.<sup>3</sup> This novelty preference has been well-documented for both visual and auditory stimuli.<sup>4-7</sup> The approach in these looking time studies is often to present one or more stimuli until infants' interest in the familiar stimuli is reduced, thereby inducing preferences for novelty.<sup>8</sup> However, infants' responses, as well as scientists' interpretations of these responses, vary substantially.<sup>2</sup> Although many studies have found that infants show a robust novelty preference<sup>9-11</sup>, there are also conditions under which infants prefer familiar stimuli over novel stimuli<sup>4,12-15</sup>. Even within specific research areas, there are usually examples of both directions of preference, and there are no clear reasons as to why<sup>6,16</sup>. Furthermore, in some cases, the same type of stimulus that draws infants' attention in one paradigm can fail to draw attention in another.<sup>17,18</sup>

Despite this variability, scientists have generally focused on novelty preferences as a basis for making inferences about infants' cognitive processing. This idea is consistent with

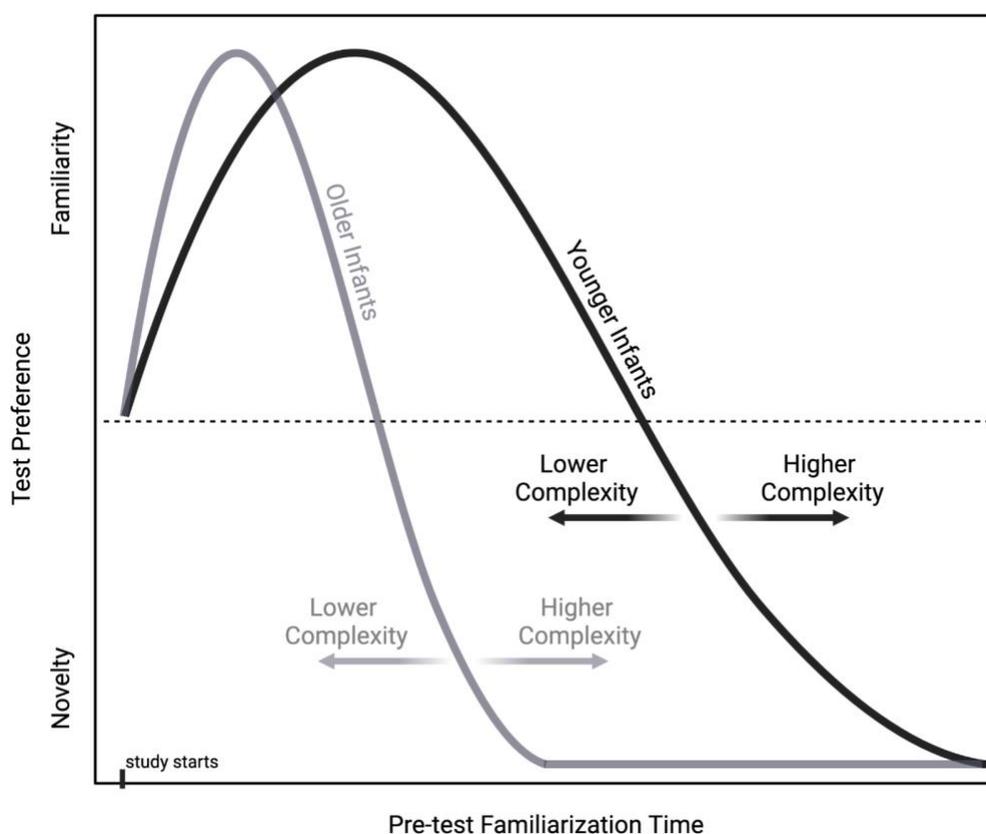
arguments from multiple disciplines suggesting that stimulus novelty may serve as a key signal for exploration and learning.<sup>19</sup> That is, the novelty of a stimulus may engage intrinsic motivation mechanisms that, in turn, drive organisms to invest cognitive resources in learning under conditions of uncertainty.<sup>20</sup> Preferences for novelty have been identified in numerous diverse species<sup>21</sup> and have been linked with both basic survival processes like locomotor adaptation<sup>22</sup> and high-level constructs like curiosity<sup>23–26</sup>. Understanding the determinants of infants' familiarity and novelty preferences, therefore, is important not only for methodological reasons but also for clarifying mechanisms of cognitive development.

The most influential account of looking preferences in the human infant literature is the conceptual model put forth by Hunter and Ames (henceforth, the Hunter and Ames model)<sup>1</sup>, which was based on several studies.<sup>4,27,28</sup> Hunter and Ames aimed to account for the circumstances under which infants show familiarity vs. novelty preferences while processing information presented in stimuli. The model assumes that when one stimulus is repeatedly presented, infants gradually develop a preference for this familiar stimulus compared to a second, novel stimulus. After continued exposure to the familiar stimulus, infants' preferences eventually switch, and they begin to show a preference for the novel stimulus (see Figure 1). The key premise of the model is that the degree to which infants prefer familiar vs. novel stimuli depends on three main factors and their interactions (see Figure 1):

- (1) **Familiarization time.** With increasing familiarization to a stimulus, infants transition from preferring a familiar stimulus to preferring a novel stimulus.
- (2) **Infant age.** The transition from preferring a familiar stimulus to preferring a novel stimulus is faster for older infants than younger infants.

**(3) Task difficulty.** As task difficulty increases (operationalized in the current study as **stimulus complexity**), infants make a slower transition from preferring a familiar stimulus to preferring a novel stimulus.

At the time of writing, this influential model has been used to explain findings within a broad range of papers examining visual attention, speech segmentation, object categorization, face recognition, quantification, memory, knowledge of the physical world, and social cognition, among others.<sup>5,12,29–31</sup>



*Figure 1.* This figure (adapted from Bergmann and Cristia<sup>6</sup>) depicts Hunter & Ames' model<sup>1</sup> of infant looking to familiar (plotted up) and novel (plotted down) stimuli at different levels of pre-test familiarization time (shown on the x-axis). The dashed line indicates equal preference for familiar and novel stimuli. The grey line represents the model's prediction for older infants, and the black line for younger infants. The "lower complexity" and "higher complexity" arrows indicate how the proposed relationship between age and familiarization time might shift with variation in stimulus complexity (i.e., the current operationalization of task difficulty).

However, this model has not undergone a rigorous and comprehensive test, and the predictions from the model have not been supported unequivocally by the existing literature. Whereas several studies have demonstrated the proposed shift from familiarity to novelty preferences<sup>32–35</sup>, other studies have found limited evidence for this shift.<sup>6,36–38</sup> More generally, there is minimal consensus on the factors that drive a familiarity preference in some conditions and a novelty preference in others. This gap in understanding about a core component of visual preferences in infancy has raised numerous interpretive challenges over the decades, including when to predict a preference for novelty versus familiarity and how to interpret unexpected, unstable, or null preferences across studies.<sup>33,34,39</sup>

A large-scale investigation into the fundamental tenets of the Hunter and Ames model is an important step toward quantifying relationships between infant age, familiarization time, task difficulty, and preference for familiar vs. novel stimuli. In the present study, we bring together researchers from around the world to create and implement a best-test of key predictions of the Hunter and Ames model. This investigation will enable a broad community of researchers to improve their interpretations of infants' visual preferences and thereby enable new and more robust insights into the infant mind.

There are two main challenges in evaluating the Hunter and Ames model. First, most behavioral studies with infants have lacked sufficient statistical power due to the inherent limitations within any given lab in testing large numbers of participants.<sup>40,41</sup> Recruiting participants and obtaining a high data 'yield' are notoriously challenging in infant research. Moving forward, infant research is in need of solutions to achieve higher statistical power in order to improve its reliability and replicability.<sup>42</sup> A second, related challenge is that most behavioral studies with infants have reported findings based on a geographically and culturally

restricted sample, causing limited generalizability across populations. This is in part a consequence of the over-representation of researchers in North America and Western Europe, who tend to recruit primarily local families who have the means and the time for a lab visit. The result of this tradition of convenience has been an overreliance on samples of White infants from middle-class families – a biased sample of the world’s population when making conclusions about both human development and human nature.<sup>43,44</sup> Overcoming these challenges is essential to the development of a robust and generalizable model of infants’ visual preferences.

We address these challenges by providing a large-scale, high-powered test of the Hunter and Ames model through an international collaboration, harnessing the infrastructure of ManyBabies.<sup>45</sup> ManyBabies is a six-continent network of scientists who are interested in understanding key theoretical questions about early child development, promoting best practices in developmental research, and broadening participation to include scientists and families from a broad range of communities and cultures. Drawing on this network, we designed an experiment testing visual preferences for familiar and novel stimuli that will include an infant sample of unprecedented size and diversity ( $N = XX$ , from  $XX$  labs in  $XX$  countries), enabling an evaluation of the extent to which the Hunter and Ames model generalizes across cultures and communities. We test three specific hypotheses (see Table 3): Infants’ preferences for novel stimuli will be stronger (1) following longer familiarization time, (2) as infants grow older, and (3) for simpler vs. more complex stimuli.

## **Method**

### **Ethics**

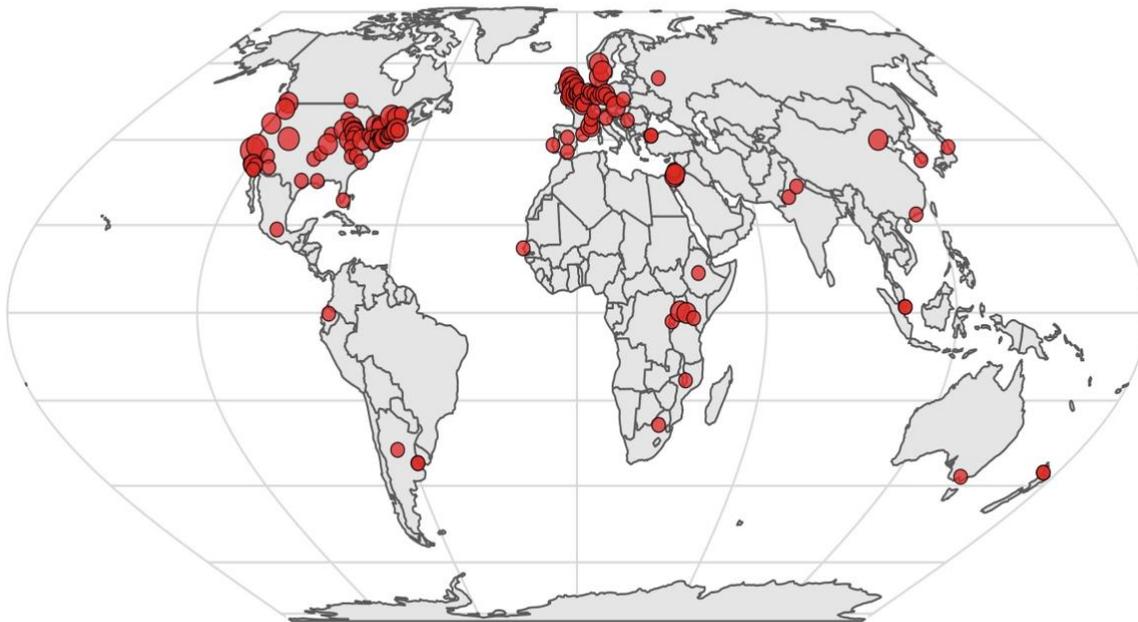
Prior to beginning data collection, all labs contributing data will be asked to complete a survey that asks for information about, among other things, their lab’s ethics approval

procedures. Labs will indicate whether they have new or existing approval from their local ethics committee or institutional review board to conduct this study or whether the current study does not require ethics approval at their institution.

For all labs, we will require the following consent procedure: A parent or legal guardian will provide informed consent for each participating infant, though the specifics of consenting procedures will vary across labs to comply with local legal requirements and cultural norms. Families will be compensated according to each lab's standard practices as approved by their local ethics board. All de-identified data will be stored on the ManyBabies5 OSF repository ([https://osf.io/g3udp/?view\\_only=0f4f6e3b48d8456999d6459c0bfe5510](https://osf.io/g3udp/?view_only=0f4f6e3b48d8456999d6459c0bfe5510)).

### **Participation Details**

An initial open call to participate in “ManyBabies 5: Hunter and Ames” was issued on July 23, 2020 via social media and developmental science listservs. Over the next two years, scientists from around the world (see Figure 2) contributed in various ways to study design. The initial Stage 1 manuscript was submitted for review on January 10, 2023. Between February 1, 2023 and June 30, 2023 we will address reviewer comments, finalize study design and procedural documents, and work with individual teams to get IRB approval. Data collection will begin on July 1, 2023 with a goal of finishing one year later on June 30, 2024. Data analysis and Stage 2 manuscript writing will begin on July 1, 2024 with a target submission date of December 31, 2024.



*Figure 2.* Map depicting the geographic locations of the ManyBabies 5 contributors.

Each participating lab will be asked to contribute a sample of at least 32 infants between the ages of 3 and 15 months; effort will be made across labs to ensure that age is distributed evenly across this age span, to the extent possible. Because many of our analyses will examine effects across labs rather than within a single lab, we will also allow labs to contribute a “half sample” of 16 infants. As in the ManyBabies1 project<sup>46</sup>, allowing “half samples” will increase the number of labs capable of contributing to data collection and will enable participation from more labs, especially those from under-resourced and/or underrepresented communities. We will ask that the sample size per lab (a full sample of 32 or half-sample of 16) includes any infant that

enters the lab and not the number of infants retained after exclusion criteria are implemented because final decisions about participant exclusion will take place centrally.

## **Participants**

The final sample will include XX infants between the ages of 3 and 15 months, recruited and tested by XX labs across XX countries and XX continents (see Table 2 for by-lab details). Based on initial surveys of potential contributors, we expect to be able to recruit a sample size of at least 1,280 infants (at least 40 labs plan to contribute data from approximately 32 infants each). The 3- to 15-month age span was chosen because it covers the majority of ages for which the Hunter and Ames model has been used to describe infants' responses to stimuli in previous research. Because the ManyBabies model makes it possible to collect a sizable sample of infants with a variety of developmental histories, we will adopt an inclusive recruitment strategy. Labs will be encouraged to recruit (using lab-standard practices) any infant who meets just two criteria: (1) the infant's age falls within the 3- to 15-month age range, and (2) the infant has no known issues that would directly impede their ability to process visual stimuli (i.e., visual impairments). However, we will additionally collect data about the criteria on which infants are typically excluded in developmental studies, such as prematurity, developmental delays or disorders, family history of colorblindness, and hearing status. For the analyses of interest in the current paper, we will exclude infants when there is a reason to expect that the predictions of our model would differ based on their developmental history (such as prematurity; see inclusion criteria below). However, all data will be retained for potential future analyses beyond the scope of the current paper (e.g., investigating the effects of prematurity on infants' preferences for novel versus familiar stimuli<sup>47</sup>).

In some labs, we expect that infants will also be tested in other experiments during their visit. In these cases, we will strongly encourage labs to administer the current study first because it is commonly observed in infant studies that “second session” experiments result in greater dropout rates. If labs absolutely must administer this study second, we will ask them to report whether infants participated in the current study first or second and, when the infant participated in the current study second, we will ask for information about the first study. All infants will be included in the primary analyses (following previous ManyBabies studies<sup>46</sup>), but information about first versus second session will be retained for exploratory analyses examining the effect of first versus second session testing.

**Table 2**

*Overview of participating labs and contributed participants (see below for additional details about method and procedure).*

<b>Lab Name</b>	<b>University</b>	<b>Method</b>	<b>Procedure</b>	<b>Mean age (days)</b>	<b>N</b>	<b>Country</b>
XX	XX	central fixation / eye-tracking	fixed-length / infant-controlled	XX	XX	XX
...	...	...	...	...	...	...

### ***Participant Demographics***

Basic demographic data will be collected for all participants using the ManyBabies Demographics questionnaire (Singh, Barokova, Baumgartner, Lopera, Okyere, et al., unpublished manuscript). This questionnaire asks about participants’ biographical information, race and ethnicity, gestational age, caregiver information, family socio-economic status, language exposure, and relevant developmental concerns. Researchers from different communities, nations, and cultures will be able to modify the content of the questionnaire in

structured ways to ensure appropriateness of the questions for local contexts while preserving, to the extent possible, comparability across test sites.

## Materials

*Visual Stimuli.* In the current experiment, we chose to operationalize *task difficulty* in terms of stimulus complexity: we assume that task difficulty increases as stimuli become more complex. Because the sample size possible in ManyBabies projects provides a unique opportunity to test generalizability across multiple stimulus types, we chose two different categories of stimuli: *fribbles*<sup>48-52</sup> and *fractals*<sup>53,54</sup>. Fribbles are colorful, computer-generated models of 3D figures. All fribbles are made up of one of three possible body shapes and up to four appendages that can be chosen from a set of possible options for heads, legs, and two parts of a tail. Fractals are geometric shapes in which the same structure is repeated iteratively at different levels of scale. These two classes of stimuli were chosen because: (1) they allow for variation in stimulus complexity within the same stimulus category (i.e., by manipulating the number of features); (2) infants are unlikely to have previous experience with either set of stimuli; and (3) the inclusion of two stimulus sets allow us to begin examining the generalizability of any observed effect.

We created 12 unique fribbles and 12 unique fractals, operationalizing complexity as the number of features present (these features are appendages in the case of fribbles and iterations of the same structure for fractals; see Figure 3). Within each stimulus type, 6 items were designed to be low-complexity and 6 were designed to be high-complexity. Specifically, low-complexity fribbles are constructed of one body shape and one appendage while high-complexity fribbles have one body shape and four appendages. In the case of fractals, low-complexity items consist

of 3 iterations of the same fractal pattern whereas high-complexity items are made of 9 iterations of the same pattern.

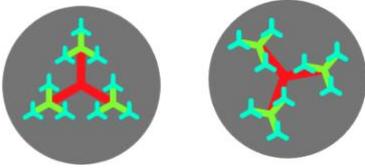
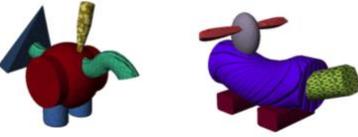
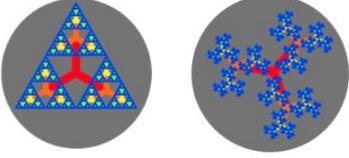
	Fribbles	Fractals
Low Complexity		
High Complexity		

Figure 3. Examples of low- and high-complexity Fribble and Fractal stimuli.

## Procedure

Full procedural instructions provided to each lab will be available in Supplementary Materials on the OSF. Before data collection begins, all labs will be asked to complete a pre-data-collection survey (which will be viewable on the OSF) and submit a walkthrough video depicting a standard testing session (described in further detail below). During the experiment, infants will be seated on their caregiver's lap or in a high-chair or car seat (whichever option corresponds to labs' standard procedures for testing infants, as reported in the pre-data-collection survey). Each trial will be made up of a familiarization phase, in which infants are first exposed to a stimulus, and two test phases where infants are presented with both the item they viewed during the familiarization phase and a novel test item. These phases will be repeated on each of 12 total trials and are described in further detail below.

***Infant-controlled versus fixed-length methods.*** Familiarization can be achieved either by presenting a fixed amount of exposure, i.e. a *fixed-length design*, or by ending the

familiarization phase of an experiment after infants have acquired a certain amount of looking at the target, i.e. an *infant-controlled design*. Both approaches have been used in the literature. In the fixed-length condition the familiarization stimulus is presented for a fixed duration regardless of how much the infant looks.<sup>55</sup> This procedure allows for variation in how much time infants look at the stimulus before their visual preference is assessed while controlling for the amount of time from the start of the familiarization phase to the test phase. Thus, in the fixed-length design, the length of the familiarization phase is fixed while the duration of visual attention directed towards the stimulus varies.

In the infant-controlled design, in contrast, the familiarization stimulus is presented until infants accumulate the same amount of looking.<sup>4,56</sup> This procedure ensures that all infants receive the same amount of familiarization before their preferences for novel stimuli are tested. It also means that the length of the familiarization phase is determined by how long it takes infants to accumulate the required amount of looking (e.g., if they have one or two long looks, with little looking away time, or multiple short looks interspersed with variable length looks away). Thus, in the infant-controlled design, the length of the familiarization phase varies while the duration of infants' visual attention directed towards the stimulus remains consistent.

The consensus across the ManyBabies 5 community was that an infant-controlled design, which facilitates equating infants' familiarization time, represented the best test of our main research questions. Participating labs will be encouraged to use and supported in their use of an infant-controlled procedure. However, because infant-controlled procedures require either well-trained testers who can reliably code infant looking time in the moment or eye-tracking systems that can support gaze-contingent protocols, it may not be possible for some labs to use this procedure. In line with ManyBabies' philosophy of inclusive participation, we will attempt to

support all labs in implementing infant-controlled familiarization, but allow for use of a fixed-length procedure in cases in which infant-controlled is not possible (i.e., labs lacking the required personnel or equipment to implement an infant-controlled procedure). This will allow labs to flexibly choose the option that works best for their resources and maximizes overall data collection, while also working towards a high proportion of labs implementing the desired procedure. In total, XX labs will use an infant-controlled procedure and XX labs will use fixed-length familiarization (as reported in the pre-data-collection survey). To ensure that any observed differences across labs using infant-controlled and fixed-length procedures will not be due to selection effects (as labs self-selected into one paradigm or the other), a subset of XX labs who use the infant-controlled design will be asked to collect a second sample of infants using a fixed-length procedure.

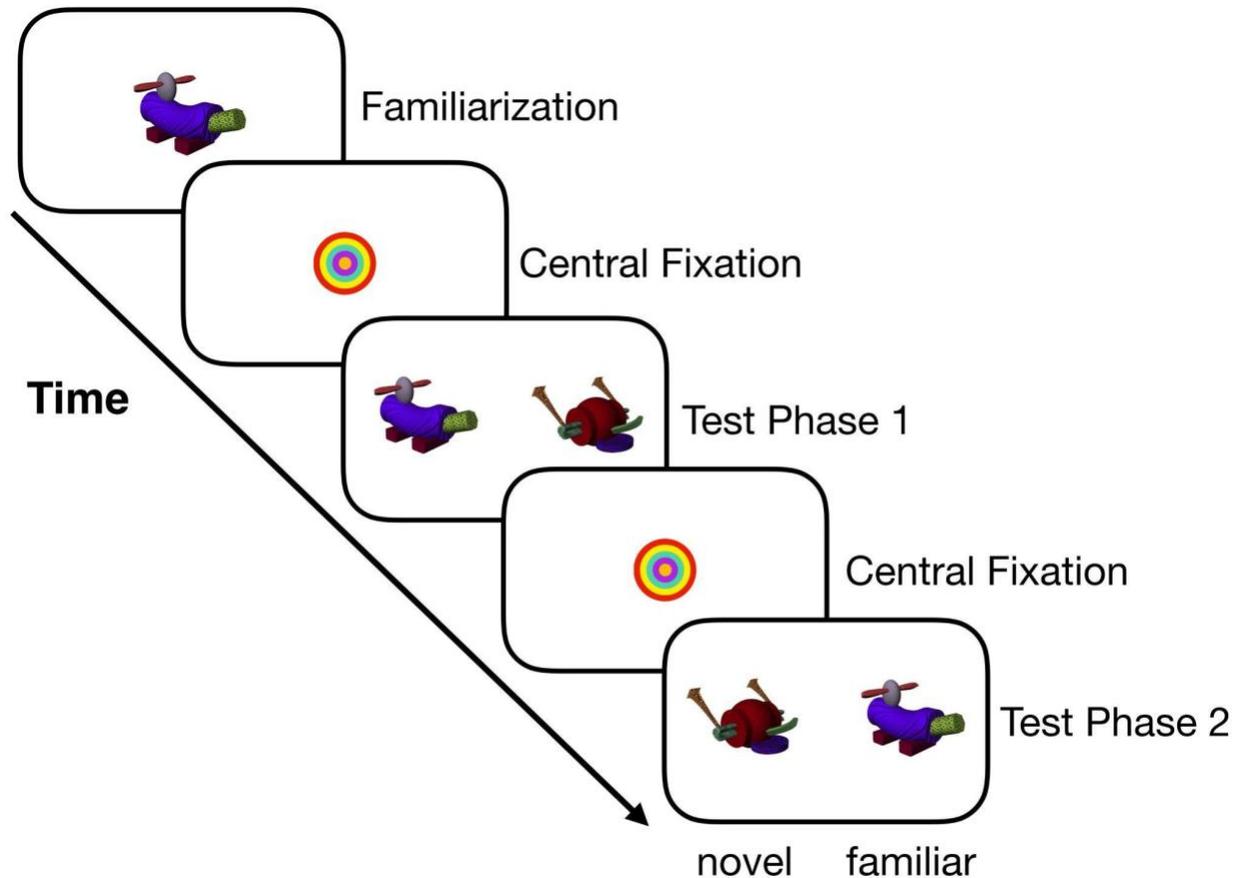
***Trial structure.*** The trial sequence is patterned after classic work by Rose and colleagues.<sup>4</sup> On each trial, a single stimulus is presented centrally on the screen for a familiarization phase, followed by two test phases in which the now-familiar stimulus is paired with a novel stimulus (the novel stimulus will be the same across the two test phases). Following Rose and colleagues<sup>4</sup>, each test phase will be 5 s in duration, and they will differ only in the left-right placement of the novel and familiar stimulus, i.e. the location of the paired test item will be switched between the two test phases.

In order to test the effect of *familiarization time*, we will vary the duration of the familiarization phase, yielding three types of trials. In the infant-controlled design, following Rose and colleagues<sup>4</sup>, the familiarized stimulus will be presented until infants accumulate 5, 10, or 15 s of looking to the familiarization stimulus (thus, the actual trial length will vary across infants depending on how long it takes each infant to accumulate the required looking time). We

will additionally implement a maximum trial length criterion; if infants do not accumulate the required looking time within twice the target familiarization time (i.e., 10, 20, or 30 s), the familiarization phase will end. In the fixed-length design, the familiarized stimulus will be presented for 5, 10, or 15 s total, regardless of the duration of infants' looking.

Immediately following familiarization (in both designs), a central-fixation stimulus – looming circles accompanied by chimes (as used in ManyBabies 1<sup>46</sup>) – will be presented in the center of the screen for a fixed duration of 500 ms to reorient infants' gaze in both the infant-controlled and the fixed-length design. The central-fixation stimulus will be immediately followed by the two test phases, which each will be a fixed length of 5 s, for a total of 10 s of test.<sup>4</sup> The test phases will be a fixed duration, even if familiarization was infant-controlled. In each test phase, infants will now see a familiar and novel item side by side, left-right position of the two stimuli reversed on the second phase. The novel item for each test will be from a different family within the same stimulus set (i.e., fribbles or fractals) that is the same level of complexity as the one they were familiarized with. The central-fixation stimulus will additionally be presented between the two test phases. Figure 4 depicts the structure of an example trial.

To retain infants' attention between trials, an attention-getter will be played before the first trial and between every trial. Labs will have the option of using a default attention-getter video of a laughing baby, which is effective in drawing infants' attention to the screen, but will be encouraged to use any attention-getter that is a better match for their lab or community. For example, a different attention-getter may be needed for researchers working with deaf or hearing-impaired infants.



*Figure 4.* This figure depicts the design of an example trial. At familiarization, a single image is shown for 5, 10, or 15 s. After the desired familiarization time is reached (based on the infant-controlled or fixed-length design criteria), infants are presented with a central fixation stimulus for 500 ms. During the two test phases (5 s each), infants view the same image to which they were familiarized paired with a new stimulus from the same set (e.g., fribbles or fractals) and the same level of complexity. The side on which the familiar and novel images are presented is counterbalanced across test phases, but the images remain the same.

**Counterbalancing.** Each infant will view both stimulus sets (fribbles and fractals), with presentation blocked by stimulus set and presentation order (fribbles first or fractals first) and counterbalanced across infants. Additionally, within each stimulus set, infants will experience all combinations of complexity level and familiarization time. As a result, infants will view a total of 6 trials per stimulus set, or 12 trials total. If infants successfully view all 12 trials, this represents approximately 4.5 minutes of looking to the screen (note that the total study time will

increase when the amount of time it takes for infants to accumulate each familiarization time is taken into account in the infant-controlled version).

We will additionally impose a number of constraints during counterbalancing. First, we will ensure that: (1) the familiar item does not occur on the same side in Test Phase 1 on more than two trials in a row, (2) no more than two trials in a row feature stimuli from the same level of complexity, (3) the same familiarization time is not used for more than two trials in a row. Additionally, each familiar and novel stimulus will only be used in one trial, as repeating stimuli will impact how familiar or novel a given stimulus is for the infant. Across infants, we will counterbalance which stimulus is used as the familiar or novel stimulus.

**Apparatus.** To facilitate standardization across labs, we will create infant-controlled and fixed-length versions of the experiment on a variety of software platforms (e.g., Habit, PyHab, EyeLink Experiment Builder, Tobii Studio; or, in the case of the fixed-length design, experimenters will be able to use video-presentation software<sup>57,58</sup>). Individual labs will be instructed to present the program using the setup with which they are most familiar (e.g., TV screen, projection screen, or computer monitor). Labs will additionally choose to collect data using central fixation methods or an eye tracker (see Table 2).

**Coding.** In labs that assess looking time by coding all or part of their videos offline, coding of infants' looking to the familiar versus novel stimulus at test (for both the infant-controlled and fixed-length procedures) and during the familiarization phase (during the fixed-length procedure only) will be conducted via standard procedures in each lab. We will additionally provide a "best practices" manual that includes information about training and reliability standards that can be used by labs without existing standard procedures or those interested in revising their current procedures.

While labs will be allowed to code videos according to their existing lab practices, labs will be asked to report inter-coder reliability by independently re-coding a minimum of 20% of videos and reporting the average frame agreement, i.e., the proportion of frames on which both coders agree on the gaze location. The target reliability for this coding is 90% frame agreement.

***Minimizing bias.*** To minimize caregiver bias, each lab will be asked to ensure (using lab-standard methods) that caregivers remain unaware of the visual stimuli being presented (e.g., by asking them to wear darkened glasses, close their eyes, or keep their gaze directed toward the infant's head). To minimize experimenter bias, experimenters making online decisions about infants' looking to or away from the screen will not be aware of which visual stimulus is being presented nor the side of the familiar stimulus during the test phase. Off-line coding, including reliability coding, will be conducted with these same efforts to minimize bias. Reliability coders will also not have access to the original coder's decisions.

### **General Lab Practices**

Each lab will be responsible for maintaining good practices, conducting this study using the same rigorous standards applied to all other studies in their lab. Labs will be asked to provide information about their standard protocols for testing, such as researcher training practices and basic practices for greeting families, obtaining consent, and debriefing. Labs that are new to infant testing will be paired with more established laboratories for support and guidance throughout the data collection process.

Prior to data collection, labs will be asked to submit a walkthrough video depicting a standard testing session. This video can be of an infant participant or of a placeholder toy (e.g., a teddy bear). Before labs are invited to move on to data collection, this video will be reviewed by the ManyBabies 5 study implementation team to address any questions or inconsistencies that

arise in study implementation. Labs will also be asked to upload a practice data file to ensure that each lab's data format is in compliance with the ManyBabies 5 data structure. After data collection is complete, participating investigators will submit their data to the analysis team. We will ask labs to refrain from analyzing the data independently prior to submission and analysis by the ManyBabies 5 team (with exceptions for trainees' educational timelines). Labs will be asked to provide experimental and participant data for any participant who enters the lab to participate regardless of whether the participant began or completed the study. The purpose of this request is to ensure that researchers do not selectively submit data based on their own impressions of data quality. For each participant, labs will be asked to report on any unusual events, such as equipment failure, parental interference, infant fussiness, or infants taking a break during the session. Whenever possible (i.e., the lab has appropriate permissions from an ethics board), sessions will be video recorded, and video recordings will be submitted together with coded data or uploaded to repositories such as Databrary.

### **Inclusion Criteria**

All data collected for the study will be handed to the analysis team for confirmatory analyses (i.e., demographic and experimental data for every infant who entered a participating lab, regardless of how many trials they complete). As mentioned previously, infants will be included in the primary analyses if they meet the following criteria: (1) their age falls within the 3- to 15-month age range, (2) they have no known vision or eye movement problems (e.g., no nystagmus), (3) they were born at 37 or more weeks gestation (if known) and/or do not meet local definitions of preterm birth, and (4) they do not have known developmental delays. In some cases, infants who do not meet these criteria will have been recruited in error (e.g., infants recruited outside of the target age range). Thus, XX infants (XX% of the total sample) will be

excluded for failing to meet the recruitment criteria. Additionally, infants will be excluded for session-level errors, such as experimenter error, equipment failure, or other forms of interference that impede infants' ability to attend to at least one trial in the experiment or that affect all trials ( $N = XX$ ,  $XX\%$  of infants remaining after recruitment-based exclusions have been implemented). All infants will otherwise be eligible for inclusion in analyses.

Finally, we will exclude individual trials for which issues are reported (e.g., infant fussiness, incorrect stimulus, single instance of parental interference, failure to accumulate the required familiarization time). A total of  $XX$  ( $XX\%$ ) trials will be excluded due to trial-level errors. To be included in the final analyses, eligible infants (as defined above) will be required to contribute nonzero looking times for at least 1 trial after trial-level exclusions are applied.  $XX$  infants ( $XX\%$ ) will be excluded for failing to meet this criterion.

Table 3.

Design Table.

Question	Hypothesis	Sampling Plan	Analysis Plan	Interpretation given to different outcomes
How does familiarization time influence infants' novelty preference?	<b>(H1A)</b> As familiarization time increases, infants' novelty preference increases.	The design ( $n=1,280$ ) provides 96.8% power to detect an effect of $d=0.5$ or larger (see Table 5).	We will use a linear mixed-effects model to predict infants' novelty preference (proportion looking to the novel stimulus) from familiarization time <b>(H1A)</b> , age <b>(H1B)</b> and stimulus complexity <b>(H1C)</b> . The model will include all two- and three-way interactions between the three predictors (these correspond to exploratory tests for interactions between the main predictors; Table 4). The model will include random effects for participant, item, and lab (see main text for full model specification). The three simple effects of familiarization time, age, and stimulus complexity correspond to the three main hypotheses of interest.	A significant coefficient in the predicted direction <b>(positive: familiarization time, age; negative: stimulus complexity)</b> will be interpreted as evidence consistent with the corresponding hypothesis. A significant coefficient in the opposite direction will be interpreted as providing evidence in the opposing direction from that predicted by the Hunter & Ames model. A non-significant coefficient will be interpreted as not providing evidence for or against the corresponding hypothesis (i.e., we cannot reject the null hypothesis that the corresponding predictor has no effect on infants' novelty preference).
How does novelty preference change as infants age?	<b>(H1B)</b> As infant age increases, infants exhibit a stronger novelty preference.	The design ( $n=1,280$ ) provides >99% power to detect an effect of $d=0.5$ or larger (see Table 5).		
How is novelty preference affected by the complexity of stimuli?	<b>(H1C)</b> Novelty preference is larger when the familiarized stimulus is simpler than when it is more complex.	The design ( $n=1,280$ ) provides >99% power to detect an effect of $d=0.5$ or larger (see Table 5).		

## Data Analysis Plan

### Dependent and Independent Variables

The analyses will include the following variables:

- **Novelty preference (model term: novelty\_preference).** Infants' preference for the novel stimulus over the familiar stimulus during the entire test phase for each trial (summed across the two test phases) will be measured as the proportion looking to the novel stimulus.<sup>4</sup>

$$\text{novelty preference} = \frac{\text{novel looking time}}{\text{novel looking time} + \text{familiar looking time}}$$

- **Infant age (age).** Infant age (mean-centered and scaled), will be a continuous predictor.
- **Familiarization time (familiarization\_time).** Familiarization time will be treated as a continuous predictor and centered. Specifically, the three familiarization times 5, 10, and 15 s will be coded as -0.5, 0, 0.5.
- **Stimulus complexity (stimulus\_complexity).** The two levels of stimulus complexity will be centered and coded as -0.5 (low) and 0.5 (high).
- **Participant.** The unique identifier for each individual infant participating in the study.
- **Item.** The unique familiar/novel stimulus pair in a given trial.
- **Lab.** The unique identifier for each individual lab that collected data for the study.

### Statistical Modeling Approach

**Modeling Approach.** In the main model, we will use a linear mixed-effects model to predict infants' novelty preference from age, familiarization time, and stimulus complexity, as well as all two- and three-way interactions between the three predictors. We will include random effects for participant, item, and lab. The planned model will include the maximal random effects structure<sup>59</sup>, which we will prune as necessary to allow the model to converge (see

[Supplementary Materials](#) S1 for details on our planned approach for handling model non-convergence). Models will be fit using the lme4 package in R<sup>60,61</sup>, and  $p$ -values will be estimated using the Satterthwaite approximation implemented in the R package lmerTest.<sup>62</sup>

**Model.** The main model will be specified as:

```
novelty_preference ~ 1 + age + familiarization_time + stimulus_complexity +
  age : familiarization_time +
  age : stimulus_complexity +
  familiarization_time : stimulus_complexity +
  age : familiarization_time : stimulus_complexity +
  (1 + familiarization_time * stimulus_complexity | participant) +
  (1 + age * familiarization_time * stimulus_complexity | lab) +
  (1 + age * familiarization_time | item)
```

### Hypothesis Tests

Our main hypotheses are that familiarization time, age, and stimulus complexity will each systematically predict infants' novelty preference (see Table 3). In addition to the main effects of interest, the model will also allow us to test for two- and three-way interactions between the main effects of interest. These possible interaction effects are not the main focus of our confirmatory analyses, and thus we will interpret these tests as exploratory (rather than confirmatory). Table 4 lists the question tested by the two- and three-way interactions in the model and how significant coefficients for each of these terms will be interpreted.

**Table 4.**

*Interpretation of the interactions in the linear mixed-effects model*

<b>Question</b>	<b>Model term</b>
To what extent does the effect of familiarization time on novelty preference change with infant age?	age * familiarization_time
To what extent does the effect of stimulus complexity on novelty preference change with infant age?	age * stimulus_complexity
To what extent does the effect of stimulus complexity on novelty preference change as familiarization time increases?	familiarization_time * stimulus_complexity
To what extent does the interaction between familiarization time and complexity of the stimulus depend on infant age?	age * familiarization_time * stimulus_complexity

### **Power Analysis**

We used a simulation-based approach to evaluate whether the expected sample size (i.e., 1,280 infants) can provide sufficient power to detect effect sizes of interest. All code associated with these simulations is publicly available on the project's OSF page ([https://osf.io/g3udp/?view\\_only=0f4f6e3b48d8456999d6459c0bfe5510](https://osf.io/g3udp/?view_only=0f4f6e3b48d8456999d6459c0bfe5510)), and the main simulation-based results are summarized in our Data Simulation and Power Analysis [Supplement](#). In order to calculate the statistical power from this expected sample size, we simulated 500 datasets with a Cohen's  $d$  effect size of 0.2, 0.3, 0.4, and 0.5 for the three main effects of interest (i.e., *stimulus type*, *age*, and *familiarization time*) as well as their two- and three-way interactions. We chose these effect size magnitudes based on estimates from previous large-scale preferential-looking studies (e.g., in ManyBabies 1, the main meta-analytic effect size was  $d = 0.35$ , with effect sizes ranging from  $d = 0.24$  to  $d = 0.51$  depending on the experimental method) as well as median effect size estimates in meta-analyses within infant populations.<sup>40,46</sup>

For each of these simulated datasets, we ran a linear mixed-effects regression model with the following structure:

$$\begin{aligned} \text{novelty\_preference} &\sim 1 + \text{stimulus\_complexity} * \text{age} * \text{familiarization\_time} + \\ &(1 + \text{stimulus\_complexity} * \text{familiarization\_time} | \text{participant}) + \\ &(1 | \text{lab}) + \\ &(1 | \text{item}) \end{aligned}$$

Based on preliminary simulations, we chose to focus on this model because it was the maximal random effects structure that reliably converged on the vast majority of simulation runs (i.e., >90%). Given past evidence that models including random slopes may not converge<sup>46</sup>, we also simulated power for a model with a simpler random effects structure that included only random intercepts for participants, labs, and items. We then evaluated the statistical power by counting the number of times the effect displayed a significantly non-zero (i.e.,  $p < .05$ ) coefficient. The main goal was to establish that we had sufficient power to detect each of the three main effects of interest (age, familiarization time, and stimulus type), i.e., to test the three main predictions of the Hunter and Ames model. As shown in Table 5, the analysis indicates ample power to detect each of these three main effects for medium effect sizes (>80% power for all main effects and  $d = 0.4$  to  $0.5$ ), with somewhat weaker power to detect a main effect of familiarization time at the lower end of the effect size range ( $d = 0.3$ : power = 63.4%;  $d = 0.4$ : power = 86.4%). The power analysis also demonstrates reasonable power to detect age and any interactions, including age, even if the effect size is small (i.e.,  $d = 0.2$ ). Notably, the current design has reasonable power to detect a three-way interaction between the main effects (>80% power for  $d = 0.2$  or larger). Overall, the power analysis indicates that the current design has sufficient power to detect each of the three main predictions of the Hunter and Ames model.

**Table 5.**

Overview of results in power simulation for the model with the maximal converging random effects structure. The numbers in each effect size column indicate the percentage of significant results in the 500 simulated datasets. *stim\_type* refers to stimulus type, *fam\_time* refers to familiarization time, and *age* refers to infant age in months. Sample size is 1,280 included participants, with 24 trials. See section 7 in the Data Simulation and Power Analysis [Supplement](#) for details.

<b>Outcome Types</b>	<b>Model Terms</b>	<b><i>d</i> = 0.2</b>	<b><i>d</i> = 0.3</b>	<b><i>d</i> = 0.4</b>	<b><i>d</i> = 0.5</b>
<b><i>main effects</i></b>	<i>stim_type</i>	47.4%	83.8%	96.6%	99.2%
	<i>age</i>	99.0%	100%	100%	100%
	<i>fam_time</i>	36.0%	63.4%	86.4%	96.8%
<b><i>two-way interactions</i></b>	<i>stim_type*fam_time</i>	12.4%	24.4%	35.0%	48.6%
	<i>stim_type*age</i>	89.4%	98.6%	100%	100%
	<i>fam_time*age</i>	86.4%	96.6%	99.4%	100%
<b><i>three-way interaction</i></b>	<i>age*stim_type*fam_time</i>	80.8%	85.6%	93.8%	95.6%

### **Robustness and Exploratory Analyses**

The analytic strategy described above represents our main confirmatory approach, consistent with contemporary analytic approaches used in the infant literature, and our main conclusions will be based on the results of these confirmatory models. However, the large dataset

collected in the current study presents a unique opportunity to explore the relationship between the underlying assumptions of these models and the distributional properties of infant-looking time data. We will therefore examine the robustness of the effects of interest across various analytic decisions and inclusion criteria. The main results from the confirmatory models will be interpreted in light of the consistency or variability of the main findings across these exploratory analyses (see S.2 and S.3 in the [Supplementary Materials](#) for details on specific robustness and exploratory analyses).

**Data Availability**

Full data will be publicly available at

[https://osf.io/g3udp/?view\\_only=0f4f6e3b48d8456999d6459c0bfe5510](https://osf.io/g3udp/?view_only=0f4f6e3b48d8456999d6459c0bfe5510).

**Code Availability**

All analysis code is publicly available at

[https://osf.io/g3udp/?view\\_only=0f4f6e3b48d8456999d6459c0bfe5510](https://osf.io/g3udp/?view_only=0f4f6e3b48d8456999d6459c0bfe5510).

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**Author Contributions**

Authorship order was determined as follows: The first two authors are the primary co-leads of the project and share first authorship. The last author was the primary senior researcher supervising the project. All other authors are listed in alphabetical order. An overview of

authorship contributions following the CRediT taxonomy can be viewed here:

<https://docs.google.com/spreadsheets/d/e/2PACX-1vRbYFrQYIUeGdXLEOw7miYzfnSBLR3QKKAQqU52F-J6nGCupIzgQZywez55NzaN9Vbm3cMh-XK-Fr/pubhtml>

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## Supplementary Materials

### S1. Model Non-Convergence and Pruning Random Effects

If the maximal model does not converge using default settings in the lme4 package, our first step will be to attempt a series of remedies to help the model including the full random effects structure achieve convergence (Brauer & Curtin, 2018). Specifically, we will (a) increase the number of iterations in the estimation procedure, (b) provide the model with improved starting values, and (c) check whether convergence is achieved using any of the other optimization functions available with the lme4 package.

If the model does not converge after attempting these remedies, we will next prune random effects until convergence is achieved. We will first remove random effects that are of lesser theoretical interest before removing random effects that are more crucial to the main hypotheses of the study (i.e., random slopes of critical main effects of interest and random intercepts). In practice, we expect that models including random slopes (although consistent with the experiment design) will be difficult to fit and will require further pruning of the random effects structure. For example, in ManyBabies 1 (ManyBabies Consortium, 2020), the maximal random effects structure that allowed the main model to converge included random intercepts only.

We will proceed as follows in pruning random effects step-by-step. In each instance, we would only proceed to the next pruning step if the model still fails to converge. Each of steps 1-3 will be repeated within each of the three types of random effects (item, lab, participant) separately until the model with the overall maximal random effects structure across the three random effect types is achieved. We will first prune random slopes (following steps 1-3) for

item-based random effects, followed by random slopes for lab-based random effects. Random slopes and covariances for participant-based random effects will be pruned last.

1. We will first sequentially remove **random slopes for interaction terms** since the interactions are of lesser theoretical interest (compared to the main effects), checking for non-convergence after each interaction term is removed. Random slopes for the three-way interaction will be removed first, followed by the interaction for stimulus complexity and age, stimulus complexity and familiarization time, and finally, familiarization time and age.
2. We will next remove **any covariances between random effects that approach 0 or 1** (likely leading to a singular fit in the main linear mixed-effects model).
3. If convergence is still not achieved, we will successively remove **random slopes for each of the main effects** of interest. We will remove random slopes in the following order, checking for convergence after each random slope is removed: (a) stimulus complexity, (b) infant age, and (c) familiarization time.

Finally, if the model still fails to converge after removing all random slopes for item, lab, and participant, we will successively remove **random intercepts**, first removing random intercepts for item, followed by random intercepts for lab and (if necessary) random intercepts for participant last.

## **S2. Analysis of Model Assumptions and Robustness**

In this section, we outline a series of analyses to test for violations of model assumptions and to investigate the robustness of the results from the main model across alternative model specifications.

### **S2.1. Model Diagnostics**

First, we will conduct diagnostic tests to inspect two key properties of the data and model: missingness and heteroskedasticity.

### **S2.1.1. Missingness**

We expect there to be a significant amount of missing data due to a variety of infant-related (e.g., “fussiness”) and experimenter-related (e.g., technical errors) factors. A relatively large proportion of missing data is common in infant research (e.g., around 20% of the data in ManyBabies1; The ManyBabies Consortium, 2020). Taking missingness into account is particularly important in the context of the current study because the main phenomenon of interest — infants’ visual attention — is systematically connected with a possible major source of missingness, namely infant inattention or lack of engagement leading to trial exclusion or failure to measure infant looking behavior. A possible risk is that our estimates of infants’ patterns of novelty preference may be biased. Therefore, we will investigate whether and how different factors influence the presence of missingness in the data. Specifically, we will fit a logistic mixed-effects model predicting the trial-by-trial presence or absence of the data (i.e., coded as 0 = data is present; 1 = data is missing) from several key predictors: the main predictors of interest (age, complexity, familiarization time) and trial number (the strongest predictor of missingness in ManyBabies1). We will include random effects for participant, lab, and item, following the same approach for specifying and pruning random effects as in the main model. The model will be specified as:

$$\text{NAs} \sim \text{familiarization\_time} + \text{age} + \text{stimulus\_complexity} + \text{trial\_number} +$$

$$(\dots | \text{participant}) + (\dots | \text{lab}) + (\dots | \text{item})$$

In exploratory analyses, we will also test a more complex model including a series of additional predictors of interest (e.g., time of testing, time of last nap). If we find evidence of systematicity

in missingness, we will explore routes for handling this systematicity, in particular conducting multiple imputation, and discuss limitations in the interpretation of the results from the main model in light of missingness issues.

### **S2.1.2. Heteroskedasticity**

A particular concern in evaluating our main model is the possibility of heteroskedasticity. Specifically, it is likely that variance in looking times will increase across trials, which may in turn lead to higher variance in our measures of preferential looking at later trials in the experiment. To check for issues related to heteroskedasticity, we will use the performance package in R (Lüdtke et al., 2021) to conduct a Breusch-Pagan test (Breusch & Pagan, 1979) of non-constant error. If we find evidence of heteroskedasticity, we will explore the use of location-scale modeling and general additive modeling approaches to model and account for sources of variability (e.g., trial number).

### **S2.2. Alternative Link Functions**

We chose the primary dependent variable, infants' novelty preference score, based on its prevalence in the literature on infant preferential looking (e.g., Rose et al., 1982). However, because this dependent measure is a proportion variable, it raises several potential analytical issues. For example, one possible concern is that the residuals (i.e., the error variance) from the model may not be normally distributed, violating an assumption of linear mixed-effects models. Therefore, in the event that the observed data do violate this assumption, we will explore alternative models that incorporate specialized link functions (i.e., beta regression) that may be better suited for the properties of proportion score data.

## **S3. Exploratory Analyses**

Below, we outline a series of analyses we plan to conduct to explore other possible sources of variation in infant-looking behavior and the impact of specific methodological choices (e.g., inclusion criteria, stimulus choices).

### **S3.1. Infant-Controlled vs. Fixed-Length Familiarization**

We will test for a moderating effect of familiarization method (infant-controlled vs. fixed-length familiarization) (a) in the overall dataset and (b) in the subsample of the data contributed by labs randomly assigned to collect both infant-controlled and fixed-length samples. Specifically, we will fit the 4-way interaction model predicting novelty preference from stimulus type, age, familiarization time, and familiarization method.

novelty\_preference ~

1 + age \* familiarization\_time \* stimulus\_complexity \* familiarization\_method +

(1 + familiarization\_time \* stimulus\_complexity | participant) +

(1 + age \* familiarization\_time \* stimulus\_complexity \* familiarization\_method | lab) +

(1 + age \* familiarization\_time \* familiarization\_method | item)

If familiarization method exerts a moderating influence on any of the main predictors of interest, we will interpret its implications in the General Discussion.

### **S3.2. Stimulus Fixation Time vs. Familiarization Manipulation**

In the main confirmatory model, we do not distinguish between infants' accumulated looking to the familiarized stimulus and the duration of the familiarization phase. Since labs can choose to either use an infant-controlled or a fixed familiarization design, this will allow us to assess each of these predictors independently. We will therefore fit a linear mixed-effects model in which we predict novelty preference from the duration of the familiarization phase

(familiarization\_duration) and infants' time fixating the stimulus during the familiarization phase (stimulus\_fixation) while controlling for age and stimulus complexity.

novelty\_preference ~ 1 +

age + familiarization\_duration + stimulus\_fixation + stimulus\_complexity +

(1 + familiarization\_duration + stimulus\_fixation + stimulus\_complexity | participant) +

(1 + age + familiarization\_duration + stimulus\_fixation + stimulus\_complexity | lab) +

(1 + age + familiarization\_duration + stimulus\_fixation | item)

We expect that familiarization duration and stimulus fixation will be correlated. We will therefore run diagnostic tests to check for issues related to multicollinearity (specifically, we will test for variance inflation). If the variance inflation factor indicates a concerning degree of multicollinearity (i.e., exceeds a value of at least 5), we will treat any results from the model with caution. If the model including both factors is high, we will instead consider a model comparisons approach in which we fit two models, one including only familiarization duration (in addition to the other main effects of age and stimulus complexity) and one with only stimulus fixation. This will allow us to investigate how well each of the two possible operationalizations of familiarization time predict preferential looking.

### **S3.3. Investigating Stimulus Types Separately**

One possible concern with the analysis outlined above is that there may be fundamental differences in how infants process the two types of stimuli, fribbles and fractals (and, e.g., their associated levels of relative complexity). In particular, fitting a model including both item kinds may lead to poor model fit if the estimates for the two item types systematically differ. To check for inconsistencies in the model predictions across the two stimulus types, we will therefore fit a linear mixed-effects model in which we predict novelty preference from the 4-way interaction

between the main effects of interest (age, familiarization time, and stimulus complexity) and stimulus type (centered; coded as fribbles = 0.5 vs. fractals = -0.5):

novelty\_preference ~ 1 +

age \* familiarization\_time \* stimulus\_complexity \* stimulus\_type +

(1 + familiarization\_time \* stimulus\_complexity \* stimulus\_type | participant) +

(1 + age \* familiarization\_time \* stimulus\_complexity \* stimulus\_type | lab) +

(1 + age \* familiarization\_time | item)

We will follow the same random effects pruning approach as with the main model. Effects of the model – especially interactions between stimulus type and the effects of age, familiarization time, and stimulus complexity – will be qualified in light of any interactions observed with stimulus type.

### S3.4. Inclusion Criteria

We will also explore the consequences of varying the inclusion criteria on the main effects of interest, in particular:

- **Minimum looking times:** We will also fit the main model at various cutoffs for minimum looking times. Specifically, we will fit the main model after excluding any trials that do not meet successively more restrictive inclusion criteria for a given trial:
  - Trials must include 1 second of looking in each of the two 5s test periods.
  - Trials must include 2 seconds of looking in each of the two 5s test periods.
- **Number of trials completed:** We will fit the main model with successively more strict inclusion criteria based on trial number. We will fit the model when only including infants who complete at least 4, 8, 12, 16, 20, and 24 trials and discuss any observed systematic changes in effect magnitude.

### S3.5. Variation across Demographic Predictors

We will explore the degree to which preferential-looking behavior varies across a range of demographic factors and participant characteristics that will be collected along with infants' looking behavior, including variation across testing location, cultural and linguistic experiences, and the backgrounds of infants and their families.

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# ManyBabies 5: The Hunter and Ames Model of Infant Looking Preference

## Supplementary Materials: Data Simulation and Power Analysis

ManyBabies Analysis Team

### Contents

<b>1</b>	<b>Data Simulation</b>	<b>2</b>
<b>2</b>	<b>Visualisation of Simulated Data</b>	<b>4</b>
2.1	Familiarization: . . . . .	4
2.2	Complexity: . . . . .	5
2.3	Age: . . . . .	6
2.4	Age*Familiarization: . . . . .	7
2.5	Age*Complexity: . . . . .	8
2.6	Familiarization*Complexity: . . . . .	9
2.7	Variability by Lab . . . . .	11
2.8	Variability by Item . . . . .	12
<b>3</b>	<b>Power Calculation with Full Data and Varying Intercepts and Varying Slopes</b>	<b>13</b>
3.1	Effect Size = 0.5 . . . . .	13
3.2	Effect Size = 0.4 . . . . .	13
3.3	Effect Size = 0.3 . . . . .	13
3.3.1	Visualise Estimates for Fixed Effects: . . . . .	14
3.3.2	Visualise Estimates for Random Effects: . . . . .	15
3.4	Effect Size = 0.2 . . . . .	16
3.5	Effect Size = 0.1 . . . . .	16
<b>4</b>	<b>Power Calculation with Full Data and Varying Intercepts</b>	<b>17</b>
4.1	Effect Size = 0.5 . . . . .	17
4.2	Effect Size = 0.4 . . . . .	17
4.3	Effect Size = 0.3 . . . . .	17
4.3.1	Visualise Estimates for Fixed Effects: . . . . .	18
4.3.2	Visualise Estimates for Random Effects: . . . . .	19
4.4	Effect Size = 0.2 . . . . .	20
4.5	Effect Size = 0.1 . . . . .	20

<b>5</b>	<b>Power Calculation with 20 pct. Missing Data and Varying Intercepts and Varying Slopes</b>	<b>21</b>
5.1	Effect Size = 0.5	21
5.2	Effect Size = 0.4	21
5.3	Effect Size = 0.3	22
5.3.1	Visualise Estimates for Fixed Effects:	22
5.3.2	Visualise Estimates for Random Effects:	23
5.4	Effect Size = 0.2	24
5.5	Effect Size = 0.1	24
<b>6</b>	<b>Power Calculation with 50 pct. Missing Data and Varying Intercepts and Varying Slopes</b>	<b>25</b>
6.1	Effect Size = 0.5	25
6.2	Effect Size = 0.4	25
6.3	Effect Size = 0.3	26
6.3.1	Visualise Estimates for Fixed Effects:	26
6.3.2	Visualise Estimates for Random Effects:	27
6.4	Effect Size = 0.2	28
6.5	Effect Size = 0.1	28
<b>7</b>	<b>Overview of Power Simulation Results</b>	<b>29</b>
7.1	Summary Statistics for Power Calculation with Full Data and Varying Intercepts and Varying Slopes:	29
7.2	Summary Statistics for Power Calculation with Full Data and Varying Intercepts:	29
7.3	Summary Statistics for Power Calculation with 20 pct. Missing Data and Varying Intercepts and Varying Slopes:	29
7.4	Summary Statistics for Power Calculation with 50 pct. Missing Data and Varying Intercepts and Varying Slopes:	30
<b>8</b>	<b>Overview of Bias Results</b>	<b>30</b>
<b>9</b>	<b>Increasing Variance with Trial Number</b>	<b>31</b>
9.1	Extracting estimates of increasing variance from the adult pilot data	31
9.2	Visualisation of adult data in proportions	32
9.3	Visualisation of adult data in z-score	33
9.4	Modification of simulation function to include variance increase	34
9.5	Visualise the variance increase	37
9.6	Model building and bias assessment	38
9.7	Test how many of the models violate homoskedasticity	42
9.8	Overview of how many of the models exhibit homoskedasticity	42
9.9	Bayesian robust location-scale regression model	42
9.9.1	Prior-Posterior Update Checks	45
<b>10</b>	<b>Data missing not completely at random</b>	<b>51</b>
10.1	Simulation of missing data according to increasing infant age	51
10.1.1	Simulation of models	52
10.1.2	Visualise Estimates for Fixed Effects:	53
10.1.3	Visualise Estimates for Random Effects:	54
10.2	Simulation of missing data according to increasing trial number	55

10.2.1	Simulation of models . . . . .	56
10.2.2	Visualise Estimates for Fixed Effects: . . . . .	57
10.2.3	Visualise Estimates for Random Effects: . . . . .	58
10.3	Simulation of missing data according to increasing familiarisation . . . . .	59
10.3.1	Simulation of models . . . . .	60
10.3.2	Visualise Estimates for Fixed Effects: . . . . .	61
<b>11</b>	<b>Summary Statistics for Power Calculation with Data missing not completely at random</b>	<b>63</b>
<b>12</b>	<b>Logistic Regression Models</b>	<b>64</b>

# 1 Data Simulation

```
my_sim_data <- function(  
  n_subj      = 1280, # number of subjects  
  n_simple   = 12,   # number of complex stimuli  
  n_complex  = 12,   # number of complex stimuli  
  n_small_fam = 8,   #small familiarization time  
  n_medium_fam = 8, #medium familiarization time  
  n_high_fam = 8,   #high familiarization time  
  n_lab      = 40,  
  
  beta_0 = 0, # intercept; i.e., the grand mean  
  beta_c = 0.3, # main effect for complexity  
  beta_f = 0.3, # main effect for familiarization time  
  beta_a = 0.3, # main effect for age  
  
  beta_ca = 0.3,  
  beta_af = 0.3,  
  beta_cf = 0.3,  
  
  beta_cfa = 0.3, #main effect for interaction between complexity and familiarization.  
  
  subject_0 = 0.2, # by-subject random intercept sd  
  
  subject_c = 0.2, # by-subject slope complexity sd  
  subject_f = 0.2, # by-subject slope familiarization sd  
  subject_a = 0.2, # by-subject slope age sd  
  
  subject_ca = 0.2, # by-subject slope for interaction between age and complexity sd  
  subject_af = 0.2, # by-subject slope for interaction between age and familiarization sd  
  subject_cf = 0.2, # by-subject slope complexity*familiarization sd  
  
  subject_cfa = 0.2, # by-subject slope for interaction between age, complexity and familiarization sd  
  
  subj_rho = .2, # correlations between by-subject random effects  
  
  lab_0 = 0.2, # by-lab random intercept sd  
  
  lab_c = 0.2, # by-lab slope complexity sd  
  lab_f = 0.2, # by-lab slope familiarization sd  
  lab_a = 0.2, # by-lab slope age sd  
  
  lab_ca = 0.2, # by-lab slope for interaction between age and complexity sd  
  lab_af = 0.2, # by-lab slope for interaction between age and familiarization sd  
  lab_cf = 0.2, # by-lab random slope complexity*familiarization sd  
  
  lab_cfa = 0.2, # by-lab slope for interaction between age, complexity and familiarization sd  
  
  lab_rho = 0.2, # correlations between by-lab random effects  
  
  item_0 = 0.2, # by-item random intercept sd  
  
  item_c = 0.2, # by-item slope complexity sd  
  item_f = 0.2, # by-item slope familiarization sd  
  item_a = 0.2, # by-item slope age sd  
  
  item_ca = 0.2, # by-item slope for interaction between age and complexity sd  
  item_af = 0.2, # by-item slope for interaction between age and familiarization sd  
  item_cf = 0.2, # by-item random slope complexity*familiarization sd  
  
  item_cfa = 0.2, # by-item slope for interaction between age, complexity and familiarization sd
```

```

item_rho = 0.2, # correlations between by-item random effects

sigma = 0.3 # residual (error) sd
) { # residual (standard deviation)

# simulate a sample of items
items <- data.frame(
  category = rep(c("simple", "complex"), c(n_simple, n_complex)),
  X_c = rep(c(-0.5, 0.5), c(n_simple, n_complex)),
  familiarization = rep(c("short", "medium", "long"), (n_simple + n_complex)/3),
  X_f = rep(c(-0.5, 0, 0.5), (n_simple + n_complex)/3),
  faux::rnorm_multi(
    n = n_simple + n_complex, mu = 0, sd = c(item_0,
      item_c,
      item_f,
      item_a,
      item_ca,
      item_af,
      item_cf,
      item_cfa), r = item_rho,
  varnames = c("I_0", "I_c", "I_f", "I_a",
    "I_ca", "I_af", "I_cf",
    "I_cfa"))
) %>%
mutate(item_id = faux::make_id(nrow(.), "I"))

# simulate a sample of subjects
subjects <-
  faux::rnorm_multi(
    n = n_subj, mu = 0, sd = c(subject_0,
      subject_c,
      subject_f,
      subject_a,
      subject_ca,
      subject_af,
      subject_cf,
      subject_cfa), r = subj_rho,
  varnames = c("S_0", "S_c", "S_f", "S_a",
    "S_ca", "S_af", "S_cf",
    "S_cfa")
) %>%
  mutate(subj_id = faux::make_id(nrow(.), "S")) %>%
  mutate(X_a = runif(n_subj, min = -0.5, max = 0.5))
#add subject age measure, sample from distribution from -0.5 to 0.5. #subjects$subj_id <- 1:n_subj

labs <- faux::rnorm_multi(
  n = n_lab, mu = 0, sd = c(lab_0, lab_c, lab_f, lab_a,
    lab_ca, lab_af, lab_cf,
    lab_cfa), r = lab_rho,
  varnames = c("L_0", "L_c", "L_f", "L_a",
    "L_ca", "L_af", "L_cf",
    "L_cfa")
) %>%
  mutate(lab_id = faux::make_id(nrow(.), "L"))

#create lab and subj nesting structure
#Number of subjects must be a multiple of number of labs
lab_multiplier = n_subj/n_lab
lab_subj_dict <- data.frame(
  subj_id = subjects$subj_id,

```

```

  lab_id = rep(labs$lab_id,lab_multiplier)
)

# cross subject and item IDs
temp <- crossing(subjects, items) %>%
  left_join(lab_subj_dict, by = "subj_id") %>%
  left_join(labs, by = "lab_id") %>%
  group_by(subj_id, item_id) %>%
  mutate(item_id = sample(item_id)) %>%
  ungroup() %>%
  mutate(trial_num = rep(seq(n_simple + n_complex), n_subj))

temp <- temp %>%
  mutate(
    B_0 = beta_0 + S_0 + L_0 + I_0,

    B_c = beta_c + S_c + L_c + I_c,
    B_f = beta_f + S_f + L_f + I_f,
    B_a = beta_a + S_a + L_a + I_a,

    B_ca = beta_ca + S_ca + L_ca + I_ca,
    B_af = beta_af + S_af + L_af + I_af,
    B_cf = beta_cf + S_cf + L_cf + I_cf,

    B_cfa = beta_cfa + S_cfa + L_cfa + I_cfa,

    e_si = rnorm(nrow(temp), mean = 0, sd = sigma),

    DV = B_0 +
      (B_a * X_a) + (B_c * X_c) + (B_f * X_f) +
      (B_cf * X_c * X_f) + (B_af * X_a * X_f) + (B_ca * X_c * X_a) +
      (B_cfa * X_c * X_f * X_a) + e_si
  )
}

dat_sim <- my_sim_data()

```

## 2 Visualisation of Simulated Data

### 2.1 Familiarization:

```

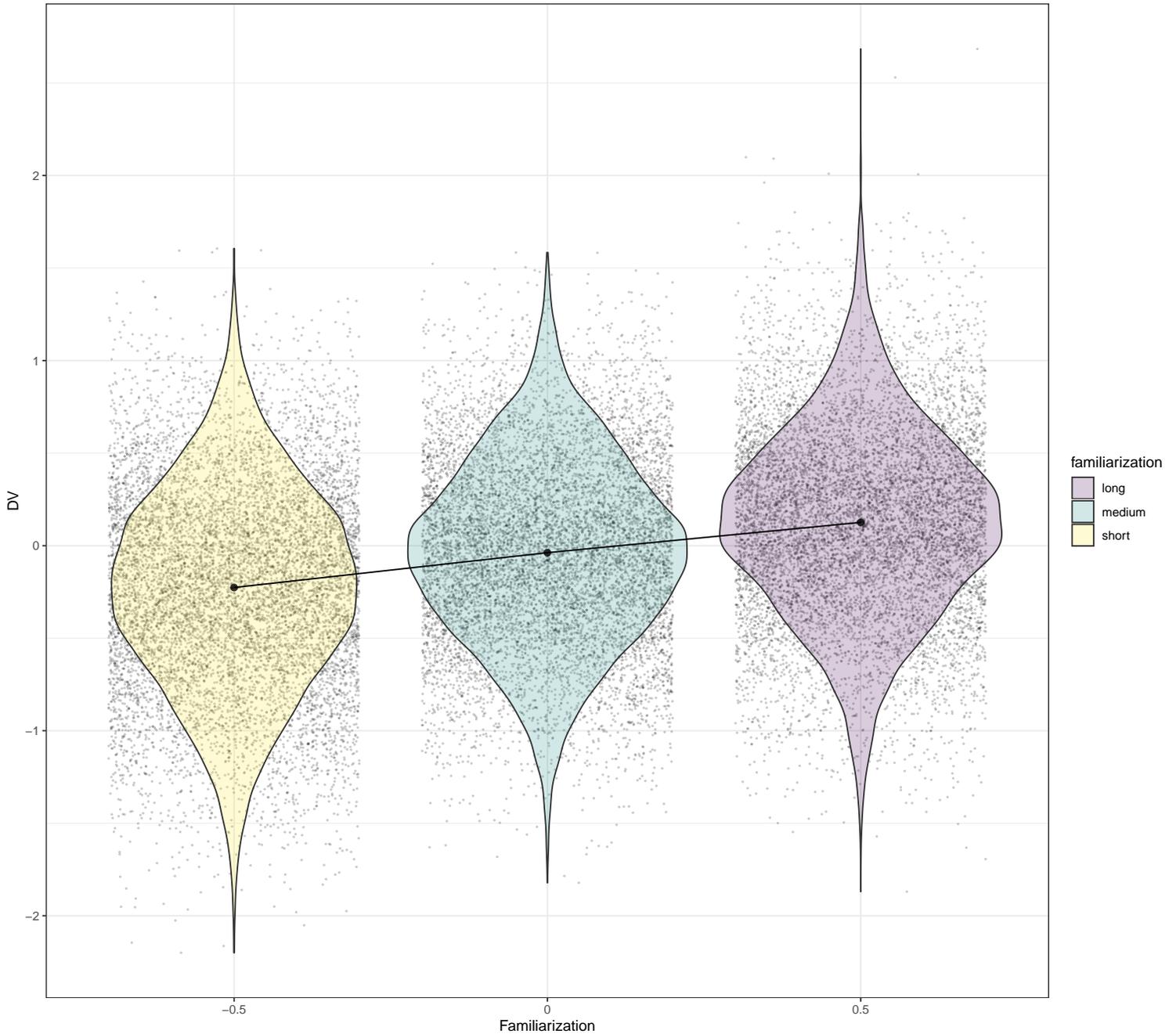
dat_sim_plot_familiarization <- dat_sim %>%
  group_by(X_f) %>%
  dplyr::summarise(med_DV = median(DV))

plot_familiarization <- dat_sim %>%
  mutate(X_f = as.factor(X_f)) %>%
  ggplot() + geom_point(aes(y = DV, x = X_f), position = "jitter",
    alpha = 0.2, size = 0.2) + geom_violin(aes(y = DV, x = X_f,
    fill = familiarization), alpha = 0.2) + geom_line(aes(y = med_DV,
    x = as.factor(X_f), group = 1), data = dat_sim_plot_familiarization) +
  geom_point(aes(y = med_DV, x = as.factor(X_f)), alpha = 0.8,
    size = 2, data = dat_sim_plot_familiarization) + scale_fill_manual(values = viridis(n = 3)) +
  ggtitle("Familiarization") + xlab("Familiarization") + theme_bw()

plot_familiarization <- plot_familiarization + theme(plot.title = element_text(hjust = 0.5,
  size = 20))
plot_familiarization

```

## Familiarization



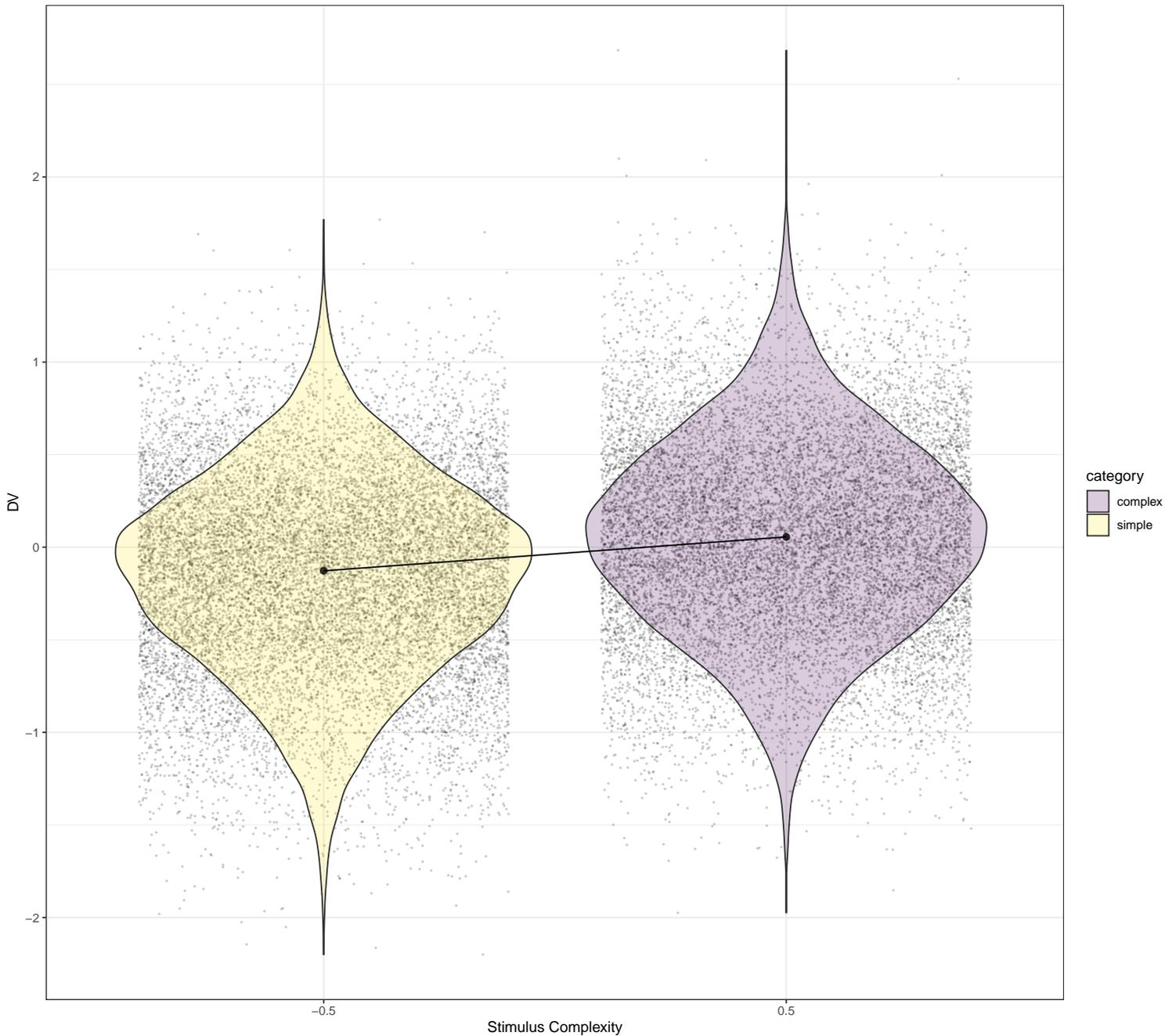
## 2.2 Complexity:

```
dat_sim_plot_complexity <- dat_sim %>%
  group_by(X_c) %>%
  dplyr::summarise(med_DV = median(DV))

plot_complexity <- dat_sim %>%
  mutate(X_c = as.factor(X_c)) %>%
  ggplot() + geom_point(aes(y = DV, x = X_c), position = "jitter",
    alpha = 0.2, size = 0.2) + geom_violin(aes(y = DV, x = X_c,
    fill = category), alpha = 0.2) + geom_line(aes(y = med_DV,
    x = as.factor(X_c), group = 1), data = dat_sim_plot_complexity) +
  geom_point(aes(y = med_DV, x = as.factor(X_c)), alpha = 0.8,
    size = 2, data = dat_sim_plot_complexity) + scale_fill_manual(values = viridis(n = 2)) +
  ggtitle("Stimulus Complexity") + xlab("Stimulus Complexity") +
  theme_bw()
```

```
plot_complexity <- plot_complexity + theme(plot.title = element_text(hjust = 0.5,
  size = 20))
plot_complexity
```

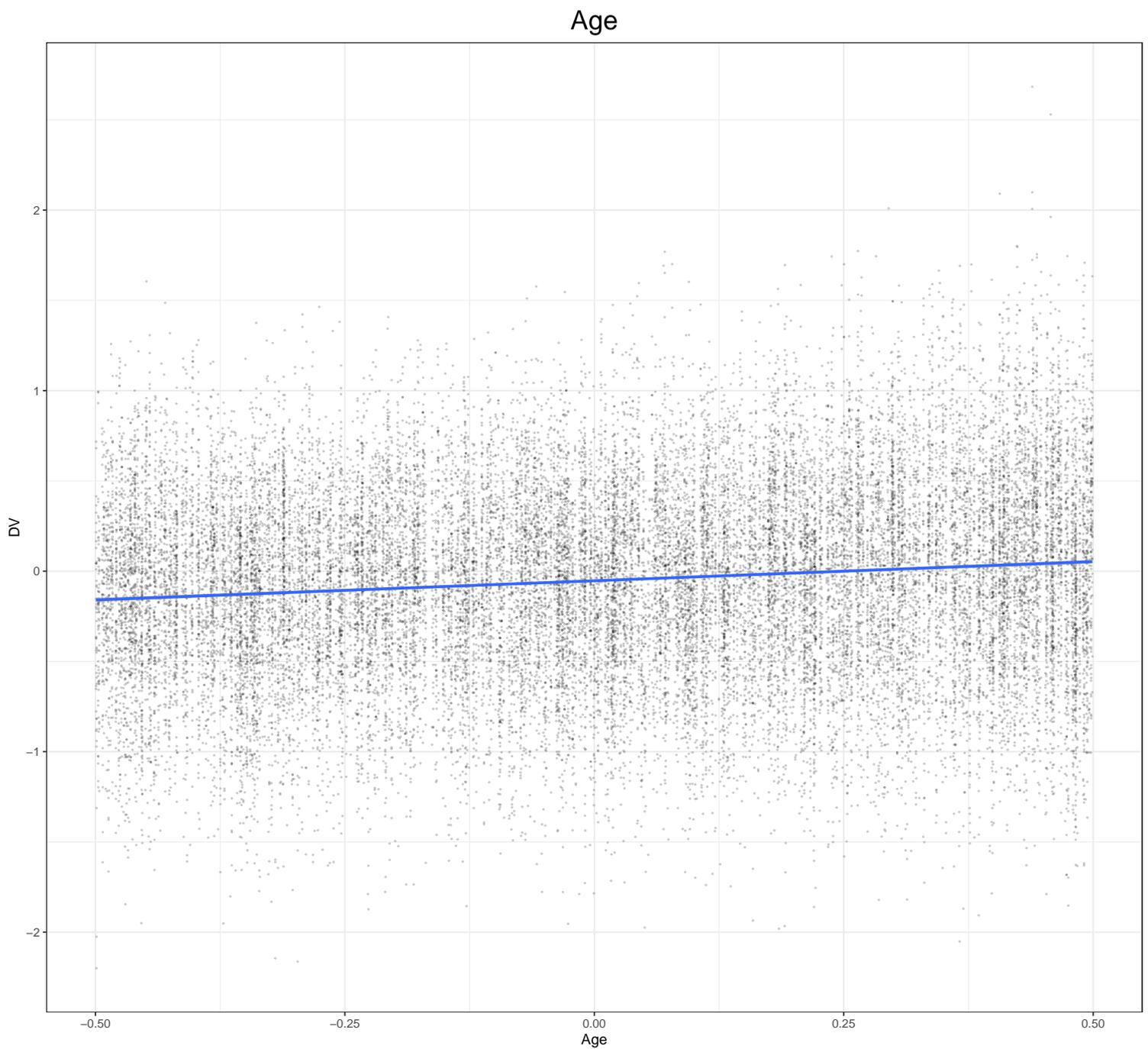
## Stimulus Complexity



### 2.3 Age:

```
plot_age <- dat_sim %>%
  ggplot() + geom_point(aes(y = DV, x = X_a), position = "jitter",
    alpha = 0.2, size = 0.2) + geom_smooth(method = "lm", se = TRUE,
    formula = y ~ x, aes(y = DV, x = X_a)) + ggtitle("Age") +
  xlab("Age") + theme_bw()
```

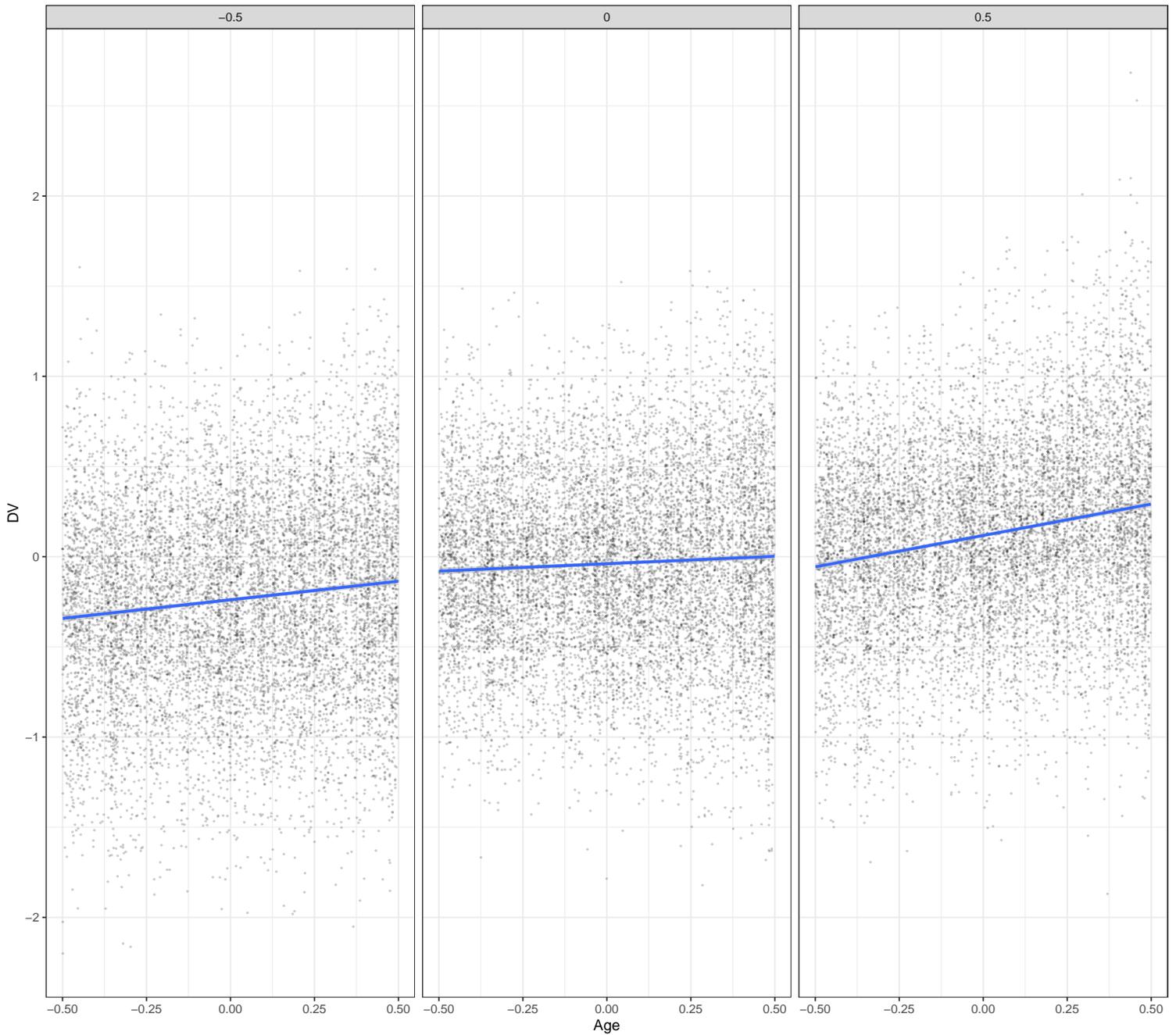
```
plot_age <- plot_age + theme(plot.title = element_text(hjust = 0.5,
  size = 20))
plot_age
```



## 2.4 Age\*Familiarization:

```
plot_age_familiarization <- dat_sim %>%  
  ggplot() + geom_point(aes(y = DV, x = X_a), position = "jitter",  
    alpha = 0.2, size = 0.2) + geom_smooth(method = "lm", formula = y ~  
    x, se = TRUE, aes(y = DV, x = X_a)) + facet_wrap(~X_f) +  
  ggtitle("Age x Familiarization Interaction") + xlab("Age") +  
  theme_bw()  
plot_age_familiarization <- plot_age_familiarization + theme(plot.title = element_text(hjust = 0.5,  
  size = 20))  
plot_age_familiarization
```

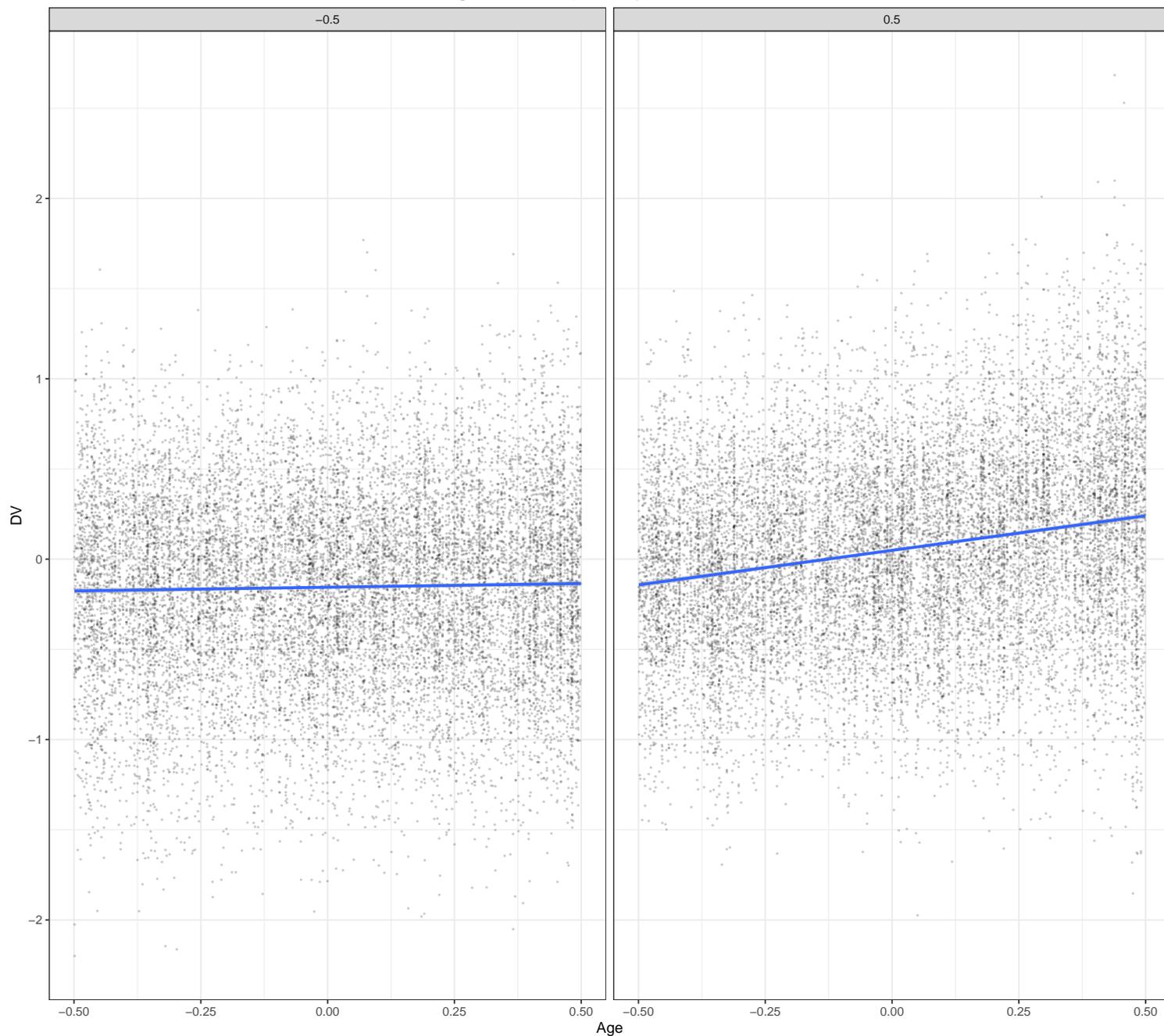
## Age x Familiarization Interaction



### 2.5 Age\*Complexity:

```
plot_age_complexity <- dat_sim %>%  
  ggplot() + geom_point(aes(y = DV, x = X_a), position = "jitter",  
    alpha = 0.2, size = 0.2) + geom_smooth(method = "lm", formula = y ~  
    x, se = TRUE, aes(y = DV, x = X_a)) + facet_wrap(~X_c) +  
  ggtitle("Age x Complexity Interaction") + xlab("Age") + theme_bw()  
  
plot_age_complexity <- plot_age_complexity + theme(plot.title = element_text(hjust = 0.5,  
  size = 20))  
plot_age_complexity
```

## Age x Complexity Interaction



## 2.6 Familiarization\*Complexity:

```
dat_f_c_interaction <- dat_sim %>%  
  mutate(X_c = as.factor(X_c)) %>%  
  mutate(X_f = as.factor(X_f)) %>%  
  group_by(X_f, X_c) %>%  
  dplyr::summarise(med_DV = median(DV))
```

```
## `summarise()` has grouped output by 'X_f'. You can override using the `.groups`  
## argument.
```

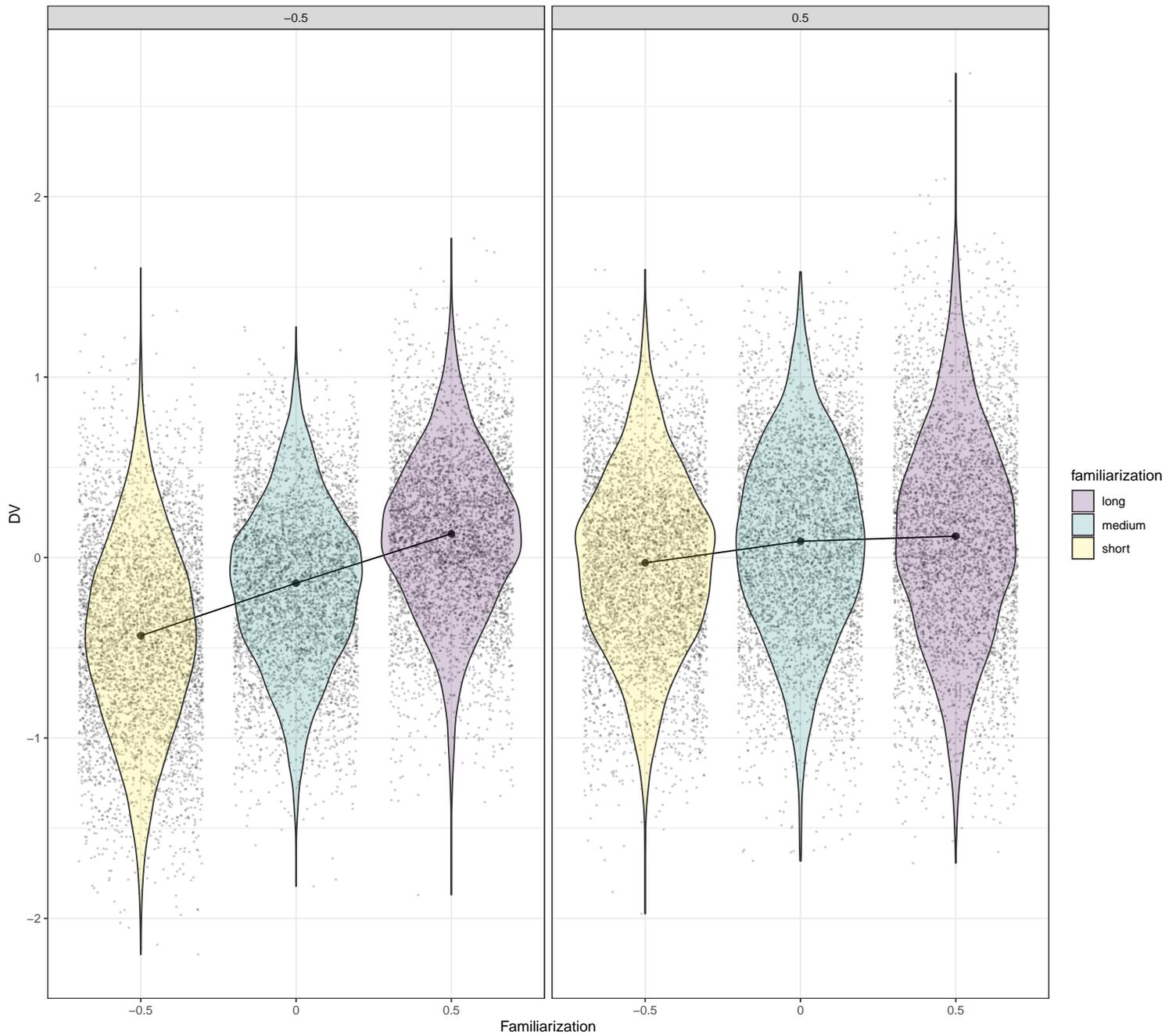
```
plot_familiarization_complexity <- dat_sim %>%  
  mutate(X_c = as.factor(X_c)) %>%  
  mutate(X_f = as.factor(X_f)) %>%  
  ggplot() + geom_point(aes(y = DV, x = X_f), position = "jitter",
```

```

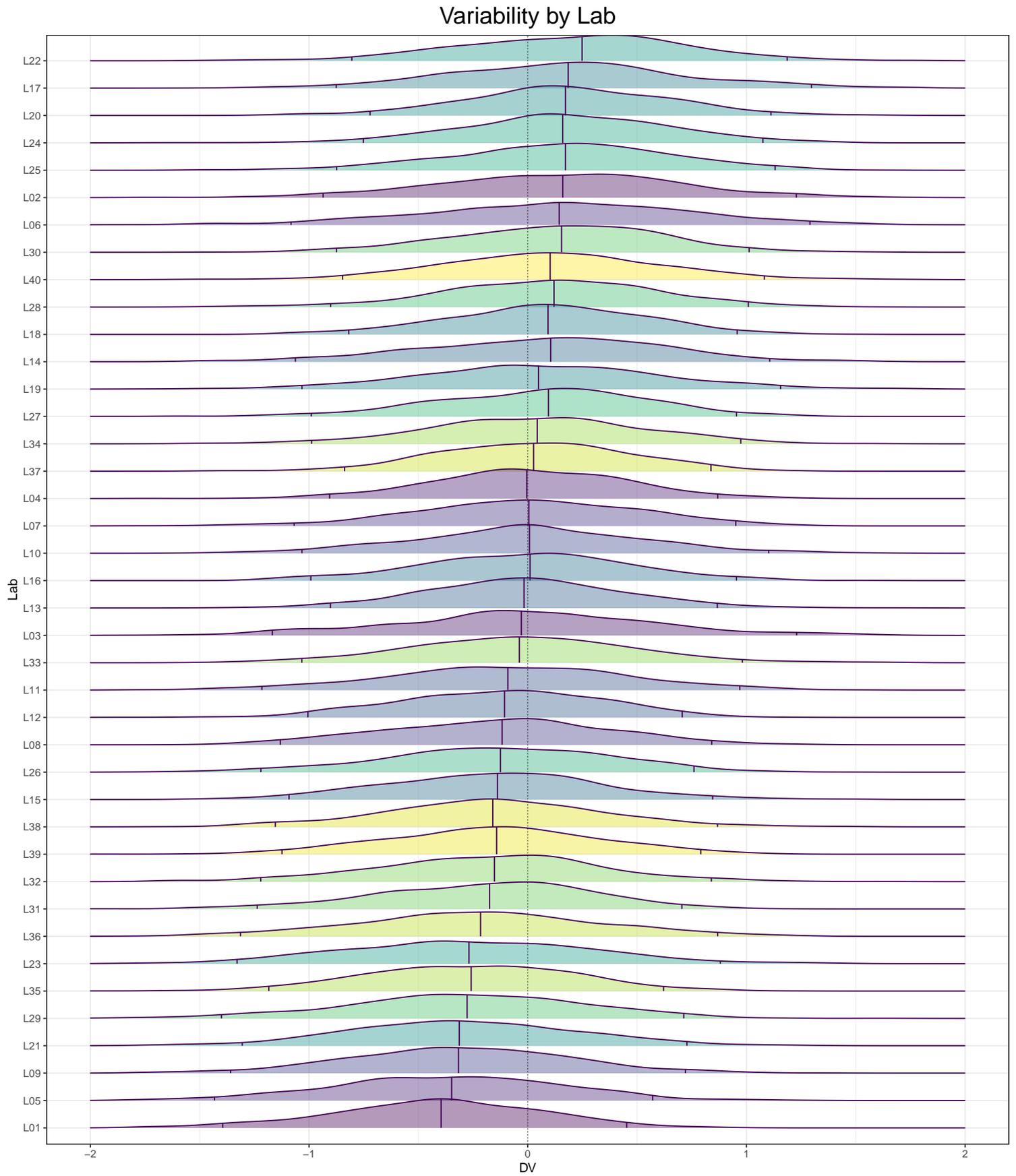
alpha = 0.2, size = 0.2) + geom_point(aes(y = med_DV, x = as.factor(X_f)),
alpha = 0.8, size = 2, data = dat_f_c_interaction) + geom_line(aes(y = med_DV,
x = as.factor(X_f), group = 1), data = dat_f_c_interaction) +
geom_violin(aes(y = DV, x = X_f, fill = familiarization),
alpha = 0.2) + scale_fill_manual(values = viridis(n = 3)) +
facet_wrap(~X_c) + ggtitle("Familiarization x Complexity Interaction") +
xlab("Familiarization") + theme_bw()
plot_familiarization_complexity <- plot_familiarization_complexity +
theme(plot.title = element_text(hjust = 0.5, size = 20))
plot_familiarization_complexity

```

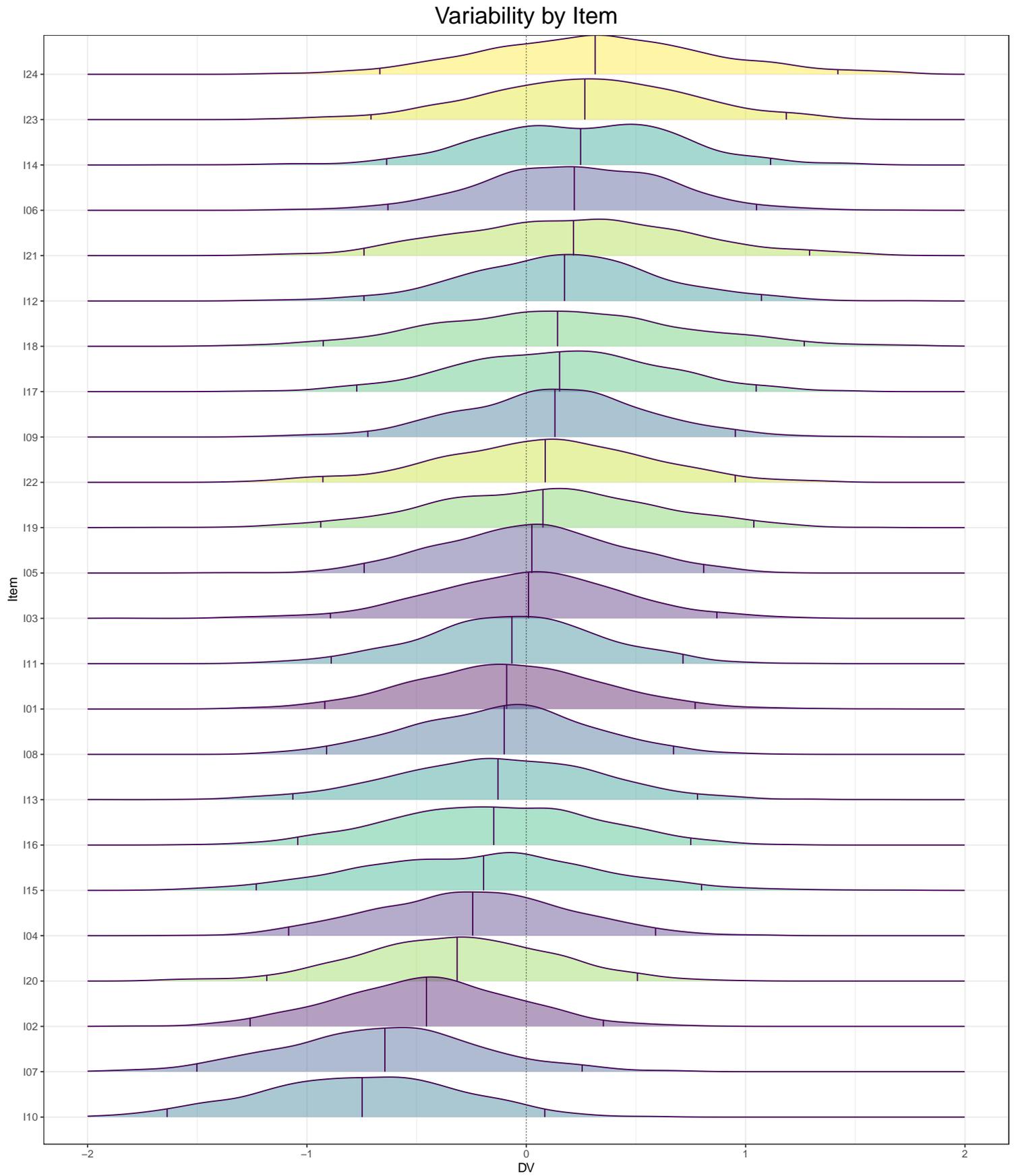
Familiarization x Complexity Interaction



## 2.7 Variability by Lab



## 2.8 Variability by Item



## 3 Power Calculation with Full Data and Varying Intercepts and Varying Slopes

### 3.1 Effect Size = 0.5

```
# Number of simulations:
reps <- 350

# Simulation function:
run_sims <- function(filename_full, ef) {

  dat_sim <- my_sim_data(beta_c = ef,
                        beta_f = ef,
                        beta_a = ef,

                        beta_ca = ef,
                        beta_af = ef,
                        beta_cf = ef,

                        beta_cfa = ef)

  mod_sim <- lmer(DV ~ 1 + X_a * X_c * X_f +
                 (1 + X_c * X_f | subj_id) +
                 (1 | lab_id) +
                 (1 | item_id),
                 data=dat_sim)

  sim_results <- broom.mixed::tidy(mod_sim)

  # append the results to a file
  append <- file.exists(filename_full)
  write_csv(sim_results, filename_full, append = append)

  # return the tidy table
  sim_results
}

filename_full_0.5 = 'run_sims_full_0.5.csv'
start_time <- Sys.time()
sims <- purrr::map_df(1:reps, ~run_sims(filename_full = filename_full_0.5, ef = 0.5))
end_time <- Sys.time()
end_time - start_time
```

### 3.2 Effect Size = 0.4

```
filename_full_0.4 = "run_sims_full_0.4.csv"
start_time <- Sys.time()
sims <- purrr::map_df(1:reps, ~run_sims(filename_full = filename_full_0.4,
                                       ef = 0.4))
end_time <- Sys.time()
end_time - start_time
```

### 3.3 Effect Size = 0.3

```
filename_full_0.3 = "run_sims_full_0.3.csv"
start_time <- Sys.time()
sims <- purrr::map_df(1:reps, ~run_sims(filename_full = filename_full_0.3,
```

```
    ef = 0.3))
end_time <- Sys.time()
end_time - start_time
```

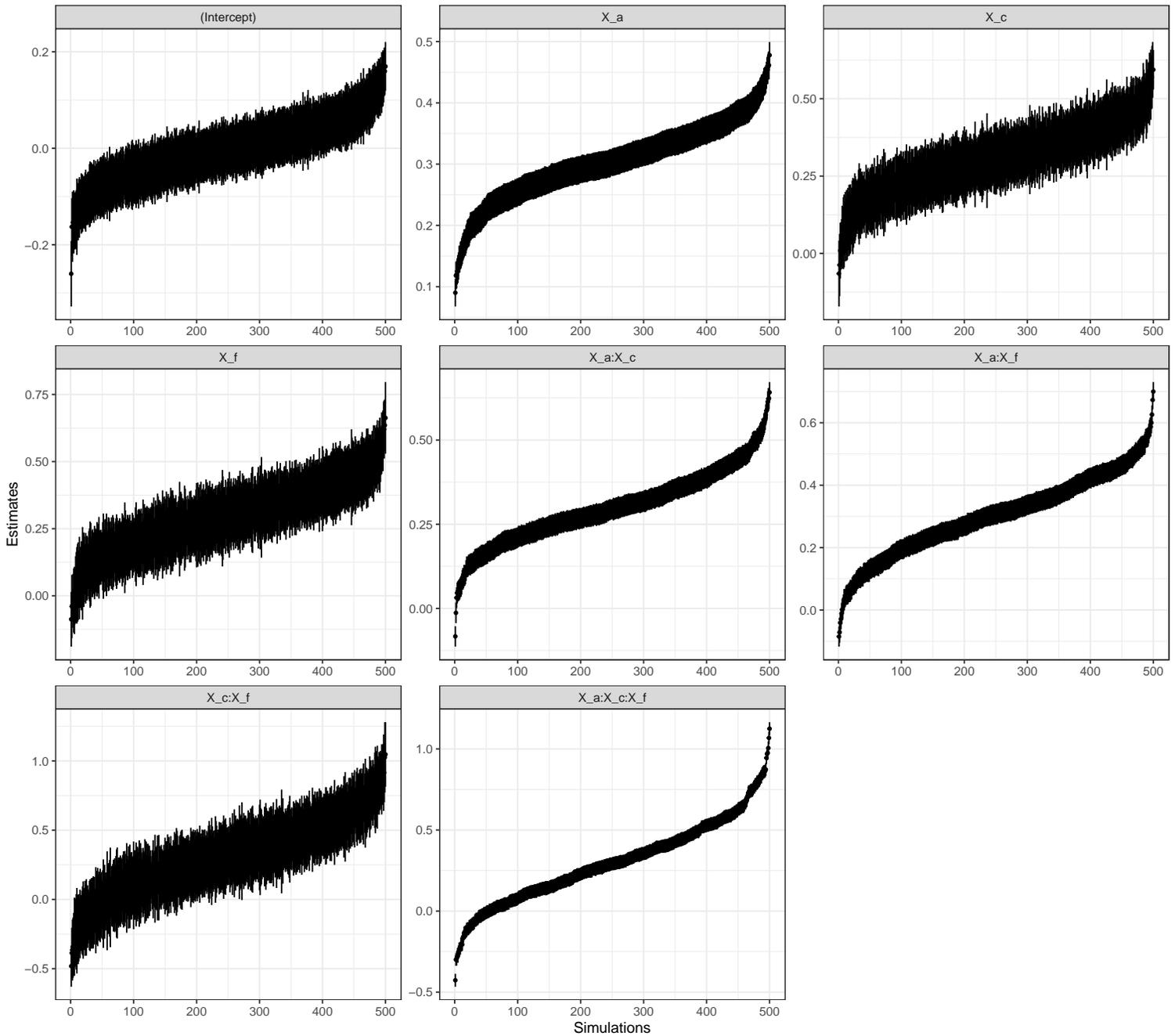
### 3.3.1 Visualise Estimates for Fixed Effects:

```
sims_full_0.3 <- read_csv(filename_full_0.3, col_types = cols(group = col_factor(ordered = TRUE),
  term = col_factor(ordered = TRUE)))

fixed_full_plot <- sims_full_0.3 %>%
  filter(effect == "fixed") %>%
  ungroup() %>%
  arrange(term, estimate) %>%
  mutate(row = rep(seq(1:reps), 8)) %>%
  ggplot(aes(x = row, y = estimate, ymin = estimate - std.error,
    ymax = estimate + std.error)) + facet_wrap(~term, scales = "free") +
  geom_pointrange(fatten = 1/2) + ylab("Estimates") + xlab("Simulations") +
  ggtitle("Estimates of Fixed Effects for Full Data and Varying Intercepts and Varying Slopes, ef = 0.3") +
  theme_bw()

fixed_full_plot <- fixed_full_plot + theme(plot.title = element_text(hjust = 0.5,
  size = 20))
fixed_full_plot
```

## Estimates of Fixed Effects for Full Data and Varying Intercepts and Varying Slopes, $ef = 0.3$

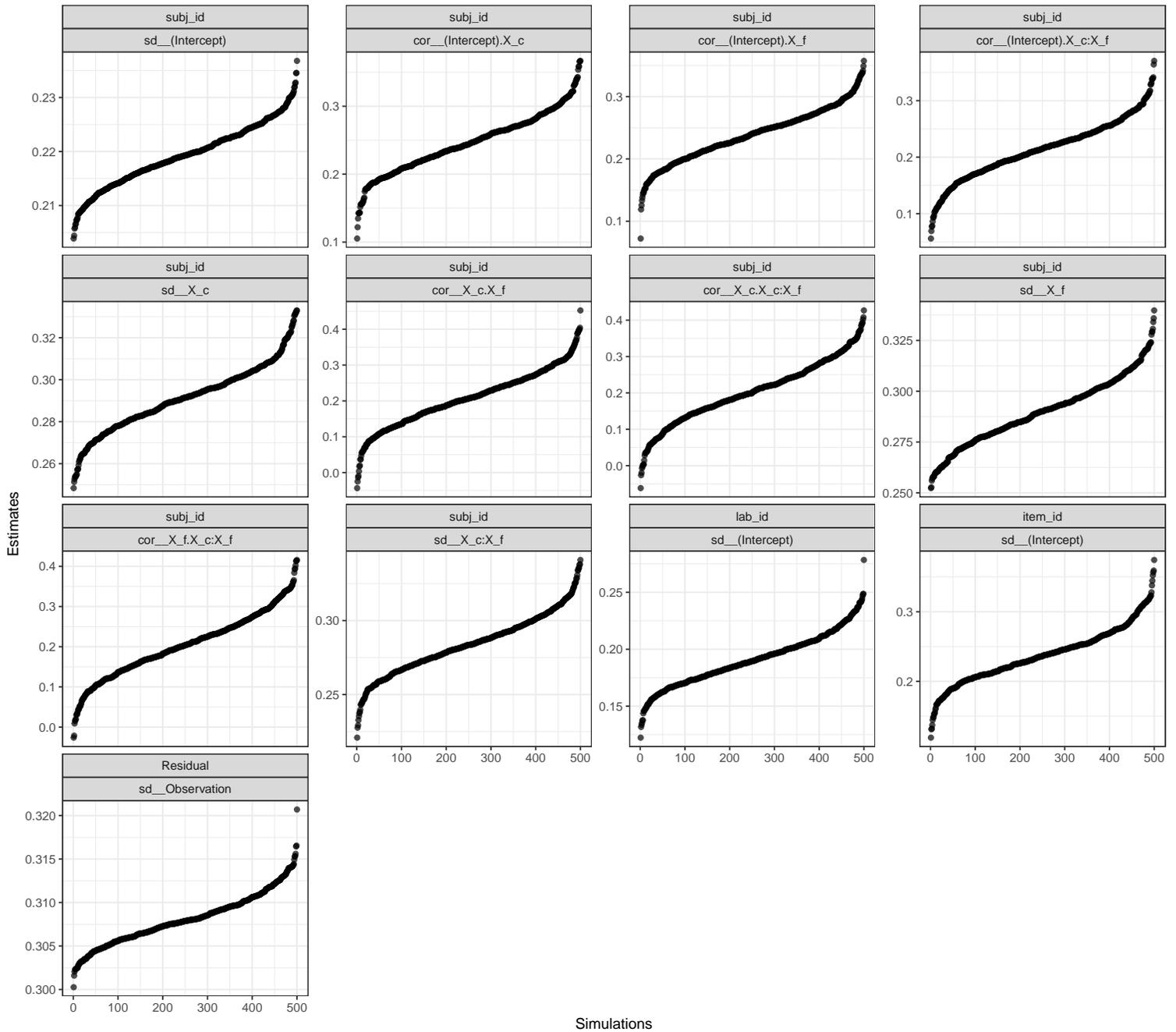


### 3.3.2 Visualise Estimates for Random Effects:

```

ran_full_plot <- sims_full_0.3 %>%
  filter(effect == "ran_pars") %>%
  ungroup() %>%
  arrange(group, term, estimate) %>%
  mutate(row = rep(seq(1:reps), 13)) %>%
  ggplot(aes(x = row, y = estimate)) + geom_point(alpha = 0.7) +
  facet_wrap(~group + term, scales = "free_y") + theme_bw() +
  ylab("Estimates") + xlab("Simulations") + ggtitle("Estimates of Random Effects Full Random-Effects Structure,
  theme_bw()
ran_full_plot <- ran_full_plot + theme(plot.title = element_text(hjust = 0.5,
  size = 20))
ran_full_plot
  
```

## Estimates of Random Effects Full Random-Effects Structure, ef = 0.3



### 3.4 Effect Size = 0.2

```
filename_full_0.2 = "run_sims_full_0.2.csv"  
start_time <- Sys.time()  
sims <- purrr::map_df(1:reps, ~run_sims(filename_full = filename_full_0.2,  
  ef = 0.2))  
end_time <- Sys.time()  
end_time - start_time
```

### 3.5 Effect Size = 0.1

```
filename_full_0.1 = "run_sims_full_0.1.csv"  
start_time <- Sys.time()  
sims <- purrr::map_df(1:reps, ~run_sims(filename_full = filename_full_0.1,
```

```

    ef = 0.1))
end_time <- Sys.time()
end_time - start_time

```

## 4 Power Calculation with Full Data and Varying Intercepts

### 4.1 Effect Size = 0.5

```

# Simulation function:
run_sims <- function(filename_full, ef) {

  dat_sim <- my_sim_data(beta_c = ef,
                        beta_f = ef,
                        beta_a = ef,

                        beta_ca = ef,
                        beta_af = ef,
                        beta_cf = ef,

                        beta_cfa = ef)

  mod_sim <- lmer(DV ~ 1 + X_a * X_c * X_f +
                 (1 | subj_id) +
                 (1 | lab_id) +
                 (1 | item_id),
                 data=dat_sim)

  sim_results <- broom.mixed::tidy(mod_sim)

  # append the results to a file
  append <- file.exists(filename_full)
  write_csv(sim_results, filename_full, append = append)

  # return the tidy table
  sim_results
}

filename_full_int_0.5 = 'run_sims_full_int_0.5.csv'
start_time <- Sys.time()
sims <- purrr::map_df(1:reps, ~run_sims(filename_full = filename_full_int_0.5, ef = 0.5))
end_time <- Sys.time()
end_time - start_time

```

### 4.2 Effect Size = 0.4

```

filename_full_int_0.4 = "run_sims_full_int_0.4.csv"
start_time <- Sys.time()
sims <- purrr::map_df(1:reps, ~run_sims(filename_full = filename_full_int_0.4,
    ef = 0.4))
end_time <- Sys.time()
end_time - start_time

```

### 4.3 Effect Size = 0.3

```

filename_full_int_0.3 = "run_sims_full_int_0.3.csv"
start_time <- Sys.time()
sims <- purrr::map_df(1:reps, ~run_sims(filename_full = filename_full_int_0.3,
  ef = 0.3))
end_time <- Sys.time()
end_time - start_time

```

#### 4.3.1 Visualise Estimates for Fixed Effects:

```

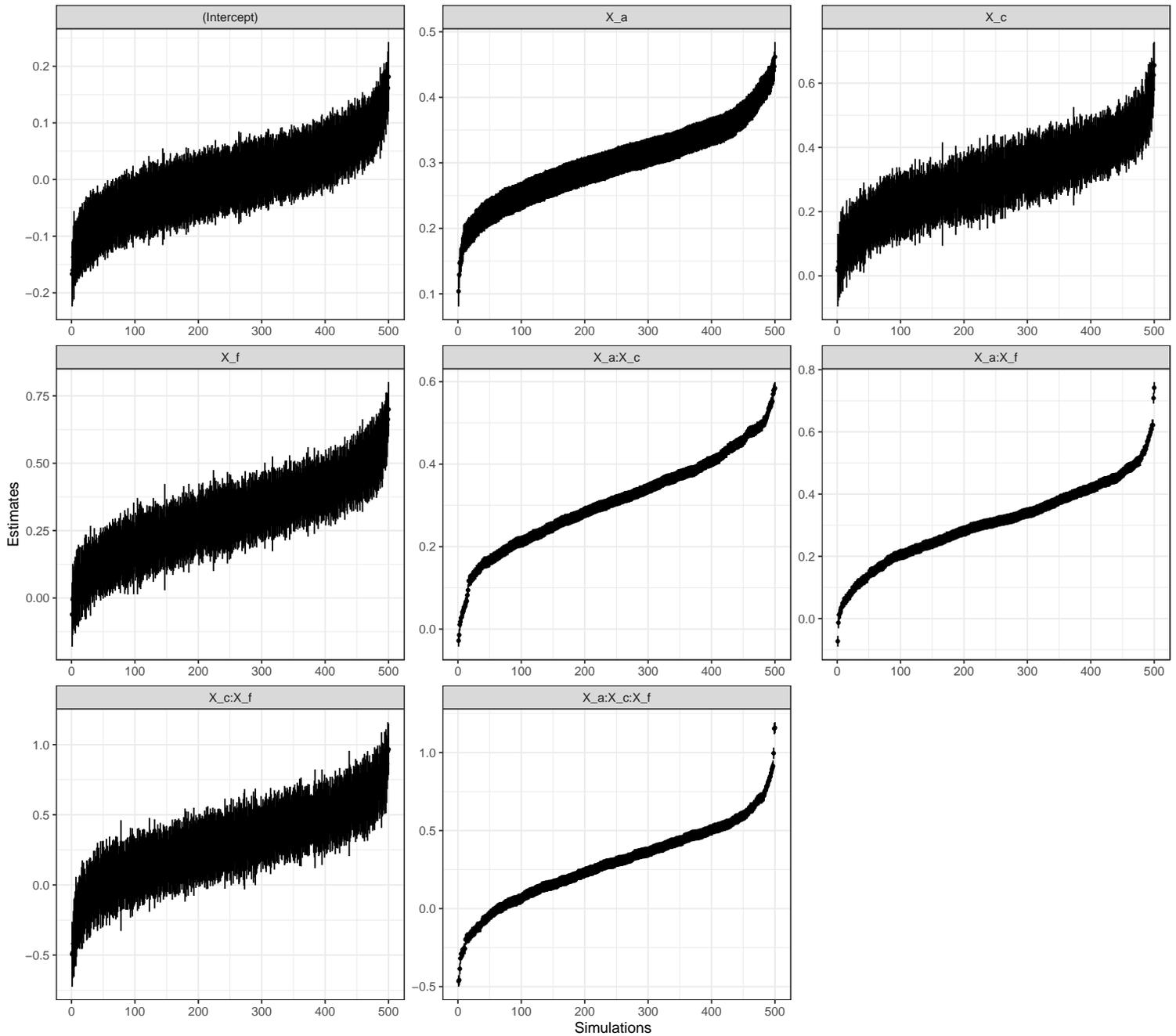
sims_full_int_0.3 <- read_csv(filename_full_int_0.3, col_types = cols(group = col_factor(ordered = TRUE),
  term = col_factor(ordered = TRUE)))

fixed_full_int_plot <- sims_full_int_0.3 %>%
  filter(effect == "fixed") %>%
  ungroup() %>%
  arrange(term, estimate) %>%
  mutate(row = rep(seq(1:reps), 8)) %>%
  ggplot(aes(x = row, y = estimate, ymin = estimate - std.error,
    ymax = estimate + std.error)) + facet_wrap(~term, scales = "free") +
  geom_pointrange(fatten = 1/2) + ylab("Estimates") + xlab("Simulations") +
  ggtitle("Estimates of Fixed Effects for Full Data and Random Intercepts, ef = 0.3") +
  theme_bw()

fixed_full_int_plot <- fixed_full_int_plot + theme(plot.title = element_text(hjust = 0.5,
  size = 20))
fixed_full_int_plot

```

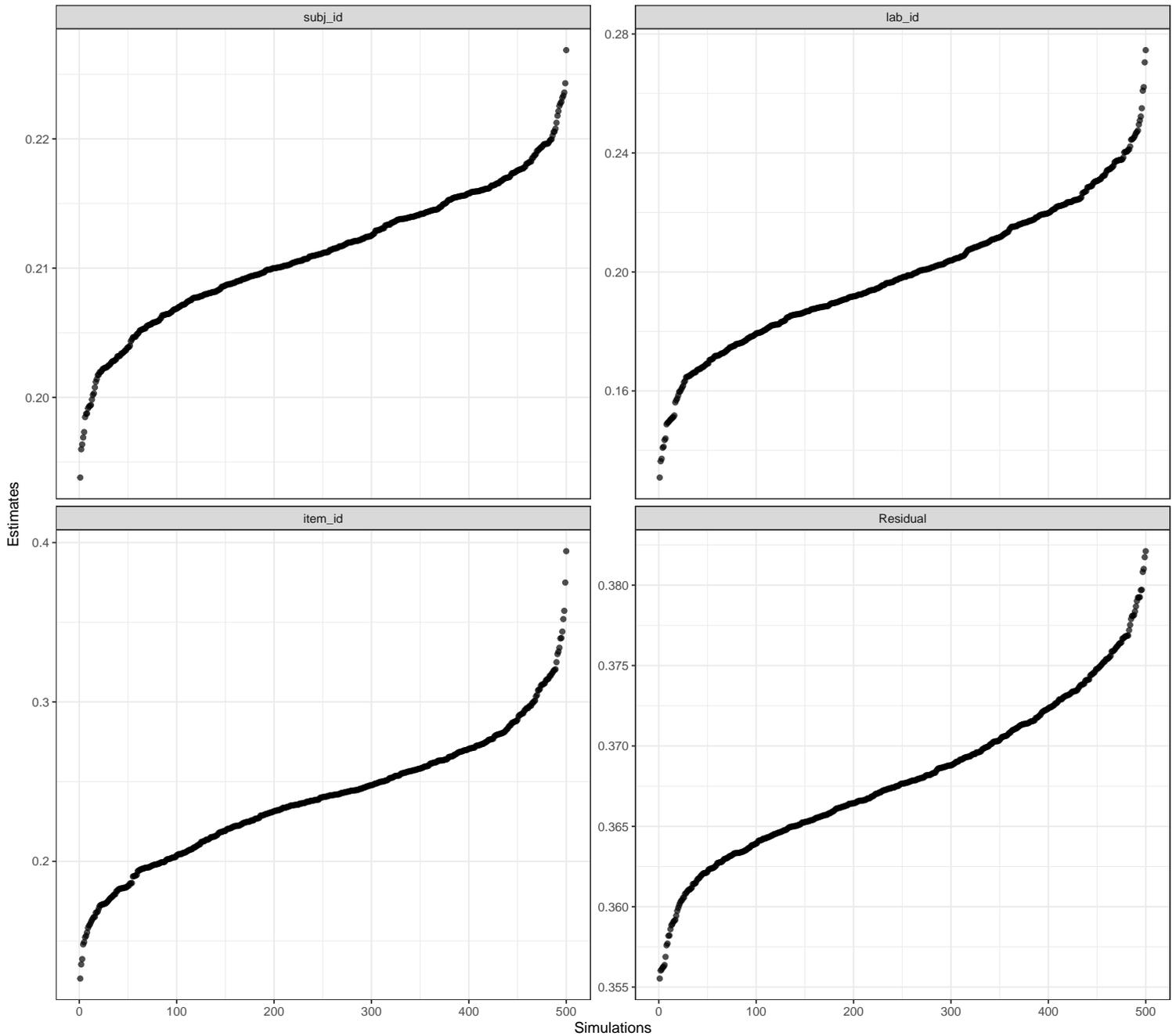
## Estimates of Fixed Effects for Full Data and Random Intercepts, $ef = 0.3$



### 4.3.2 Visualise Estimates for Random Effects:

```
ran_full_int_plot <- sims_full_int_0.3 %>%  
  filter(effect == "ran_pars") %>%  
  ungroup() %>%  
  arrange(group, estimate) %>%  
  mutate(row = rep(seq(1:reps), 4)) %>%  
  ggplot(aes(x = row, y = estimate)) + geom_point(alpha = 0.7) +  
  facet_wrap(~group, scales = "free_y") + theme_bw() + ylab("Estimates") +  
  xlab("Simulations") + ggtitle("Estimates of Random Effects for Full Data, ef = 0.3") +  
  theme_bw()  
ran_full_int_plot <- ran_full_int_plot + theme(plot.title = element_text(hjust = 0.5,  
  size = 20))  
ran_full_int_plot
```

## Estimates of Random Effects for Full Data, ef = 0.3



### 4.4 Effect Size = 0.2

```
filename_full_int_0.2 = "run_sims_full_int_0.2.csv"  
start_time <- Sys.time()  
sims <- purrr::map_df(1:reps, ~run_sims(filename_full = filename_full_int_0.2,  
  ef = 0.2))  
end_time <- Sys.time()  
end_time - start_time
```

### 4.5 Effect Size = 0.1

```
filename_full_int_0.1 = "run_sims_full_int_0.1.csv"  
start_time <- Sys.time()  
sims <- purrr::map_df(1:reps, ~run_sims(filename_full = filename_full_int_0.1,
```

```

    ef = 0.1))
end_time <- Sys.time()
end_time - start_time

```

## 5 Power Calculation with 20 pct. Missing Data and Varying Intercepts and Varying Slopes

### 5.1 Effect Size = 0.5

```

run_sims_missing <- function(filename_missing, ef) {

  dat_sim <- my_sim_data(beta_c = ef,
                        beta_f = ef,
                        beta_a = ef,

                        beta_ca = ef,
                        beta_af = ef,
                        beta_cf = ef,

                        beta_cfa = ef)

  missing_samples <- dat_sim %>%
    mutate(nas = rbinom(nrow(dat_sim), 1, 1 - .20)) %>%
    mutate(DV = ifelse(nas == 1, DV, NA)) %>%
    drop_na()

  mod_sim <- lmer(DV ~ 1 + X_a * X_c * X_f +
                 (1 + X_c * X_f | subj_id) +
                 (1 | lab_id) +
                 (1 | item_id),
                 data=missing_samples)

  sim_results <- broom.mixed::tidy(mod_sim)

  # append the results to a file
  append <- file.exists(filename_missing)
  write_csv(sim_results, filename_missing, append = append)

  # return the tidy table
  sim_results
}

filename_20_missing_0.5 = 'run_sims_20_missing_0.5.csv'
start_time <- Sys.time()
sims_missing <- purrr::map_df(1:reps, ~run_sims_missing(filename_missing = filename_20_missing_0.5, ef = 0.5))
end_time <- Sys.time()
end_time - start_time

```

### 5.2 Effect Size = 0.4

```

filename_20_missing_0.4 = "run_sims_20_missing_0.4.csv"
start_time <- Sys.time()
sims_missing <- purrr::map_df(1:reps, ~run_sims_missing(filename_missing = filename_20_missing_0.4,
                                                       ef = 0.4))
end_time <- Sys.time()
end_time - start_time

```

## 5.3 Effect Size = 0.3

```
filename_20_missing_0.3 = "run_sims_20_missing_0.3.csv"
start_time <- Sys.time()
sims_missing <- purrr::map_df(1:reps, ~run_sims_missing(filename_missing = filename_20_missing_0.3,
  ef = 0.3))
end_time <- Sys.time()
end_time - start_time
```

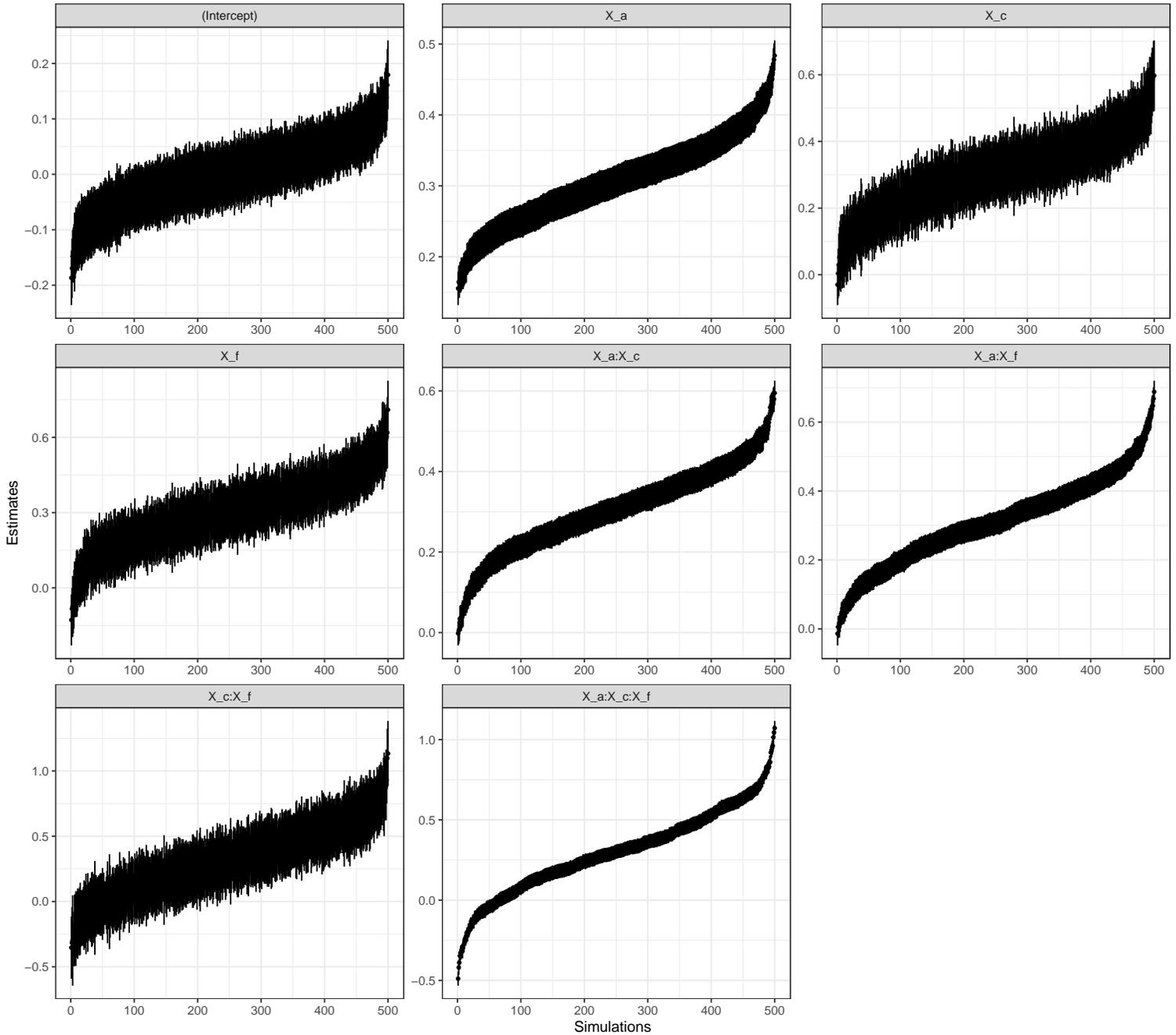
### 5.3.1 Visualise Estimates for Fixed Effects:

```
# read saved simulation data
sims_20_missing_0.3 <- read_csv(filename_20_missing_0.3, col_types = cols(
  # makes sure plots display in this order
  group = col_factor(ordered = TRUE),
  term = col_factor(ordered = TRUE)
))

fixed_missing_plot <- sims_20_missing_0.3 %>%
  filter(effect == "fixed") %>%
  ungroup() %>%
  arrange(term, estimate) %>%
  mutate(row = rep(seq(1:reps), 8)) %>%
  ggplot(aes(x = row, y = estimate, ymin = estimate-std.error, ymax = estimate+std.error)) +
  facet_wrap(~term, scales = "free") +
  geom_pointrange(fatten = 1/2) +
  ylab("Estimates") +
  xlab("Simulations") +
  ggtitle('Estimates of Fixed Effects for 20 pct. Missing Data, ef = 0.3') +
  theme_bw()

fixed_missing_plot <- fixed_missing_plot + theme(plot.title = element_text(hjust = 0.5, size=20))
fixed_missing_plot
```

## Estimates of Fixed Effects for 20 pct. Missing Data, ef = 0.3



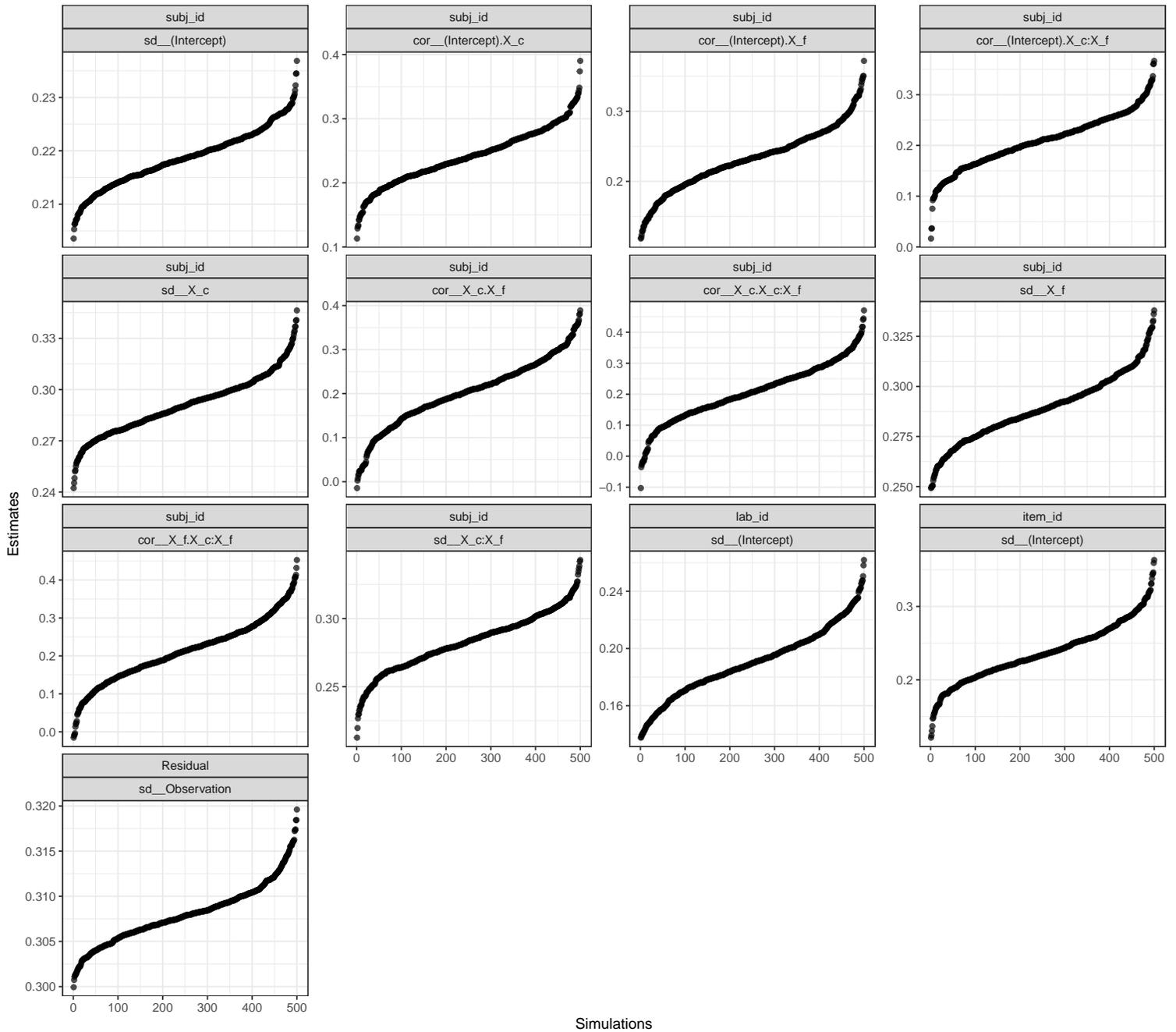
### 5.3.2 Visualise Estimates for Random Effects:

```

ran_missing_plot <- sims_20_missing_0.3 %>%
  filter(effect == "ran_pars") %>%
  ungroup() %>%
  arrange(group, term, estimate) %>%
  mutate(row = rep(seq(1:reps), 13)) %>%
  ggplot(aes(x = row, y = estimate)) + geom_point(alpha = 0.7) +
  facet_wrap(~group + term, scales = "free_y") + theme_bw() +
  ylab("Estimates") + xlab("Simulations") + ggtitle("Estimates of Random Effects for 20 pct. Missing Data, ef = ") +
  theme_bw()
ran_missing_plot <- ran_missing_plot + theme(plot.title = element_text(hjust = 0.5,
  size = 20))
ran_missing_plot

```

## Estimates of Random Effects for 20 pct. Missing Data, ef = 0.3



### 5.4 Effect Size = 0.2

```
filename_20_missing_0.2 = "run_sims_20_missing_0.2.csv"
start_time <- Sys.time()
sims_missing <- purrr::map_df(1:reps, ~run_sims_missing(filename_missing = filename_20_missing_0.2,
  ef = 0.2))
end_time <- Sys.time()
end_time - start_time
```

### 5.5 Effect Size = 0.1

```
filename_20_missing_0.1 = "run_sims_20_missing_0.1.csv"
start_time <- Sys.time()
sims_missing <- purrr::map_df(1:reps, ~run_sims_missing(filename_missing = filename_20_missing_0.1,
```

```

    ef = 0.1))
end_time <- Sys.time()
end_time - start_time

```

## 6 Power Calculation with 50 pct. Missing Data and Varying Intercepts and Varying Slopes

### 6.1 Effect Size = 0.5

```

run_sims_missing <- function(filename_missing, ef) {

  dat_sim <- my_sim_data(beta_c = ef,
                        beta_f = ef,
                        beta_a = ef,

                        beta_ca = ef,
                        beta_af = ef,
                        beta_cf = ef,

                        beta_cfa = ef)

  missing_samples <- dat_sim %>%
    mutate(nas = rbinom(nrow(dat_sim), 1, 1 - .50)) %>%
    mutate(DV = ifelse(nas == 1, DV, NA)) %>%
    drop_na()

  mod_sim <- lmer(DV ~ 1 + X_a * X_c * X_f +
                 (1 + X_c * X_f | subj_id) +
                 (1 | lab_id) +
                 (1 | item_id),
                 data=missing_samples)

  sim_results <- broom.mixed::tidy(mod_sim)

  # append the results to a file
  append <- file.exists(filename_missing)
  write_csv(sim_results, filename_missing, append = append)

  # return the tidy table
  sim_results
}

filename_50_missing_0.5 = 'run_sims_50_missing_0.5.csv'
start_time <- Sys.time()
sims_missing <- purrr::map_df(1:reps, ~run_sims_missing(filename_missing = filename_50_missing_0.5, ef = 0.5))
end_time <- Sys.time()
end_time - start_time

```

### 6.2 Effect Size = 0.4

```

filename_50_missing_0.4 = "run_sims_50_missing_0.4.csv"
start_time <- Sys.time()
sims_missing <- purrr::map_df(1:reps, ~run_sims_missing(filename_missing = filename_50_missing_0.4,
                                                       ef = 0.4))
end_time <- Sys.time()
end_time - start_time

```

## 6.3 Effect Size = 0.3

```
filename_50_missing_0.3 = "run_sims_50_missing_0.3.csv"
start_time <- Sys.time()
sims_missing <- purrr::map_df(1:reps, ~run_sims_missing(filename_missing = filename_50_missing_0.3,
  ef = 0.3))
end_time <- Sys.time()
end_time - start_time
```

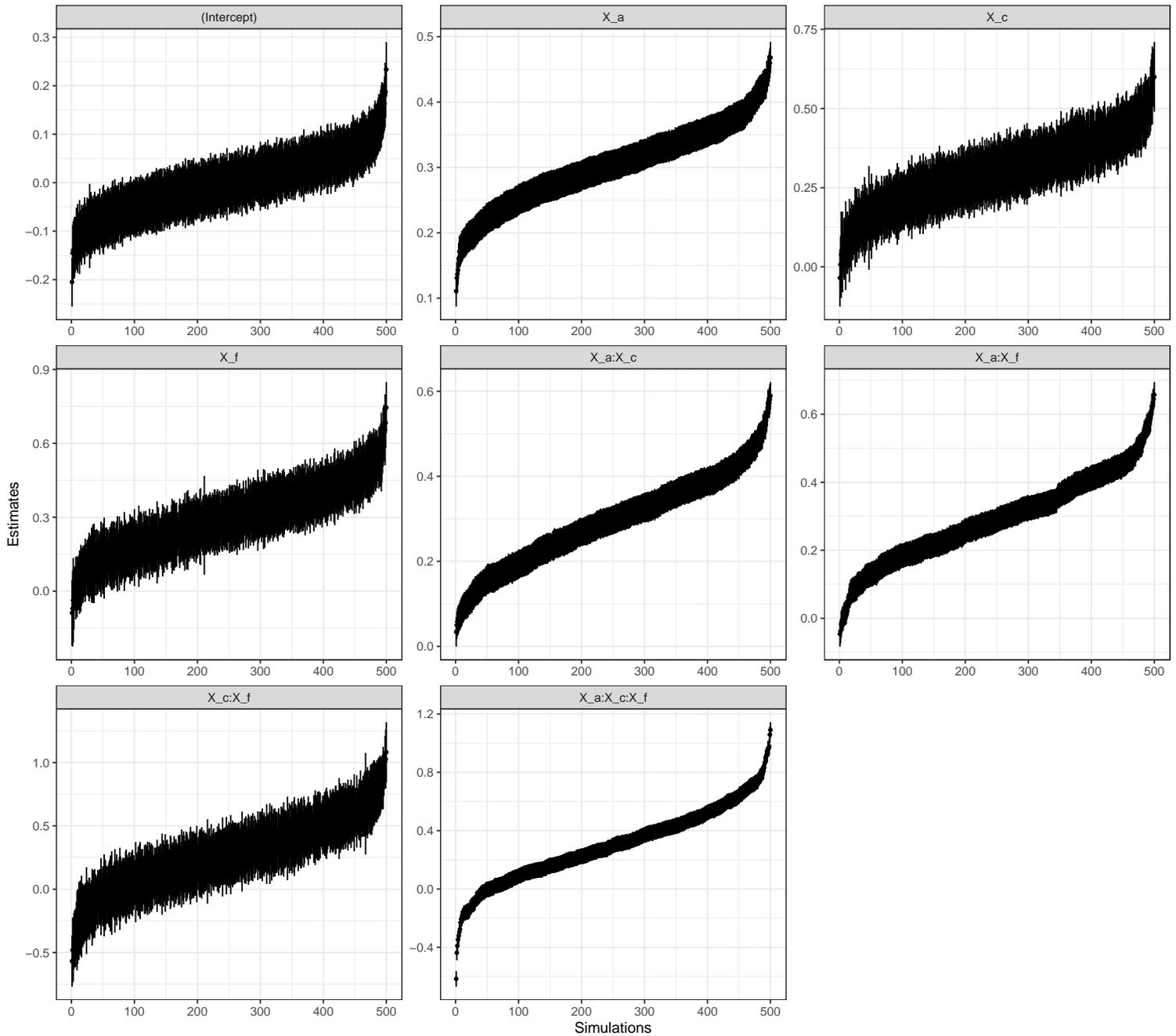
### 6.3.1 Visualise Estimates for Fixed Effects:

```
# read saved simulation data
sims_50_missing_0.3 <- read_csv(filename_50_missing_0.3, col_types = cols(
  # makes sure plots display in this order
  group = col_factor(ordered = TRUE),
  term = col_factor(ordered = TRUE)
))

fixed_missing_plot <- sims_50_missing_0.3 %>%
  filter(effect == "fixed") %>%
  ungroup() %>%
  arrange(term, estimate) %>%
  mutate(row = rep(seq(1:reps), 8)) %>%
  ggplot(aes(x = row, y = estimate, ymin = estimate-std.error, ymax = estimate+std.error)) +
  facet_wrap(~term, scales = "free") +
  geom_pointrange(fatten = 1/2) +
  ylab("Estimates") +
  xlab("Simulations") +
  ggtitle('Estimates of Fixed Effects for 50 pct. Missing Data, ef = 0.3') +
  theme_bw()

fixed_missing_plot <- fixed_missing_plot + theme(plot.title = element_text(hjust = 0.5, size=20))
fixed_missing_plot
```

## Estimates of Fixed Effects for 50 pct. Missing Data, ef = 0.3



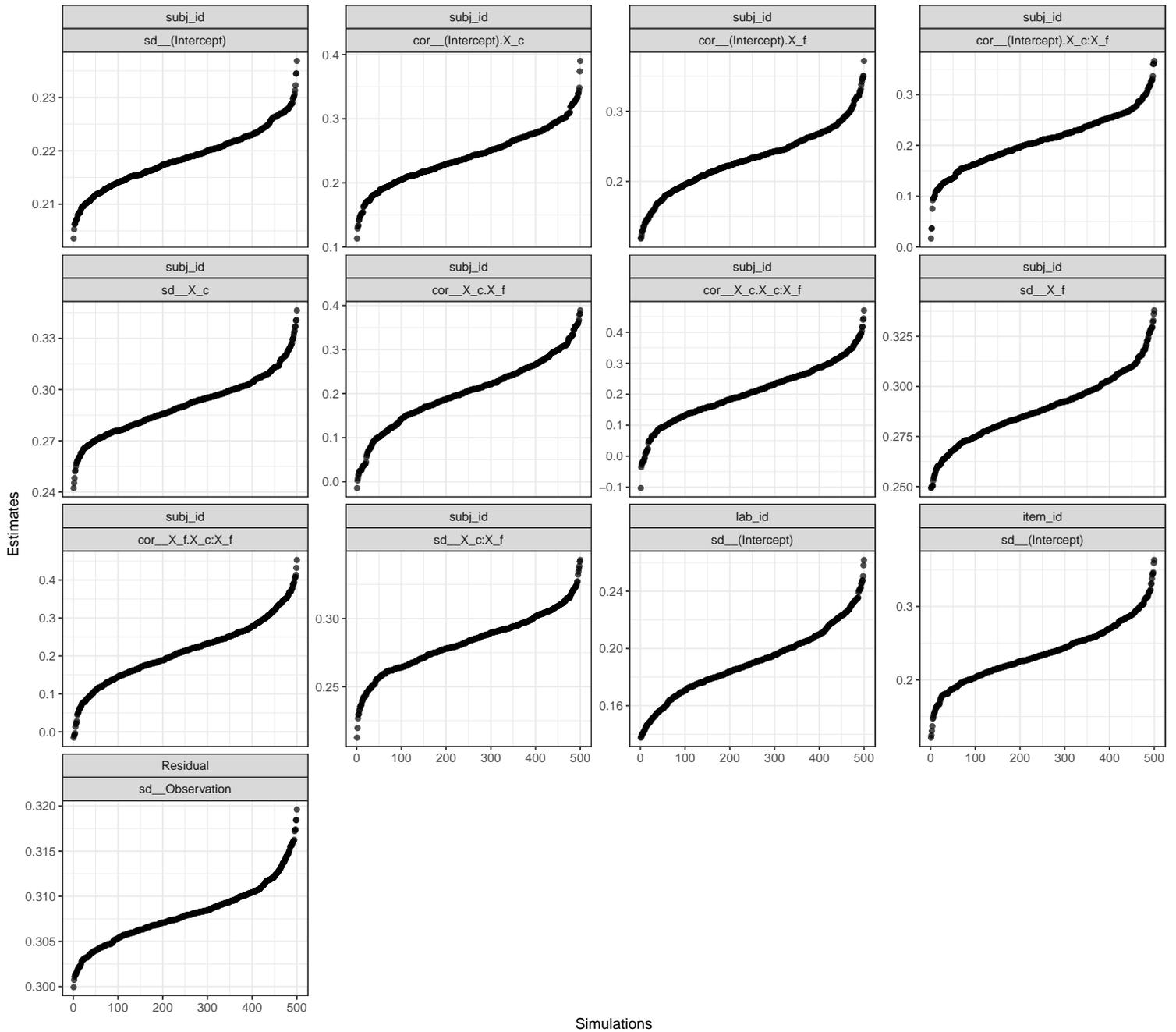
### 6.3.2 Visualise Estimates for Random Effects:

```

ran_missing_plot <- sims_20_missing_0.3 %>%
  filter(effect == "ran_pars") %>%
  ungroup() %>%
  arrange(group, term, estimate) %>%
  mutate(row = rep(seq(1:reps), 13)) %>%
  ggplot(aes(x = row, y = estimate)) + geom_point(alpha = 0.7) +
  facet_wrap(~group + term, scales = "free_y") + theme_bw() +
  ylab("Estimates") + xlab("Simulations") + ggtitle("Estimates of Random Effects for 50 pct. Missing Data, ef = ") +
  theme_bw()
ran_missing_plot <- ran_missing_plot + theme(plot.title = element_text(hjust = 0.5,
  size = 20))
ran_missing_plot

```

## Estimates of Random Effects for 50 pct. Missing Data, ef = 0.3



### 6.4 Effect Size = 0.2

```
filename_50_missing_0.2 = "run_sims_50_missing_0.2.csv"
start_time <- Sys.time()
sims_missing <- purrr::map_df(1:reps, ~run_sims_missing(filename_missing = filename_50_missing_0.2,
  ef = 0.2))
end_time <- Sys.time()
end_time - start_time
```

### 6.5 Effect Size = 0.1

```
filename_50_missing_0.1 = "run_sims_50_missing_0.1.csv"
start_time <- Sys.time()
sims_missing <- purrr::map_df(1:reps, ~run_sims_missing(filename_missing = filename_50_missing_0.1,
```

```

ef = 0.1))
end_time <- Sys.time()
end_time - start_time

```

## 7 Overview of Power Simulation Results

### 7.1 Summary Statistics for Power Calculation with Full Data and Varying Intercepts and Varying Slopes:

Table 1: Power for Simulations with Full Data and Varying Intercepts and Varying Slopes

term	power, ef = 0.1	power, ef = 0.2	power, ef = 0.3	power, ef = 0.4	power, ef = 0.5
(Intercept)	0.050	0.056	0.070	0.056	0.026
X_a	0.844	0.990	1.000	1.000	1.000
X_c	0.170	0.474	0.838	0.966	0.992
X_f	0.118	0.360	0.634	0.864	0.968
X_a:X_c	0.690	0.894	0.986	1.000	1.000
X_a:X_f	0.748	0.864	0.966	0.994	1.000
X_c:X_f	0.072	0.124	0.244	0.350	0.486
X_a:X_c:X_f	0.770	0.808	0.856	0.938	0.956

### 7.2 Summary Statistics for Power Calculation with Full Data and Varying Intercepts:

Table 2: Power for Simulations with Full Data and Varying Intercepts

term	power, ef = 0.1	power, ef = 0.2	power, ef = 0.3	power, ef = 0.4	power, ef = 0.5
(Intercept)	0.050	0.056	0.050	0.080	0.058
X_a	0.822	0.994	1.000	1.000	1.000
X_c	0.162	0.482	0.810	0.998	0.966
X_f	0.136	0.352	0.644	0.970	0.884
X_a:X_c	0.868	0.958	0.990	1.000	0.998
X_a:X_f	0.848	0.932	0.990	1.000	0.998
X_c:X_f	0.092	0.136	0.220	0.564	0.348
X_a:X_c:X_f	0.798	0.832	0.882	0.964	0.934

### 7.3 Summary Statistics for Power Calculation with 20 pct. Missing Data and Varying Intercepts and Varying Slopes:

Table 3: Power for Simulations with 20 pct. Missing Data and Varying Intercepts and Slopes

term	power, ef = 0.1	power, ef = 0.2	power, ef = 0.3	power, ef = 0.4	power, ef = 0.5
(Intercept)	0.040	0.054	0.042	0.064	0.052
X_a	0.832	0.996	1.000	1.000	1.000
X_c	0.162	0.476	0.806	0.950	0.998
X_f	0.122	0.348	0.658	0.842	0.972
X_a:X_c	0.686	0.928	0.982	1.000	1.000
X_a:X_f	0.674	0.872	0.974	0.996	1.000
X_c:X_f	0.062	0.132	0.266	0.376	0.498
X_a:X_c:X_f	0.762	0.800	0.862	0.926	0.950

## 7.4 Summary Statistics for Power Calculation with 50 pct. Missing Data and Varying Intercepts and Varying Slopes:

Table 4: Power for Simulations with 50 pct. Missing Data and Varying Intercepts and Slopes

term	power, ef = 0.1	power, ef = 0.2	power, ef = 0.3	power, ef = 0.4	power, ef = 0.5
(Intercept)	0.036	0.058	0.054	0.044	0.046
X_a	0.822	0.986	1.000	1.000	1.000
X_c	0.168	0.540	0.840	0.970	0.996
X_f	0.120	0.366	0.658	0.836	0.964
X_a:X_c	0.666	0.902	0.992	0.998	1.000
X_a:X_f	0.680	0.826	0.966	0.992	1.000
X_c:X_f	0.066	0.174	0.232	0.372	0.460
X_a:X_c:X_f	0.690	0.754	0.840	0.896	0.942

## 8 Overview of Bias Results

```
sim_stats_full_bias_int_slope %>%
  kbl(caption = "Bias for Simulations with Full Data and Varying Intercepts and Varying Slopes",
      digits = 3, align = "c") %>%
  kable_styling(full_width = T, latex_options = c("striped",
    "HOLD_position"))
```

Table 5: Bias for Simulations with Full Data and Varying Intercepts and Varying Slopes

term	bias, ef = 0.1	bias, ef = 0.2	bias, ef = 0.3	bias, ef = 0.4	bias, ef = 0.5
(Intercept)	0.000	0.003	0.003	-0.003	0.003
X_a	0.000	-0.001	-0.003	0.002	-0.003
X_c	0.001	-0.003	-0.007	0.000	-0.004
X_f	0.012	0.002	0.002	0.014	-0.021
X_a:X_c	-0.001	0.004	0.005	0.006	0.006
X_a:X_f	-0.007	0.013	-0.007	0.000	-0.002
X_c:X_f	0.024	-0.001	-0.015	0.001	0.011
X_a:X_c:X_f	0.009	0.003	0.008	0.018	-0.018

```
sim_stats_full_bias_int %>%
  kbl(caption = "Bias for Simulations with Full Data and Varying Intercepts",
      digits = 3, align = "c") %>%
  kable_styling(full_width = T, latex_options = c("striped",
    "HOLD_position"))
```

Table 6: Bias for Simulations with Full Data and Varying Intercepts

term	bias, ef = 0.1	bias, ef = 0.2	bias, ef = 0.3	bias, ef = 0.4	bias, ef = 0.5
(Intercept)	0.002	0.006	0.001	0.003	-0.003
X_a	0.001	0.002	-0.002	-0.098	0.094
X_c	0.007	0.000	-0.003	-0.103	0.095
X_f	0.004	0.012	-0.001	-0.094	0.094
X_a:X_c	-0.011	0.008	-0.013	-0.105	0.105
X_a:X_f	-0.003	0.006	-0.011	-0.105	0.093
X_c:X_f	-0.005	0.014	-0.005	-0.136	0.101
X_a:X_c:X_f	0.004	0.027	-0.003	-0.075	0.113

```
sim_stats_20_missing_bias %>%
  kbl(caption = "Bias for Simulations with 20 pct. Missing Data and Varying Intercepts and Slopes",
      digits = 3, align = "c") %>%
```

```
kable_styling(full_width = T, latex_options = c("striped",
"HOLD_position"))
```

Table 7: Bias for Simulations with 20 pct. Missing Data and Varying Intercepts and Slopes

term	bias, ef = 0.1	bias, ef = 0.2	bias, ef = 0.3	bias, ef = 0.4	bias, ef = 0.5
(Intercept)	0.001	0.002	0.002	-0.005	0.001
X_a	0.010	0.000	-0.006	0.006	0.001
X_c	0.002	-0.003	-0.011	0.007	0.000
X_f	0.000	0.001	-0.006	0.005	-0.002
X_a:X_c	-0.001	-0.009	-0.005	0.004	-0.014
X_a:X_f	0.014	0.005	-0.007	0.001	0.000
X_c:X_f	0.008	0.002	-0.008	0.011	-0.008
X_a:X_c:X_f	0.009	-0.014	-0.003	0.003	0.020

```
sim_stats_50_missing_bias %>%
  kbl(caption = "Bias for Simulations with 50 pct. Missing Data and Varying Intercepts and Slopes",
      digits = 3, align = "c") %>%
  kable_styling(full_width = T, latex_options = c("striped",
"HOLD_position"))
```

Table 8: Bias for Simulations with 50 pct. Missing Data and Varying Intercepts and Slopes

term	bias, ef = 0.1	bias, ef = 0.2	bias, ef = 0.3	bias, ef = 0.4	bias, ef = 0.5
(Intercept)	-0.006	-0.003	0.002	0.000	-0.003
X_a	-0.002	0.002	-0.004	0.004	0.000
X_c	0.006	-0.011	0.000	-0.005	0.000
X_f	-0.003	-0.008	0.007	0.016	0.005
X_a:X_c	0.000	-0.002	0.002	0.004	-0.007
X_a:X_f	0.004	0.004	0.010	0.012	0.007
X_c:X_f	0.010	-0.031	0.029	-0.012	0.014
X_a:X_c:X_f	0.000	0.016	0.000	0.020	-0.009

## 9 Increasing Variance with Trial Number

### 9.1 Extracting estimates of increasing variance from the adult pilot data

```
# download data from adult pilot study:
adult_data <- read.csv("02_processed_lt_data.csv")

stimulus_info_df <- adult_data %>%
  filter(phase == "pref") %>%
  mutate(familiar_item = case_when(left_stimulus == "familiar" ~
    stimulus_processed_left, right_stimulus == "familiar" ~
    stimulus_processed_right), novel_item = case_when(left_stimulus ==
    "novel" ~ stimulus_processed_left, right_stimulus ==
    "novel" ~ stimulus_processed_right)) %>%
  select(subject, block_number, familiar_item, novel_item) %>%
  distinct(subject, block_number, .keep_all = TRUE)

lt_comparison_df <- adult_data %>%
  filter(phase == "pref") %>%
  group_by(subject, exposure_time, block_number, complexity,
    gaze_location_type) %>%
  summarise(sum_dwell_time = sum(dwell_time)) %>%
  pivot_wider(names_from = gaze_location_type, values_from = sum_dwell_time) %>%
  ungroup() %>%
```

```

left_join(stimulus_info_df, by = c("subject", "block_number")) %>%
rename(familiarization_time = exposure_time, stimulus_complexity = complexity,
      trial_number = block_number, familiar_looking_time = familiar,
      novel_looking_time = novel, participant = subject) %>%
# currently excluding all the empty trials
filter(!is.na(familiar_looking_time) & !is.na(novel_looking_time))

```

## `summarise()` has grouped output by 'subject', 'exposure\_time', 'block\_number',  
## 'complexity'. You can override using the `.groups` argument.

```

lt_comparison_df <- lt_comparison_df %>%
  mutate(familiarization_time_scaled_centered = log10(familiarization_time) -
         log10(2000), contrast_coded_complexity = ifelse(stimulus_complexity ==
         "complex", 0.5, -0.5))

proportion_df <- adult_data %>%
  filter(phase == "pref") %>%
  group_by(block_number, exposure_time, complexity, gaze_location_type,
          subject) %>%
  summarise(sum_dwell_time = sum(dwell_time, na.rm = TRUE)) %>%
  pivot_wider(names_from = gaze_location_type, values_from = sum_dwell_time) %>%
  mutate(familiar = ifelse(is.na(familiar), 0, familiar), novel = ifelse(is.na(novel),
  0, novel)) %>%
  mutate(novelty_looking_proportion = novel/(familiar + novel))

```

## `summarise()` has grouped output by 'block\_number', 'exposure\_time',  
## 'complexity', 'gaze\_location\_type'. You can override using the `.groups`  
## argument.

```

dwell_proportion_mean <- mean(proportion_df$novelty_looking_proportion,
  na.rm = T)
dwell_proportion_sd <- sd(proportion_df$novelty_looking_proportion,
  na.rm = T)

adult_data_z <- proportion_df %>%
  mutate(dwell_time_z = (novelty_looking_proportion - dwell_proportion_mean)/dwell_proportion_sd)

data_on_increase_in_sd <- adult_data_z %>%
  group_by(block_number) %>%
  summarise(n = n(), dwell_time_median = median(dwell_time_z,
  na.rm = T), sd_block = sd(dwell_time_z, na.rm = T))

increase_sd_value <- data_on_increase_in_sd$sd_block[12] - data_on_increase_in_sd$sd_block[1]
increase_sd_value

## [1] 0.3098641

```

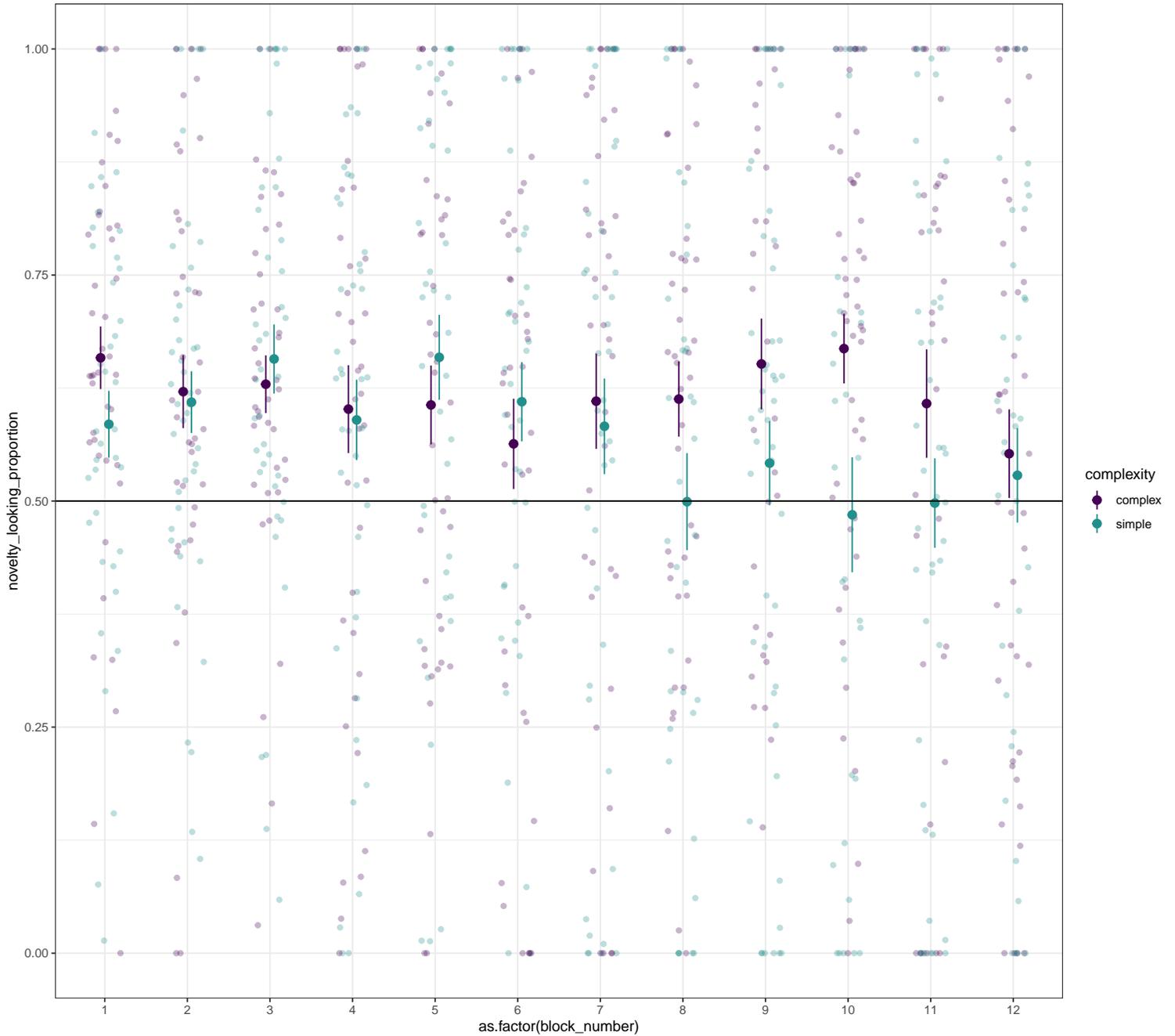
## 9.2 Visualisation of adult data in proportions

```

adult_proportion_plot <- proportion_df %>%
  ggplot(aes(x = as.factor(block_number), y = novelty_looking_proportion,
            color = complexity)) + geom_jitter(alpha = 0.3, width = 0.2) +
  stat_summary(aes(x = as.factor(block_number), y = novelty_looking_proportion,
            color = complexity), position = position_dodge(width = 0.2)) +
  geom_hline(yintercept = 0.5) + scale_color_manual(values = viridis(n = 3)) +
  ggtitle("Adult Pilot Data in Proportions") + theme_bw()
adult_proportion_plot <- adult_proportion_plot + theme(plot.title = element_text(hjust = 0.5,
  size = 20))
adult_proportion_plot

```

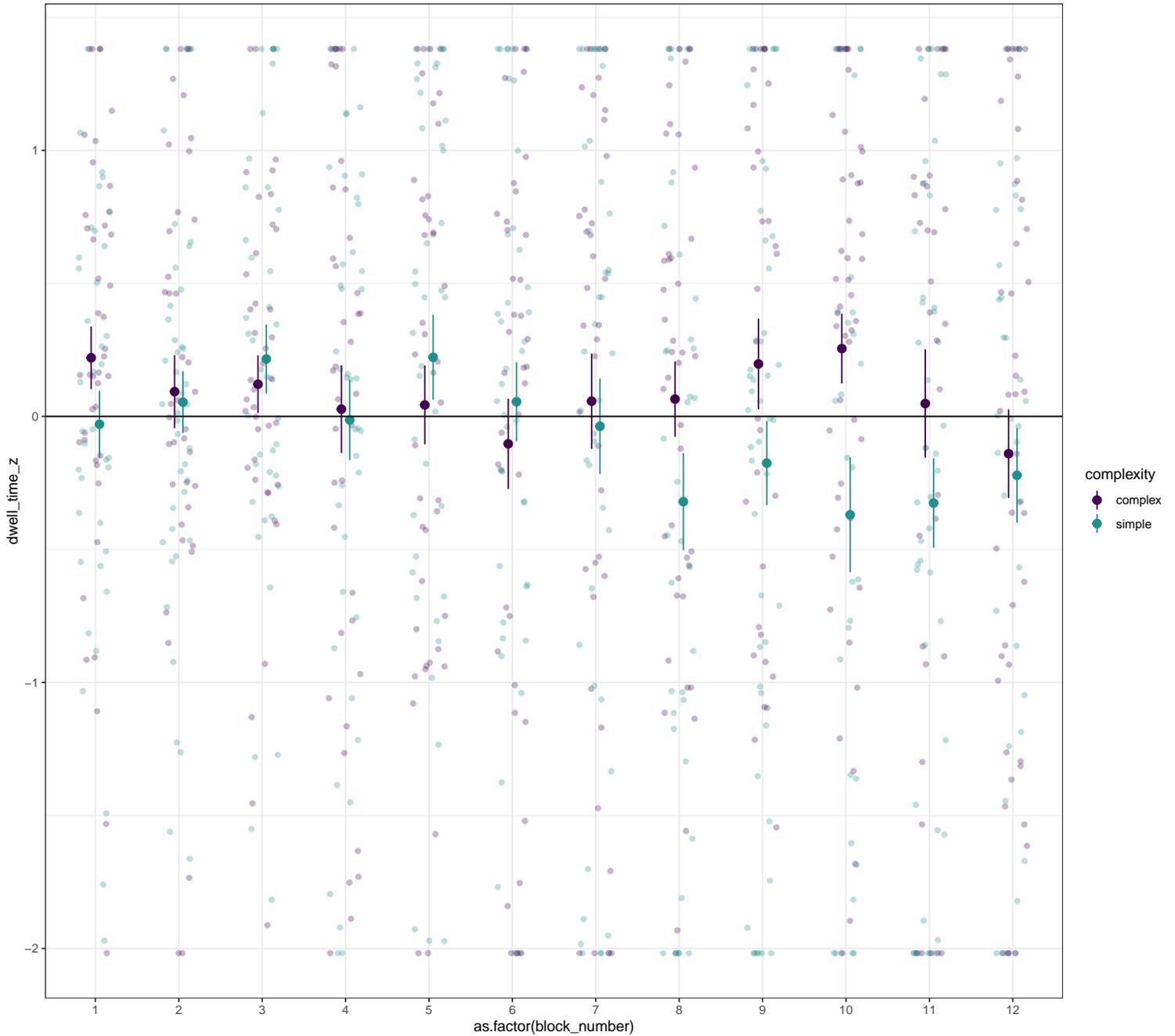
## Adult Pilot Data in Proportions



### 9.3 Visualisation of adult data in z-score

```
adult_data_z_plot <- adult_data_z %>%  
  ggplot(aes(x = as.factor(block_number), y = dwell_time_z,  
            color = complexity)) + geom_jitter(alpha = 0.3, width = 0.2) +  
  stat_summary(aes(x = as.factor(block_number), y = dwell_time_z,  
                  color = complexity), position = position_dodge(width = 0.2)) +  
  geom_hline(yintercept = 0) + scale_color_manual(values = viridis(n = 3)) +  
  ggtitle("Adult Pilot Data in z-scores") + theme_bw()  
  
adult_data_z_plot <- adult_data_z_plot + theme(plot.title = element_text(hjust = 0.5,  
  size = 20))  
adult_data_z_plot
```

## Adult Pilot Data in z-scores



### 9.4 Modification of simulation function to include variance increase

```
my_sim_data_variance <- function(  
  increase_sd = increase_sd_value, #Add variance to increase with trial number  
  
  n_subj      = 1280, # number of subjects  
  n_simple   = 12,   # number of complex stimuli  
  n_complex  = 12,   # number of complex stimuli  
  n_small_fam = 8,   #small familiarization time  
  n_medium_fam = 8, #medium familiarization time  
  n_high_fam  = 8,   #high familiarization time  
  n_lab      = 40,  
  
  beta_0 = 0, # intercept; i.e., the grand mean  
  beta_c = 0.3, # main effect for complexity  
  beta_f = 0.3, # main effect for familiarization time
```

```

beta_a = 0.3, # main effect for age

beta_ca = 0.3,
beta_af = 0.3,
beta_cf = 0.3,

beta_cfa = 0.3, #main effect for interaction between complexity and familiarization.

subject_0 = 0.2, # by-subject random intercept sd

subject_c = 0.2, # by-subject slope complexity sd
subject_f = 0.2, # by-subject slope familiarization sd
subject_a = 0.2, # by-subject slope age sd

subject_ca = 0.2, # by-subject slope for interaction between age and complexity sd
subject_af = 0.2, # by-subject slope for interaction between age and familiarization sd
subject_cf = 0.2, # by-subject slope complexity*familiarization sd

subject_cfa = 0.2, # by-subject slope for interaction between age, complexity and familiarization sd

subj_rho = .2, # correlations between by-subject random effects

lab_0 = 0.2, # by-lab random intercept sd

lab_c = 0.2, # by-lab slope complexity sd
lab_f = 0.2, # by-lab slope familiarization sd
lab_a = 0.2, # by-lab slope age sd

lab_ca = 0.2, # by-lab slope for interaction between age and complexity sd
lab_af = 0.2, # by-lab slope for interaction between age and familiarization sd
lab_cf = 0.2, # by-lab random slope complexity*familiarization sd

lab_cfa = 0.2, # by-lab slope for interaction between age, complexity and familiarization sd

lab_rho = 0.2, # correlations between by-lab random effects

item_0 = 0.2, # by-item random intercept sd

item_c = 0.2, # by-item slope complexity sd
item_f = 0.2, # by-item slope familiarization sd
item_a = 0.2, # by-item slope age sd

item_ca = 0.2, # by-item slope for interaction between age and complexity sd
item_af = 0.2, # by-item slope for interaction between age and familiarization sd
item_cf = 0.2, # by-item random slope complexity*familiarization sd

item_cfa = 0.2, # by-item slope for interaction between age, complexity and familiarization sd

item_rho = 0.2, # correlations between by-item random effects

sigma = 0.3 # residual (error) sd
) { # residual (standard deviation)

# simulate a sample of items
items <- data.frame(
  item_id = seq_len(n_simple + n_complex),
  category = rep(c("simple", "complex"), c(n_simple, n_complex)),
  X_c = rep(c(-0.5, 0.5), c(n_simple, n_complex)),
  familiarization = rep(c("short", "medium", "long"), (n_simple + n_complex)/3),
  X_f = rep(c(-0.5, 0, 0.5), (n_simple + n_complex)/3),
  faux::rnorm_multi(

```

```

n = n_simple + n_complex, mu = 0, sd = c(item_0,
      item_c,
      item_f,
      item_a,
      item_ca,
      item_af,
      item_cf,
      item_cfa), r = item_rho,
varnames = c("I_0", "I_c", "I_f", "I_a",
      "I_ca", "I_af", "I_cf",
      "I_cfa"))
) %>%
mutate(item_id = faux::make_id(nrow(.), "I"))

# simulate a sample of subjects
subjects <-
  faux::rnorm_multi(
    n = n_subj, mu = 0, sd = c(subject_0,
      subject_c,
      subject_f,
      subject_a,
      subject_ca,
      subject_af,
      subject_cf,
      subject_cfa), r = subj_rho,
    varnames = c("S_0", "S_c", "S_f", "S_a",
      "S_ca", "S_af", "S_cf",
      "S_cfa"))
) %>%
  mutate(subj_id = faux::make_id(nrow(.), "S")) %>%
  mutate(X_a = runif(n_subj, min = -0.5, max = 0.5))
#add subject age measure, sample from distribution from -0.5 to 0.5. #subjects$subj_id <- 1:n_subj

labs <- faux::rnorm_multi(
  n = n_lab, mu = 0, sd = c(lab_0, lab_c, lab_f, lab_a,
    lab_ca, lab_af, lab_cf,
    lab_cfa), r = lab_rho,
  varnames = c("L_0", "L_c", "L_f", "L_a",
    "L_ca", "L_af", "L_cf",
    "L_cfa"))
) %>%
  mutate(lab_id = faux::make_id(nrow(.), "L"))

#create lab and subj nesting structure
#Number of subjects must be a multiple of number of labs
lab_multiplier = n_subj/n_lab
lab_subj_dict <- data.frame(
  subj_id = subjects$subj_id,
  lab_id = rep(labs$lab_id, lab_multiplier)
)

# cross subject and item IDs
temp <- crossing(subjects, items) %>%
  left_join(lab_subj_dict, by = "subj_id") %>%
  left_join(labs, by = "lab_id") %>%
  group_by(subj_id, item_id) %>% mutate(item_id = sample(item_id)) %>%
  ungroup() %>%
  mutate(trial_num = rep(seq(n_simple + n_complex), n_subj))

temp <- temp %>%
  mutate(trial_i_var = ifelse(trial_num > 0, rnorm(nrow(temp), 0, trial_num*(increase_sd / 24)), 0))

```

```

temp %>%
  mutate(
    B_0 = beta_0 + S_0 + L_0 + I_0,

    B_c = beta_c + S_c + L_c + I_c,
    B_f = beta_f + S_f + L_f + I_f,
    B_a = beta_a + S_a + L_a + I_a,

    B_ca = beta_ca + S_ca + L_ca + I_ca,
    B_af = beta_af + S_af + L_af + I_af,
    B_cf = beta_cf + S_cf + L_cf + I_cf,

    B_cfa = beta_cfa + S_cfa + L_cfa + I_cfa,

    e_si = rnorm(nrow(temp), mean = 0, sd = sigma) + trial_i_var,

    DV = B_0 +
      (B_a * X_a) + (B_c * X_c) + (B_f * X_f) +
      (B_cf * X_c * X_f) + (B_af * X_a * X_f) + (B_ca * X_c * X_a) +
      (B_cfa * X_c * X_f * X_a) + e_si
  )
}

dat_sim <- my_sim_data_variance()

```

## 9.5 Visualise the variance increase

```

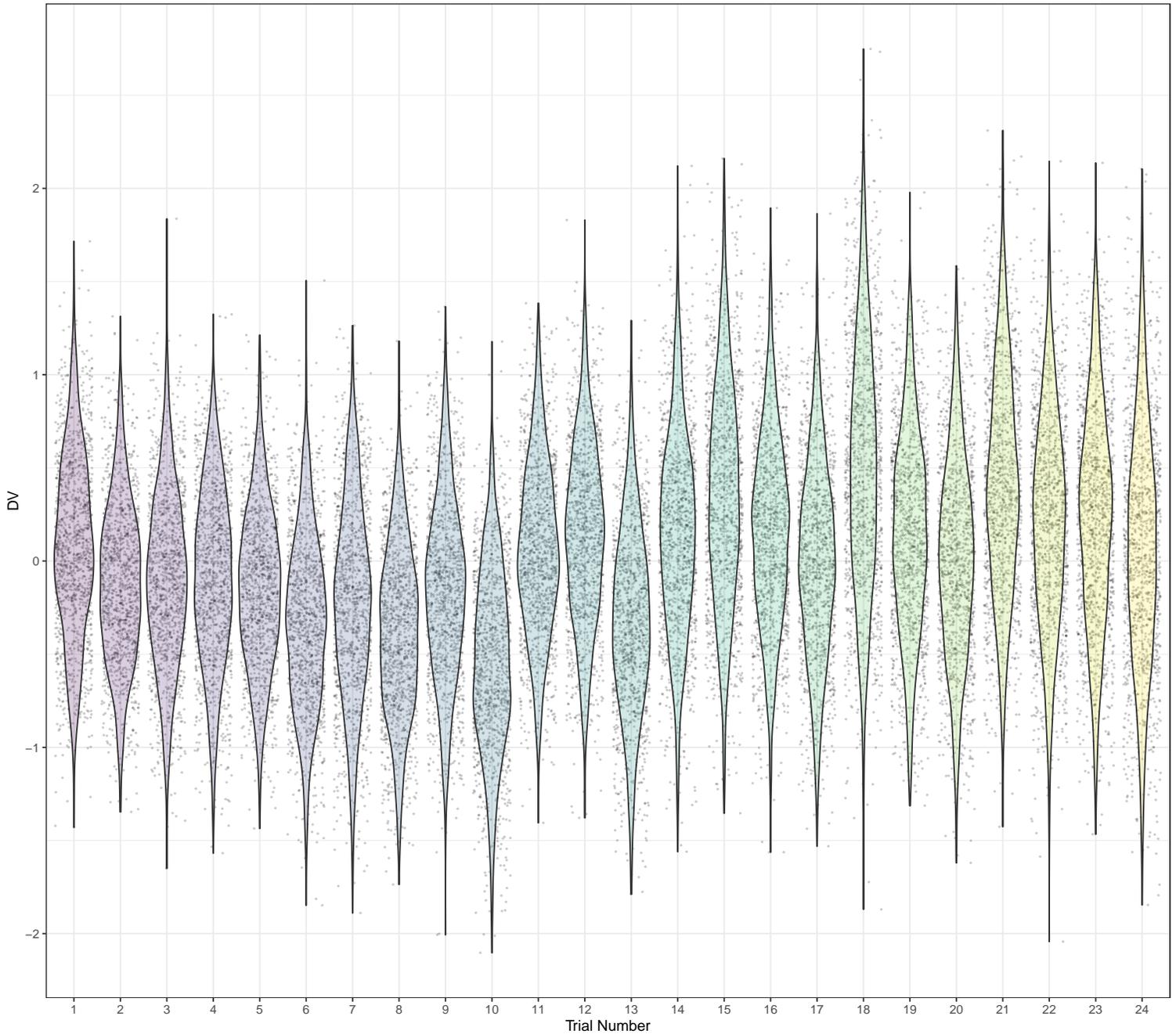
# The increase in variance is over 24 trials in this
# ManyBabies5 case:
dat_sim <- my_sim_data_variance(increase_sd = increase_sd_value)

plot_trial_num <- dat_sim %>%
  mutate(trial_num = as.factor(trial_num)) %>%
  ggplot() + geom_point(aes(y = DV, x = trial_num), position = "jitter",
    alpha = 0.2, size = 0.2) + geom_violin(aes(y = DV, x = trial_num,
    fill = trial_num), alpha = 0.2, show.legend = FALSE) + scale_fill_manual(values = viridis(n = 24)) +
  ggtitle("Increase in Variance across Trial Number") + xlab("Trial Number") +
  theme_bw()

plot_trial_num <- plot_trial_num + theme(plot.title = element_text(hjust = 0.5,
  size = 20))
plot_trial_num

```

## Increase in Variance across Trial Number



### 9.6 Model building and bias assessment

```
dat_sim <- my_sim_data_variance(increase_sd = increase_sd_value)

mod_sim <- lmer(DV ~ 1 + X_a * X_c * X_f + (1 | subj_id) + (1 |
  lab_id) + (1 | item_id), data = dat_sim)

summary(mod_sim)
```

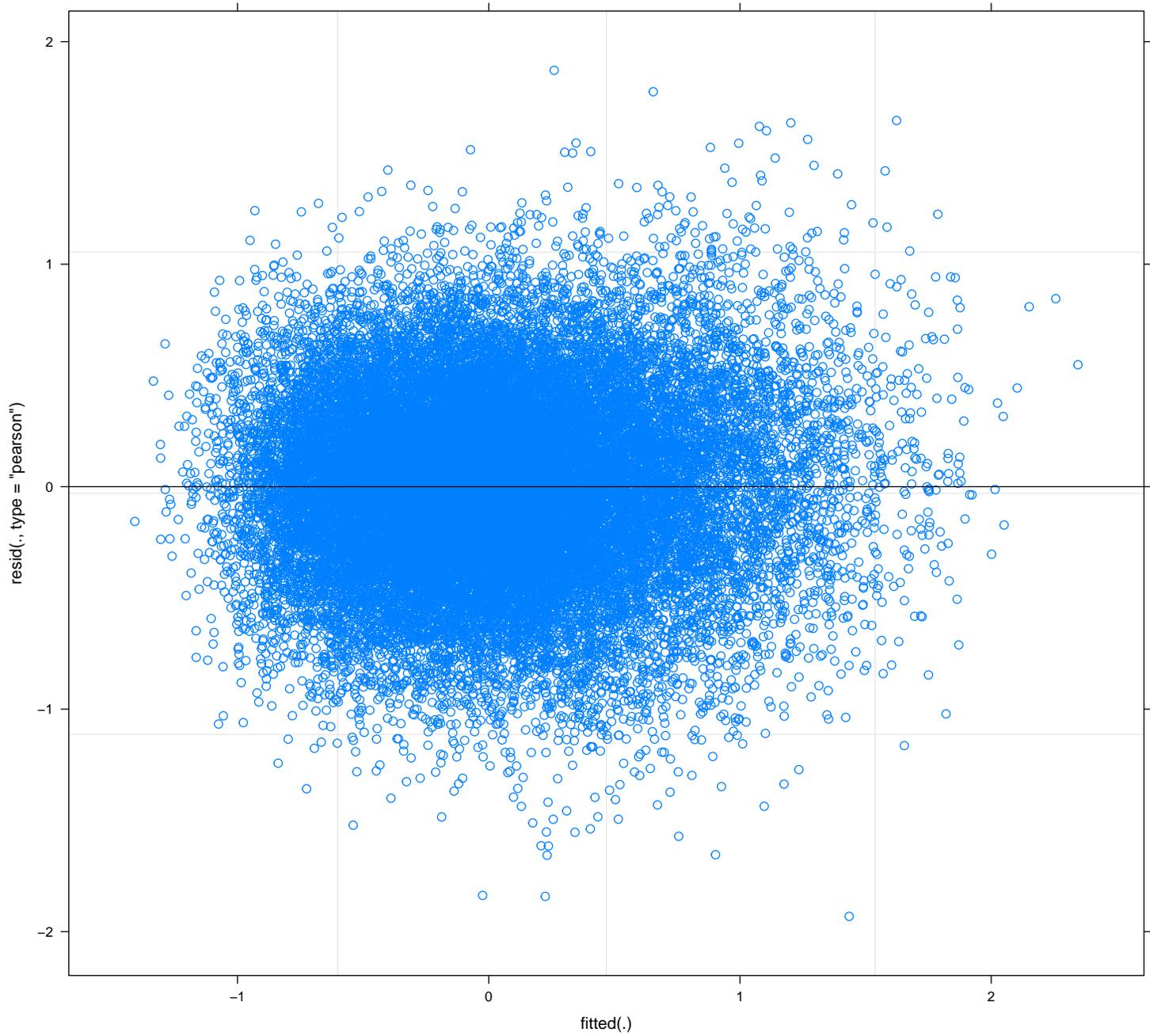
```
## Linear mixed model fit by REML. t-tests use Satterthwaite's method [
## lmerModLmerTest]
## Formula: DV ~ 1 + X_a * X_c * X_f + (1 | subj_id) + (1 | lab_id) + (1 |
##   item_id)
## Data: dat_sim
##
```

```

## REML criterion at convergence: 35957.5
##
## Scaled residuals:
##      Min       1Q   Median       3Q      Max
## -4.6460 -0.6486 -0.0005  0.6571  4.5024
##
## Random effects:
## Groups   Name                Variance Std.Dev.
## subj_id  (Intercept) 0.04078  0.2019
## lab_id   (Intercept) 0.02656  0.1630
## item_id  (Intercept) 0.06894  0.2626
## Residual                    0.17278  0.4157
## Number of obs: 30720, groups:  subj_id, 1280; lab_id, 40; item_id, 24
##
## Fixed effects:
##              Estimate Std. Error      df t value Pr(>|t|)
## (Intercept) 2.737e-02  5.978e-02 2.993e+01  0.458  0.65040
## X_a          3.720e-01  2.133e-02 1.243e+03 17.437 < 2e-16 ***
## X_c          5.855e-01  1.073e-01 2.000e+01  5.457 2.43e-05 ***
## X_f          4.687e-01  1.314e-01 2.000e+01  3.567 0.00193 **
## X_a:X_c      1.976e-01  1.627e-02 2.941e+04 12.144 < 2e-16 ***
## X_a:X_f      3.844e-01  1.992e-02 2.941e+04 19.294 < 2e-16 ***
## X_c:X_f      6.772e-01  2.628e-01 2.000e+01  2.577 0.01801 *
## X_a:X_c:X_f 7.447e-01  3.985e-02 2.941e+04 18.690 < 2e-16 ***
## ---
## Signif. codes:  0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1
##
## Correlation of Fixed Effects:
##              (Intr) X_a    X_c    X_f    X_:X_c X_:X_f X_c:X_
## X_a          -0.002
## X_c           0.000  0.000
## X_f           0.000  0.000  0.000
## X_a:X_c       0.000  0.000 -0.001  0.000
## X_a:X_f       0.000  0.000  0.000 -0.001  0.000
## X_c:X_f       0.000  0.000  0.000  0.000  0.000  0.000
## X_a:X_c:X_f  0.000  0.000  0.000  0.000  0.000  0.000 -0.001

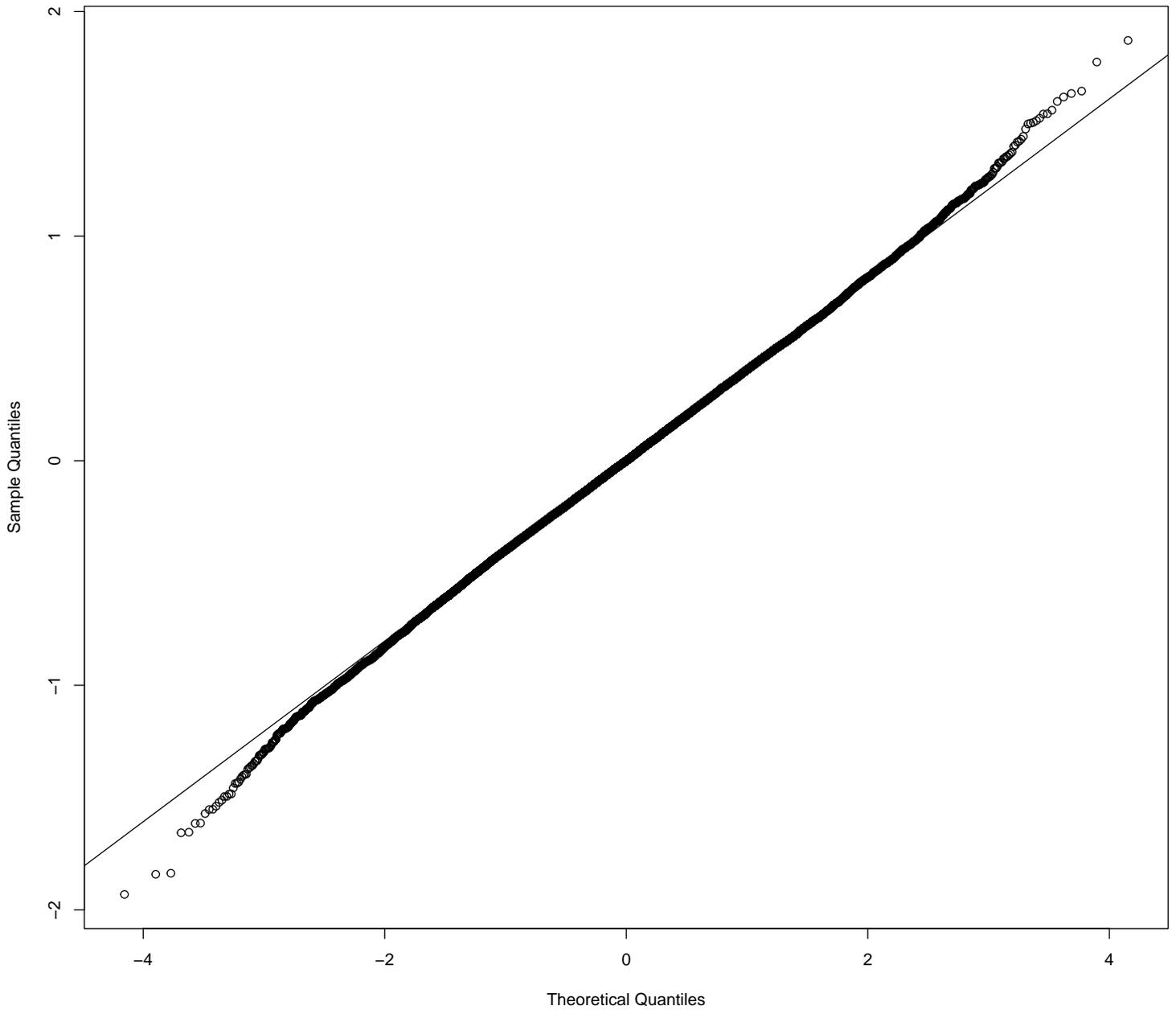
```

```
plot(mod_sim)
```



```
qqnorm(resid(mod_sim))  
qqline(resid(mod_sim))
```

Normal Q-Q Plot



```
check_normality(mod_sim)
```

```
## OK: residuals appear as normally distributed (p = 0.250).
```

```
check_outliers(mod_sim)
```

```
## OK: No outliers detected.
```

```
## - Based on the following method and threshold: cook (0.92).
```

```
## - For variable: (Whole model)
```

```
check_collinearity(mod_sim)
```

```
## # Check for Multicollinearity
```

```
##
```

```
## Low Correlation
```

```
##
```

##	Term	VIF	VIF 95% CI	Increased SE	Tolerance	Tolerance 95% CI
##	X_a	1.00	[1.00, ]	1.00	1.00	[ , 1.00]
##	X_c	1.00	[1.00, Inf]	1.00	1.00	[0.00, 1.00]
##	X_f	1.00	[1.00, Inf]	1.00	1.00	[0.00, 1.00]
##	X_a:X_c	1.00	[1.00, Inf]	1.00	1.00	[0.00, 1.00]
##	X_a:X_f	1.00	[1.00, Inf]	1.00	1.00	[0.00, 1.00]
##	X_c:X_f	1.00	[1.00, Inf]	1.00	1.00	[0.00, 1.00]
##	X_a:X_c:X_f	1.00	[1.00, Inf]	1.00	1.00	[0.00, 1.00]

```
check_heteroscedasticity(mod_sim)
```

```
## Warning: Heteroscedasticity (non-constant error variance) detected (p = 0.014).
```

## 9.7 Test how many of the models violate homoskedasticity

```
reps <- 200
```

```
run_sims_heteroskedasticity <- function(filename_heteroskedasticity) {
```

```
  dat_sim <- my_sim_data_variance(increase_sd = increase_sd_value)
```

```
  mod_sim <- lmer(DV ~ 1 + X_a * X_c * X_f + (1 | subj_id) +
    (1 | lab_id) + (1 | item_id), data = dat_sim)
```

```
  heteroskedasticity_results <- as_tibble(check_heteroscedasticity(mod_sim)[1])
```

```
  # append the results to a file
```

```
  append <- file.exists(filename_heteroskedasticity)
  write_csv(heteroskedasticity_results, filename_heteroskedasticity,
    append = append)
```

```
  # return the tidy table
```

```
  heteroskedasticity_results
```

```
}
```

```
filename_heteroskedasticity = "sims/filename_heteroskedasticity.csv"
```

```
start_time <- Sys.time()
```

```
heteroskedasticity_results_data <- purrr::map_df(1:reps, ~run_sims_heteroskedasticity(filename_heteroskedasticity))
```

```
end_time <- Sys.time()
```

```
end_time - start_time
```

## 9.8 Overview of how many of the models exhibit homoskedasticity

```
heteroskedasticity_results <- read.csv(filename_heteroskedasticity)
```

```
homoskedastic_models <- heteroskedasticity_results %>%
  filter(value > 0.05)
```

```
print(paste0(nrow(homoskedastic_models), "/", nrow(heteroskedasticity_results),
  " of the models exhibit homoskedasticity"))
```

```
## [1] "156/200 of the models exhibit homoskedasticity"
```

## 9.9 Bayesian robust location-scale regression model

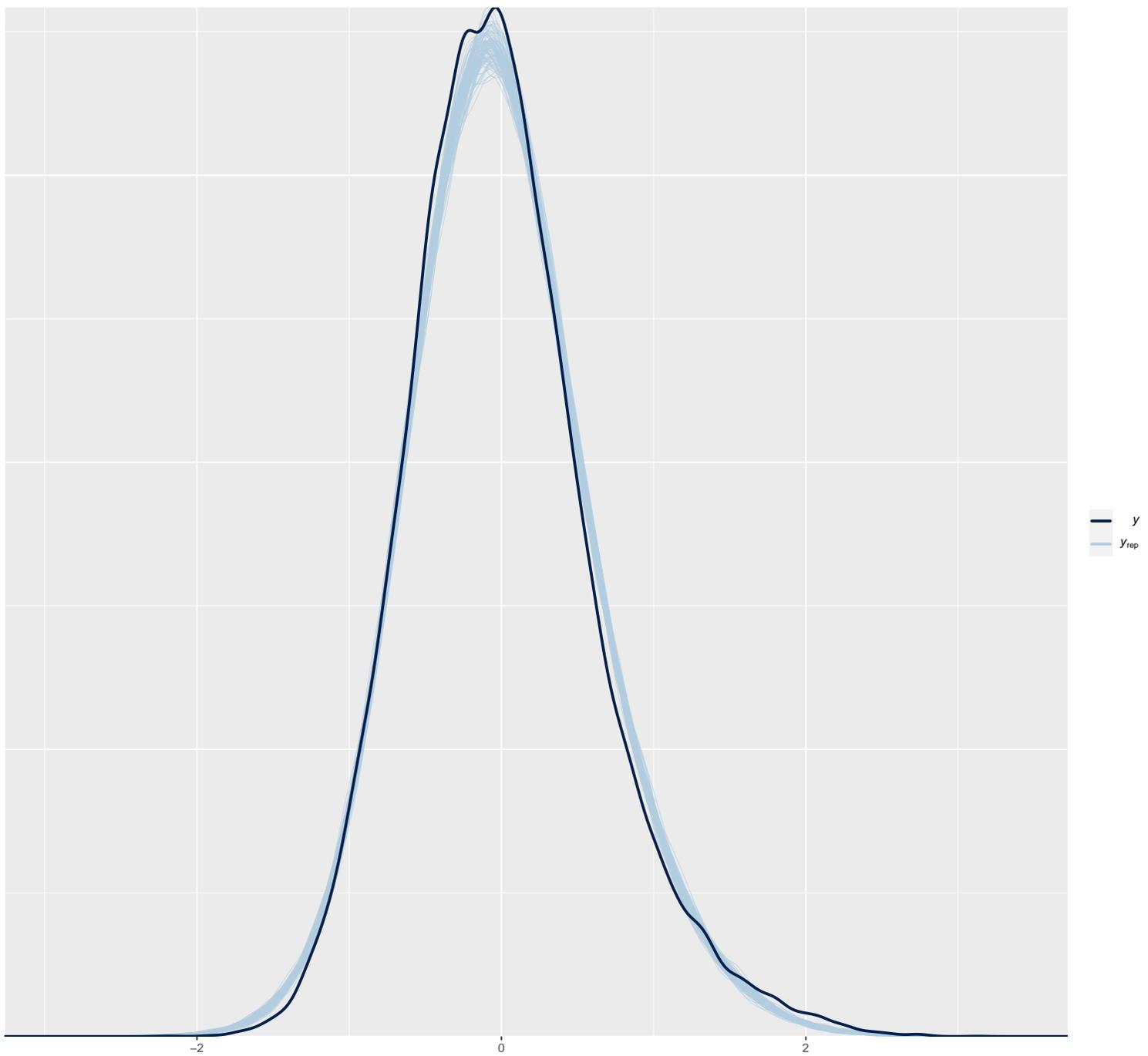
```
model_formula <- bf(DV ~ 1 + X_a * X_c * X_f +
  (1 | subj_id) + (1 | lab_id) + (1 | item_id),
  sigma ~ 1 + trial_num + (1 | subj_id) + (1 | lab_id) + (1 | item_id))

#get_prior(model_formula, data = dat_sim, family = student)

priors1 <- c(prior(normal(0, 0.5), class = Intercept),
  prior(normal(0, 0.5), class = b),
  prior(normal(0, 0.5), class = b, dpar = sigma),
  prior(normal(0.25, 0.3), class = sd),
  prior(gamma(2, 0.1), class = nu)
)

heteroskedasticity_fit <-
  brm(data = dat_sim,
    family = student,
    model_formula,
    prior = priors1,
    sample_prior = "yes",
    iter = 4000,
    warmup = 500,
    #backend = "cmdstanr",
    #threads = threading(2),
    file = "heteroskedasticity_fit",
    cores = 64,
    chains = 2,
    save_pars = save_pars(all = TRUE))

pp_check(heteroskedasticity_fit, ndraws = 100)
```



heteroskedasticity\_fit

```
## Family: student
## Links: mu = identity; sigma = log; nu = identity
## Formula: DV ~ 1 + X_a * X_c * X_f + (1 | subj_id) + (1 | lab_id) + (1 | item_id)
##          sigma ~ 1 + trial_num + (1 | subj_id) + (1 | lab_id) + (1 | item_id)
## Data: dat_sim (Number of observations: 30720)
## Draws: 2 chains, each with iter = 4000; warmup = 500; thin = 1;
##        total post-warmup draws = 7000
##
## Group-Level Effects:
## ~item_id (Number of levels: 24)
##          Estimate Est.Error 1-95% CI u-95% CI Rhat Bulk_ESS Tail_ESS
## sd(Intercept)      0.00     0.00    0.00    0.01 1.00    3967    4291
## sd(sigma_Intercept) 0.01     0.01    0.00    0.02 1.00    3111    3651
##
## ~lab_id (Number of levels: 40)
```

```

##           Estimate Est.Error 1-95% CI u-95% CI Rhat Bulk_ESS Tail_ESS
## sd(Intercept)      0.17    0.02    0.13    0.22 1.00    1881    3066
## sd(sigma_Intercept) 0.04    0.01    0.03    0.06 1.00    3176    5234
##
## ~subj_id (Number of levels: 1280)
##           Estimate Est.Error 1-95% CI u-95% CI Rhat Bulk_ESS Tail_ESS
## sd(Intercept)      0.21    0.01    0.20    0.22 1.00    2481    3755
## sd(sigma_Intercept) 0.05    0.01    0.03    0.07 1.00    1260    1282
##
## Population-Level Effects:
##           Estimate Est.Error 1-95% CI u-95% CI Rhat Bulk_ESS Tail_ESS
## Intercept          -0.01    0.03   -0.06    0.05 1.00    1051    1816
## sigma_Intercept    -1.07    0.01   -1.09   -1.04 1.00    6208    5953
## X_a                 0.26    0.02    0.22    0.31 1.00    2180    3628
## X_c                 0.30    0.01    0.29    0.32 1.00   11577    5326
## X_f                 0.39    0.01    0.38    0.40 1.00   14033    5762
## X_a:X_c             0.45    0.02    0.42    0.49 1.00   17307    5375
## X_a:X_f             0.67    0.02    0.62    0.71 1.00   14751    5860
## X_c:X_f             0.73    0.01    0.70    0.75 1.00   16307    5246
## X_a:X_c:X_f        0.36    0.04    0.27    0.44 1.00   15476    5133
## sigma_trial_num    0.02    0.00    0.02    0.02 1.00    7882    5137
##
## Family Specific Parameters:
##           Estimate Est.Error 1-95% CI u-95% CI Rhat Bulk_ESS Tail_ESS
## nu      81.74     19.02     52.07  125.80 1.00   11984    5331
##
## Draws were sampled using sampling(NUTS). For each parameter, Bulk_ESS
## and Tail_ESS are effective sample size measures, and Rhat is the potential
## scale reduction factor on split chains (at convergence, Rhat = 1).

```

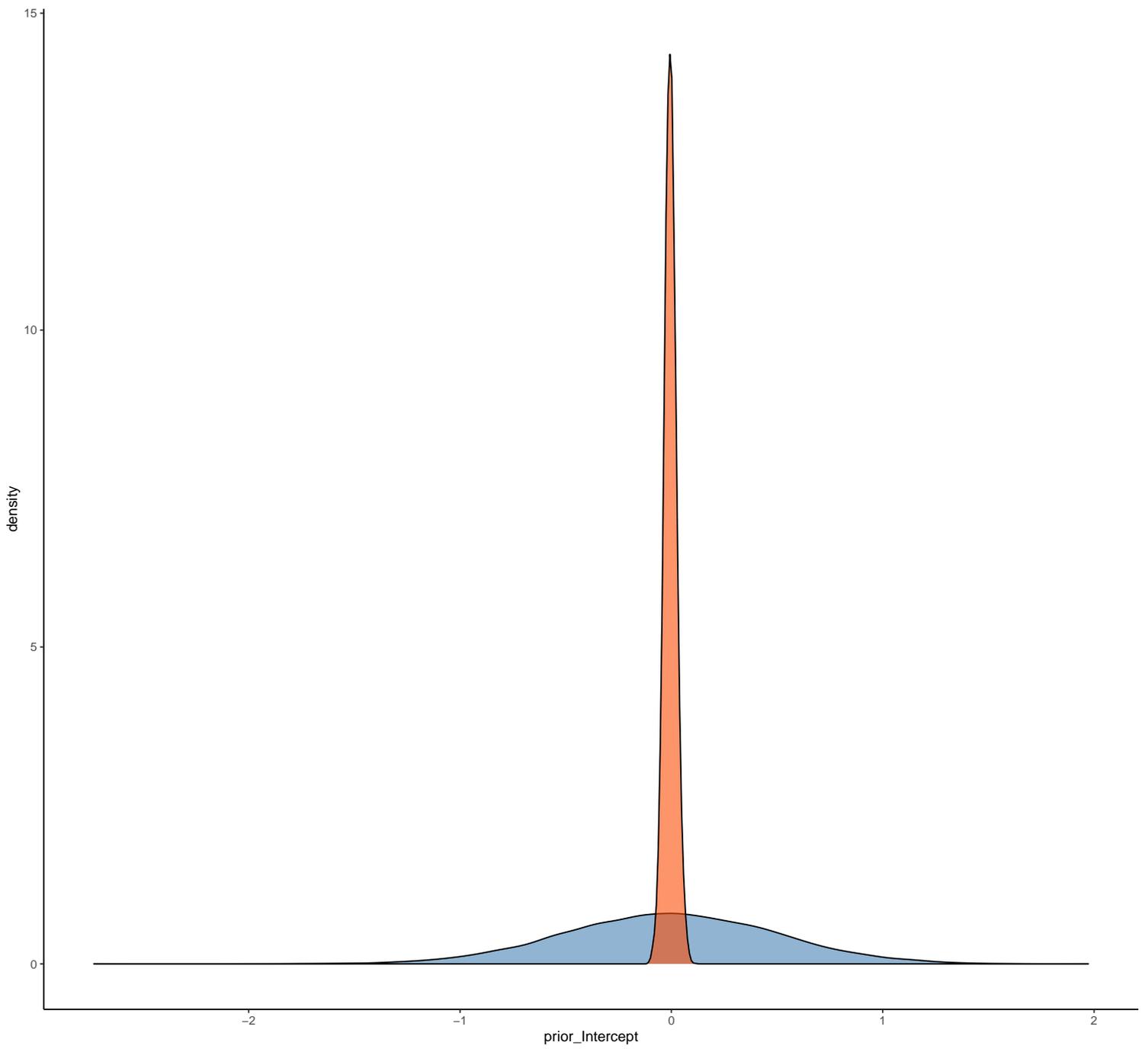
### 9.9.1 Prior-Posterior Update Checks

```

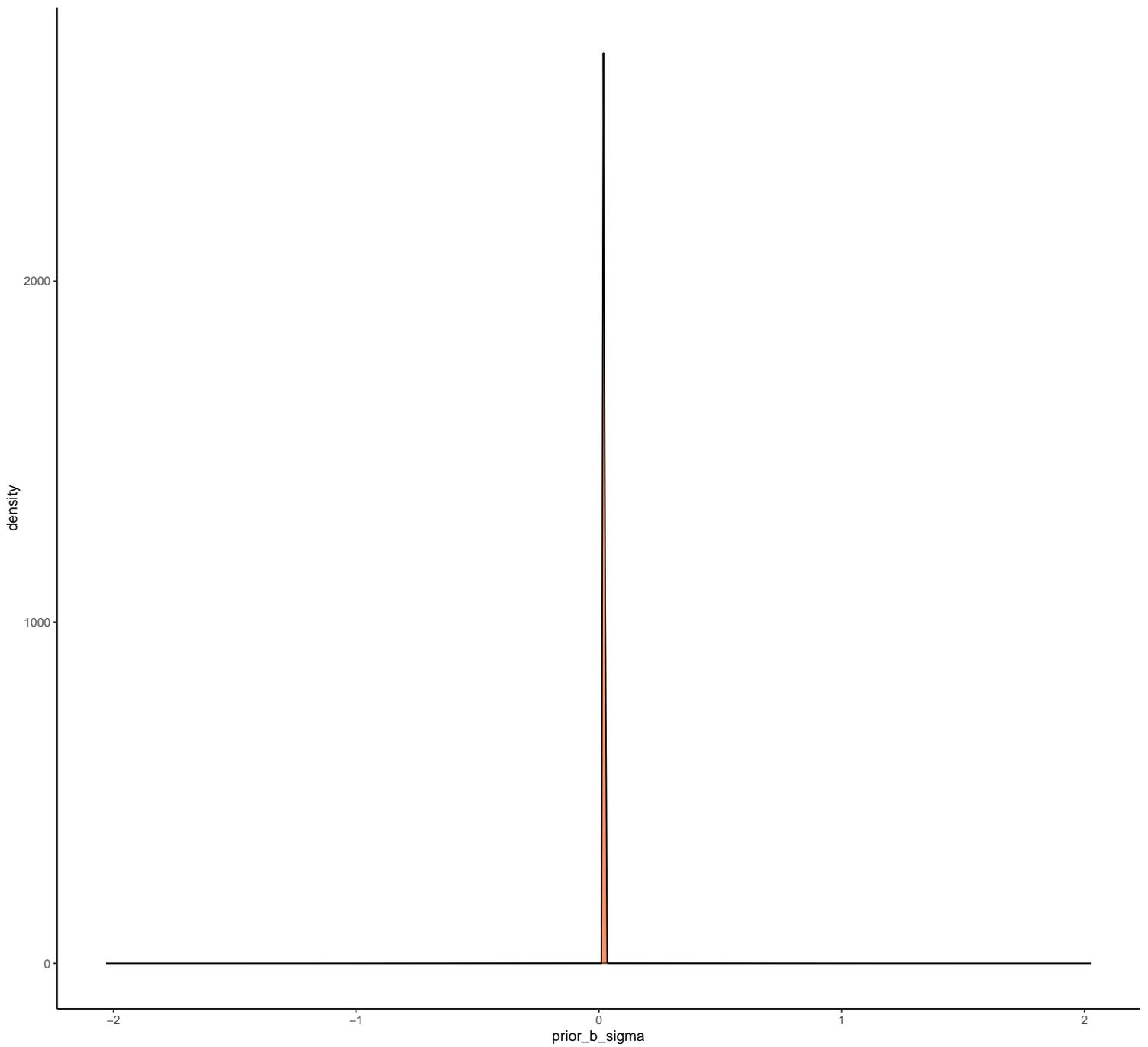
# Sample the parameters of interest:
Posterior_m1 <- as_draws_df(heteroskedasticity_fit)

# Plot the prior-posterior update plot for the intercept:
ggplot(Posterior_m1) + geom_density(aes(prior_Intercept), fill = "steelblue",
  color = "black", alpha = 0.6) + geom_density(aes(b_Intercept),
  fill = "#FC4E07", color = "black", alpha = 0.6) + theme_classic()

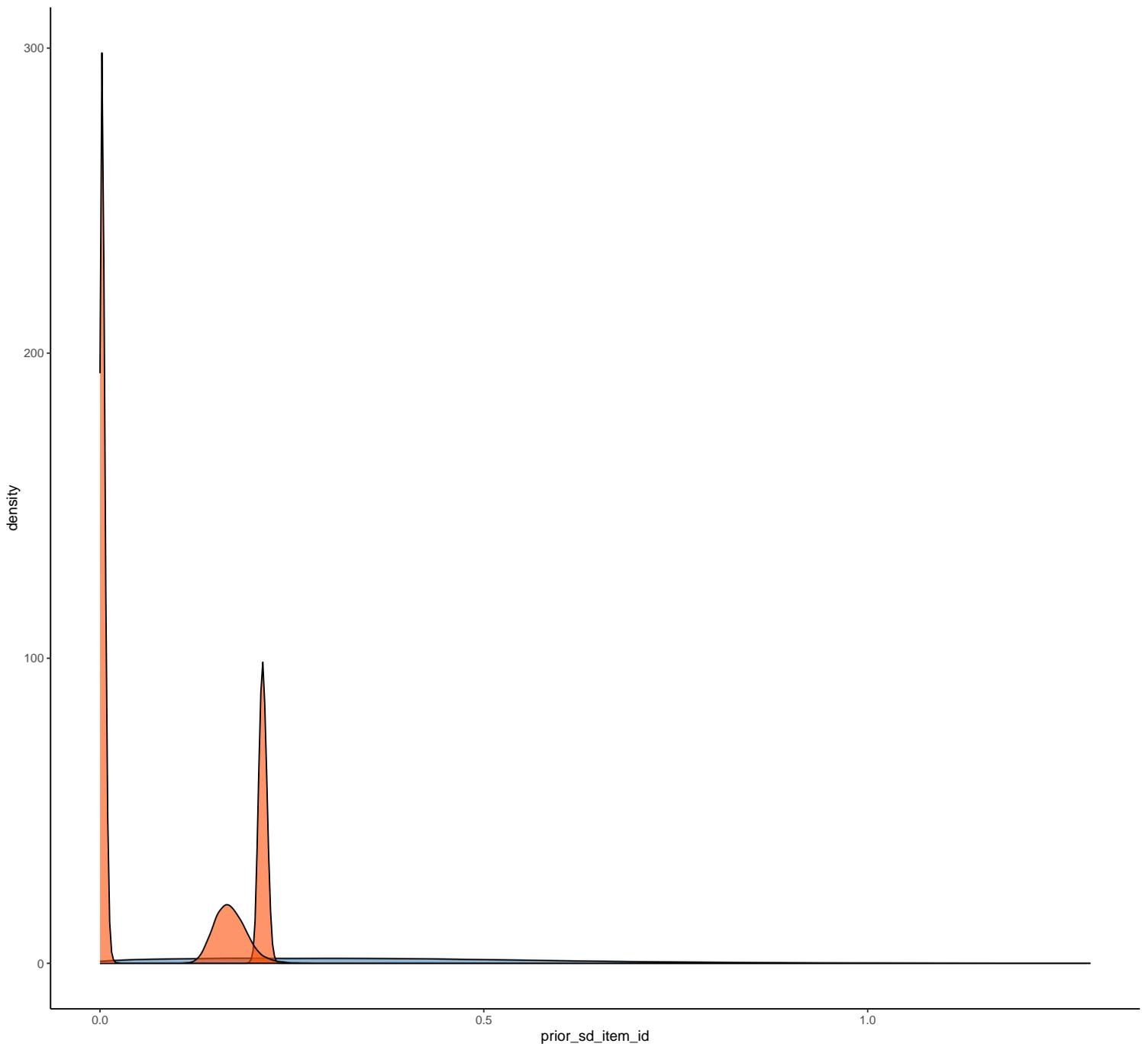
```



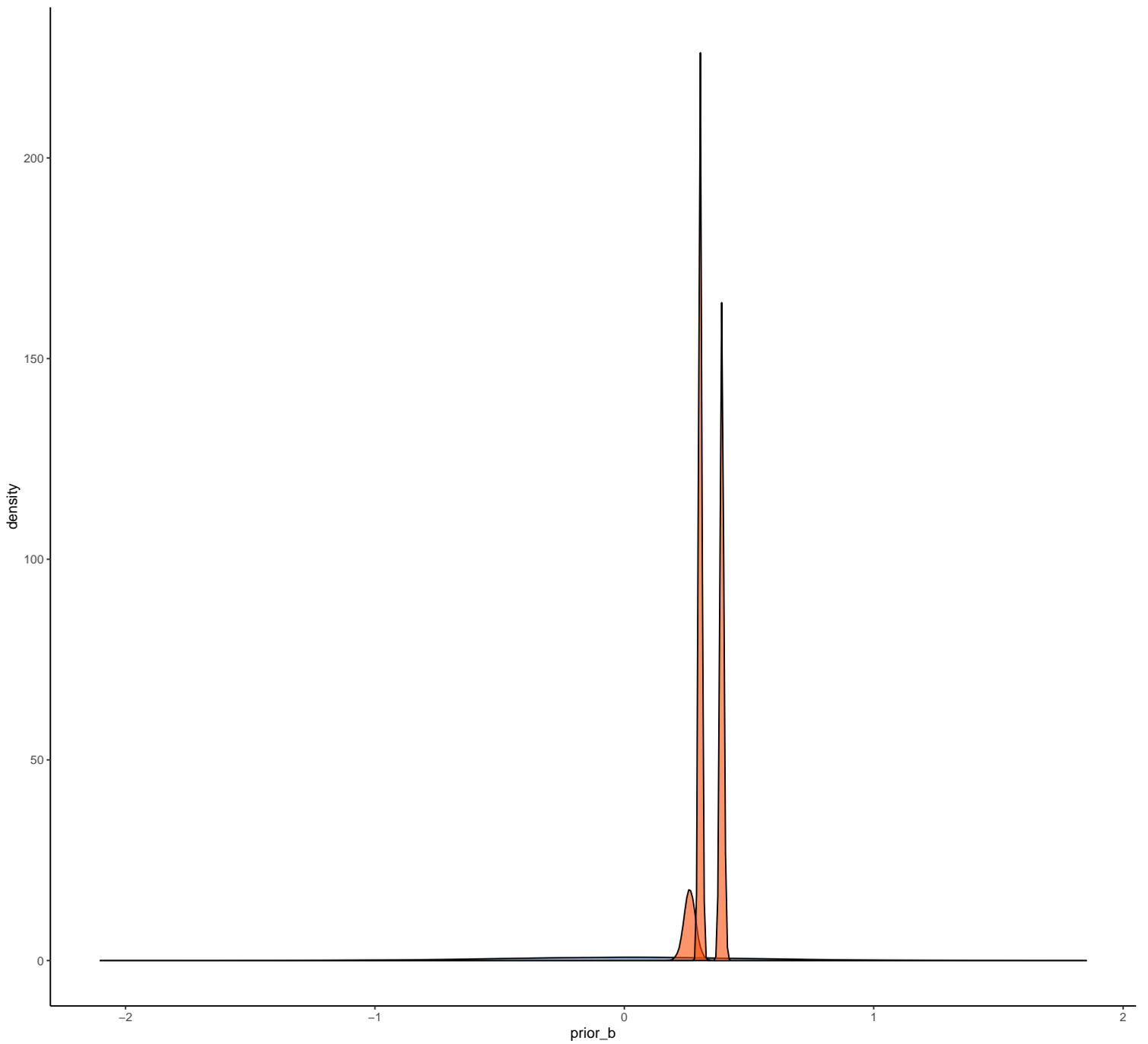
```
ggplot(Posterior_m1) + geom_density(aes(prior_b_sigma), fill = "steelblue",  
  color = "black", alpha = 0.6) + geom_density(aes(b_sigma_trial_num),  
  fill = "#FC4E07", color = "black", alpha = 0.6) + theme_classic()
```



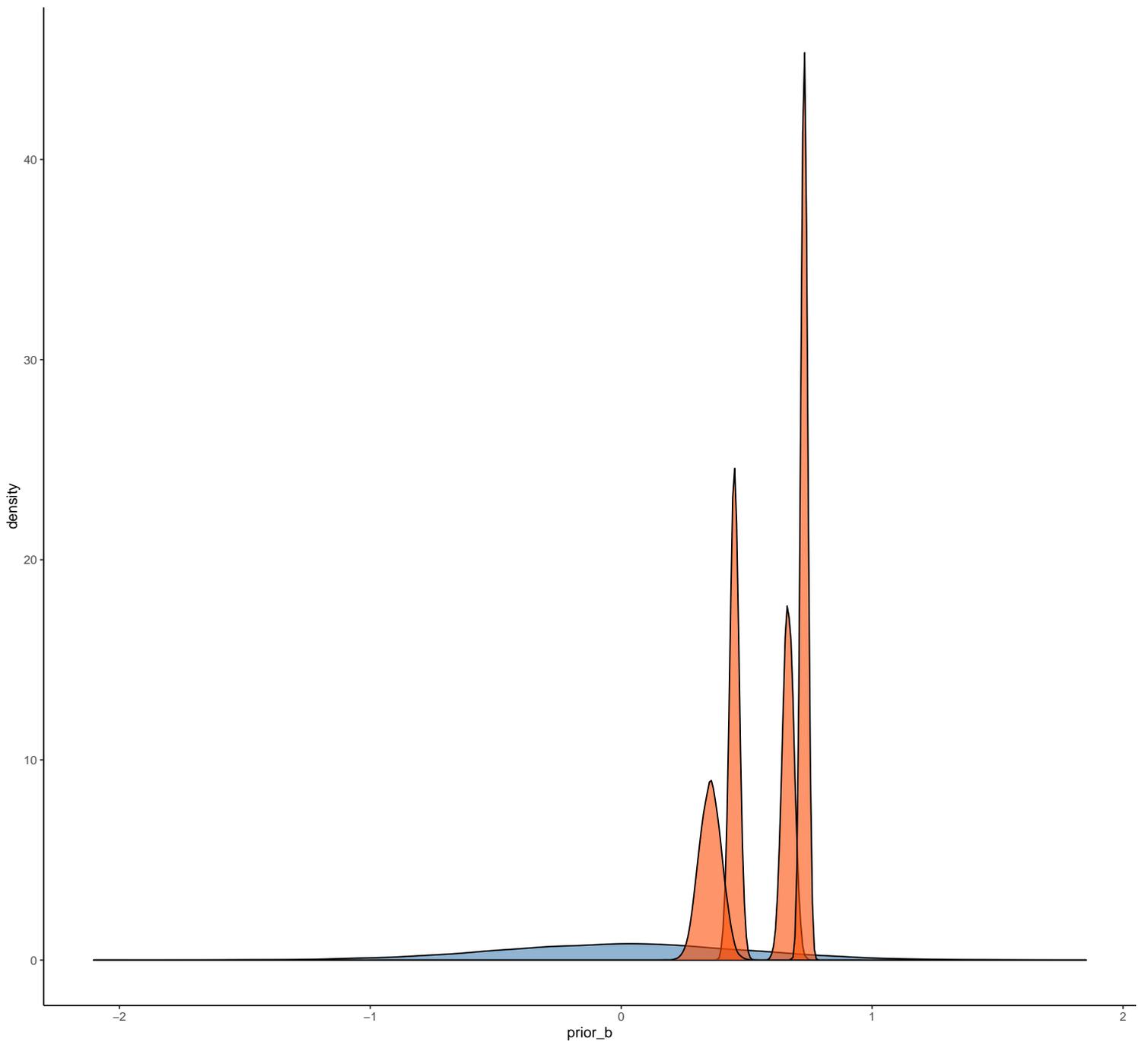
```
ggplot(Posterior_m1) + geom_density(aes(prior_sd_item_id), fill = "steelblue",  
  color = "black", alpha = 0.6) + geom_density(aes(sd_subj_id__Intercept),  
  fill = "#FC4E07", color = "black", alpha = 0.6) + geom_density(aes(sd_lab_id__Intercept),  
  fill = "#FC4E07", color = "black", alpha = 0.6) + geom_density(aes(sd_item_id__Intercept),  
  fill = "#FC4E07", color = "black", alpha = 0.6) + theme_classic()
```



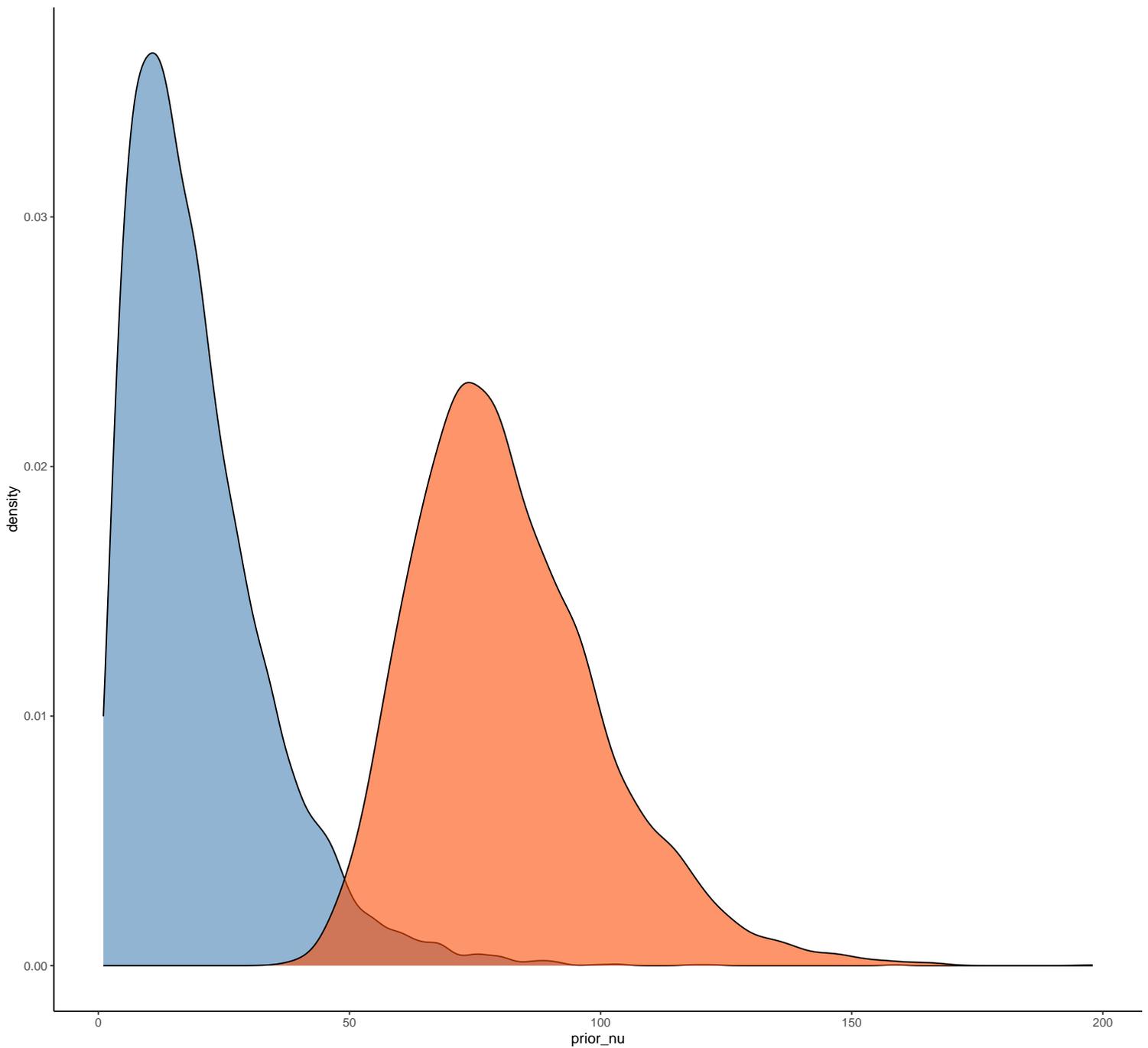
```
ggplot(Posterior_m1) + geom_density(aes(prior_b), fill = "steelblue",  
  color = "black", alpha = 0.6) + geom_density(aes(b_X_a),  
  fill = "#FC4E07", color = "black", alpha = 0.6) + geom_density(aes(b_X_c),  
  fill = "#FC4E07", color = "black", alpha = 0.6) + geom_density(aes(b_X_f),  
  fill = "#FC4E07", color = "black", alpha = 0.6) + theme_classic()
```



```
ggplot(Posterior_m1) + geom_density(aes(prior_b), fill = "steelblue",  
  color = "black", alpha = 0.6) + geom_density(aes(`b_X_a:X_c`),  
  fill = "#FC4E07", color = "black", alpha = 0.6) + geom_density(aes(`b_X_a:X_f`),  
  fill = "#FC4E07", color = "black", alpha = 0.6) + geom_density(aes(`b_X_c:X_f`),  
  fill = "#FC4E07", color = "black", alpha = 0.6) + geom_density(aes(`b_X_a:X_c:X_f`),  
  fill = "#FC4E07", color = "black", alpha = 0.6) + theme_classic()
```



```
ggplot(Posterior_m1) + geom_density(aes(prior_nu), fill = "steelblue",  
  color = "black", alpha = 0.6) + geom_density(aes(nu), fill = "#FC4E07",  
  color = "black", alpha = 0.6) + theme_classic()
```



## 10 Data missing not completely at random

### 10.1 Simulation of missing data according to increasing infant age

```
# back to assumptions of equal variance across trials:
dat_sim <- my_sim_data()

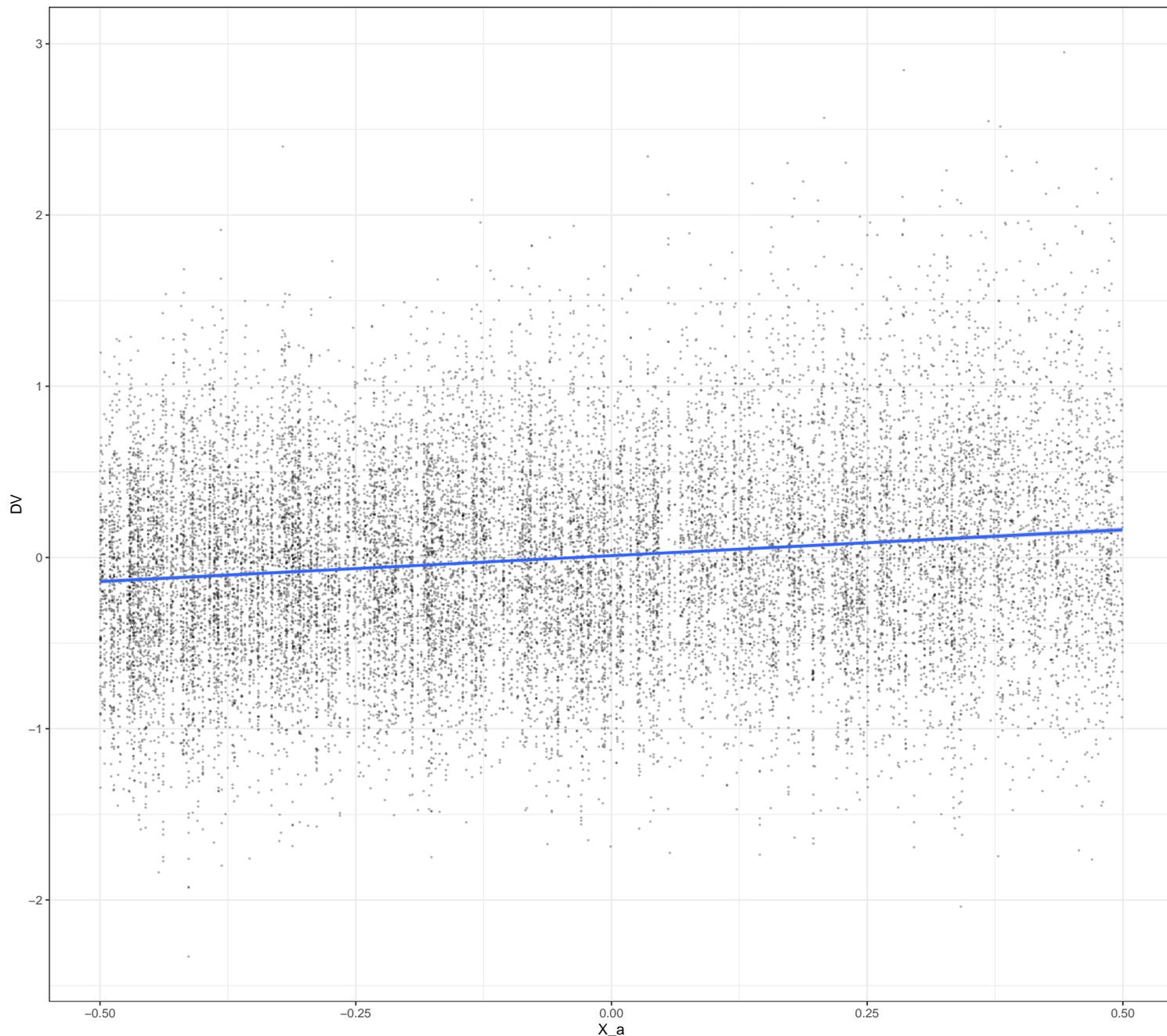
# Proportion of missing data increases with age:
missing_samples_age <- dat_sim %>%
  mutate(nas = rbinom(n(), 1, 0.95 - ifelse(X_a > -0.5, (X_a +
    0.5) * 0.5, 0))) %>%
  mutate(DV = ifelse(nas == 1, DV, NA))

missing_age_plot <- missing_samples_age %>%
  drop_na() %>%
```

```
ggplot() + ggtitle("DV as a function of age, missing data") +  
geom_point(aes(x = X_a, y = DV), alpha = 0.3, size = 0.2,  
  position = "jitter") + geom_smooth(aes(x = X_a, y = DV),  
  method = "lm", se = TRUE, formula = y ~ x) + theme_bw()
```

```
missing_age_plot <- missing_age_plot + theme(plot.title = element_text(hjust = 0.5,  
  size = 20))  
missing_age_plot
```

DV as a function of age, missing data



### 10.1.1 Simulation of models

```
# Number of simulations:  
reps <- 100  
  
# Simulation function:
```

```

run_sims <- function(filename_full, ef) {

  dat_sim <- my_sim_data(beta_c = ef,
                        beta_f = ef,
                        beta_a = ef,

                        beta_ca = ef,
                        beta_af = ef,
                        beta_cf = ef,

                        beta_cfa = ef)

  missing_samples_age <- dat_sim %>%
    mutate(nas = rbinom(n(), 1, 0.95 - ifelse(X_a > -0.5, (X_a+0.5)*0.50, 0))) %>%
    mutate(DV = ifelse(nas == 1, DV, NA))

  mod_sim <- lmer(DV ~ 1 + X_a * X_c * X_f +
                 (1 | subj_id) +
                 (1 | lab_id) +
                 (1 | item_id),
                 data=missing_samples_age)

  sim_results <- broom.mixed::tidy(mod_sim)

  # append the results to a file
  append <- file.exists(filename_full)
  write_csv(sim_results, filename_full, append = append)

  # return the tidy table
  sim_results
}

filename_full_0.3_missing_age = 'sims/run_sims_0.3_age_missing.csv'
start_time <- Sys.time()
sims <- purrr::map_df(1:reps, ~run_sims(filename_full = filename_full_0.3_missing_age, ef = 0.3))
end_time <- Sys.time()
end_time - start_time

```

### 10.1.2 Visualise Estimates for Fixed Effects:

```

# read saved simulation data
sims_50_missing_age_0.3 <- read_csv(filename_full_0.3_missing_age, col_types = cols(
  # makes sure plots display in this order
  group = col_factor(ordered = TRUE),
  term = col_factor(ordered = TRUE)
))

reps <- 100

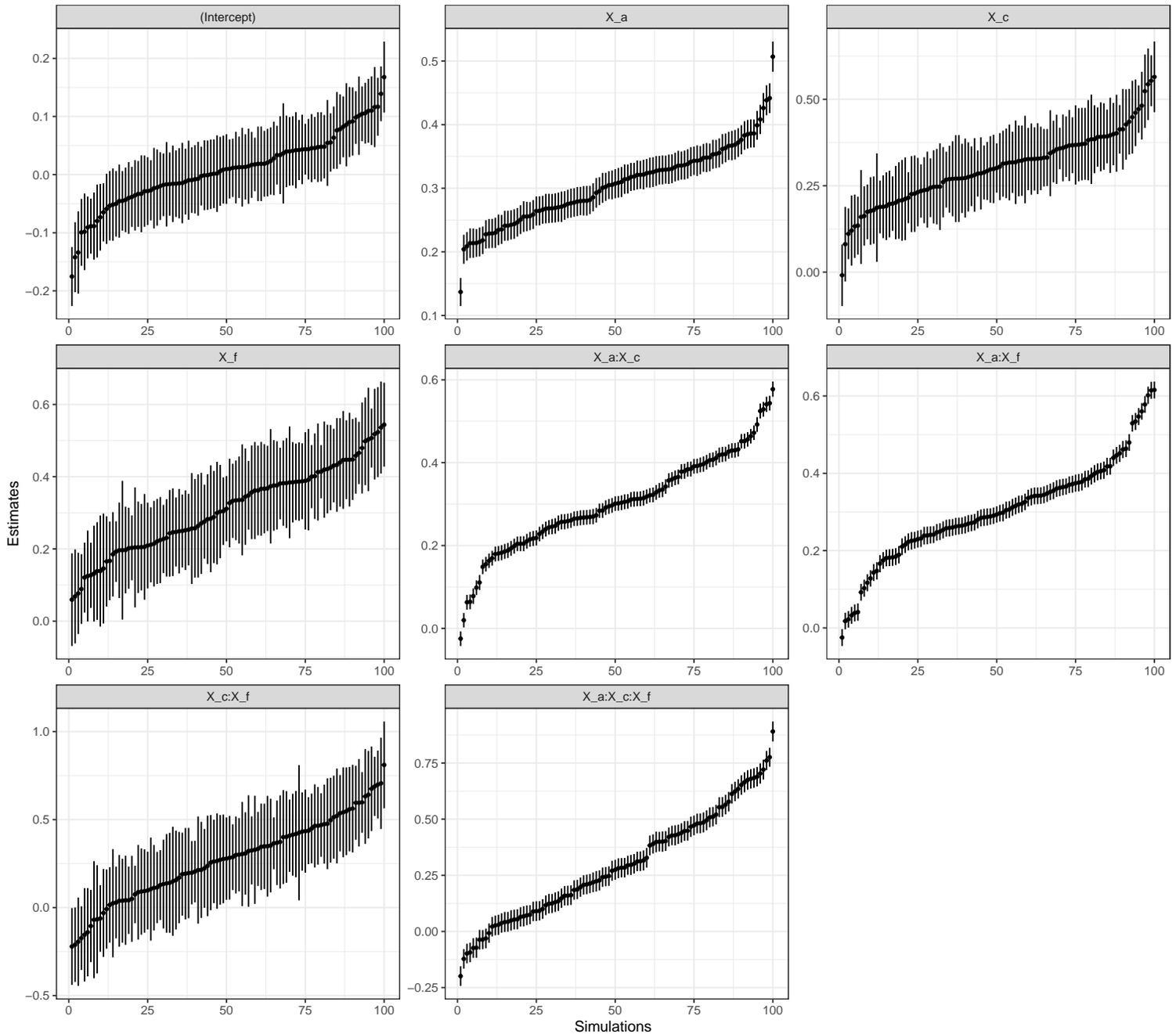
fixed_missing_age_plot <- sims_50_missing_age_0.3 %>%
  filter(effect == "fixed") %>%
  ungroup() %>%
  arrange(term, estimate) %>%
  mutate(row = rep(seq(1:reps), 8)) %>%
  ggplot(aes(x = row, y = estimate, ymin = estimate-std.error, ymax = estimate+std.error)) +
  facet_wrap(~term, scales = "free") +
  geom_pointrange(fatten = 1/2) +
  ylab("Estimates") +
  xlab("Simulations") +
  ggtitle('Estimates of Fixed Effects for Missing Data with Age, ef = 0.3') +

```

```
theme_bw()
```

```
fixed_missing_age_plot <- fixed_missing_age_plot + theme(plot.title = element_text(hjust = 0.5, size=20))  
fixed_missing_age_plot
```

### Estimates of Fixed Effects for Missing Data with Age, ef = 0.3



#### 10.1.3 Visualise Estimates for Random Effects:

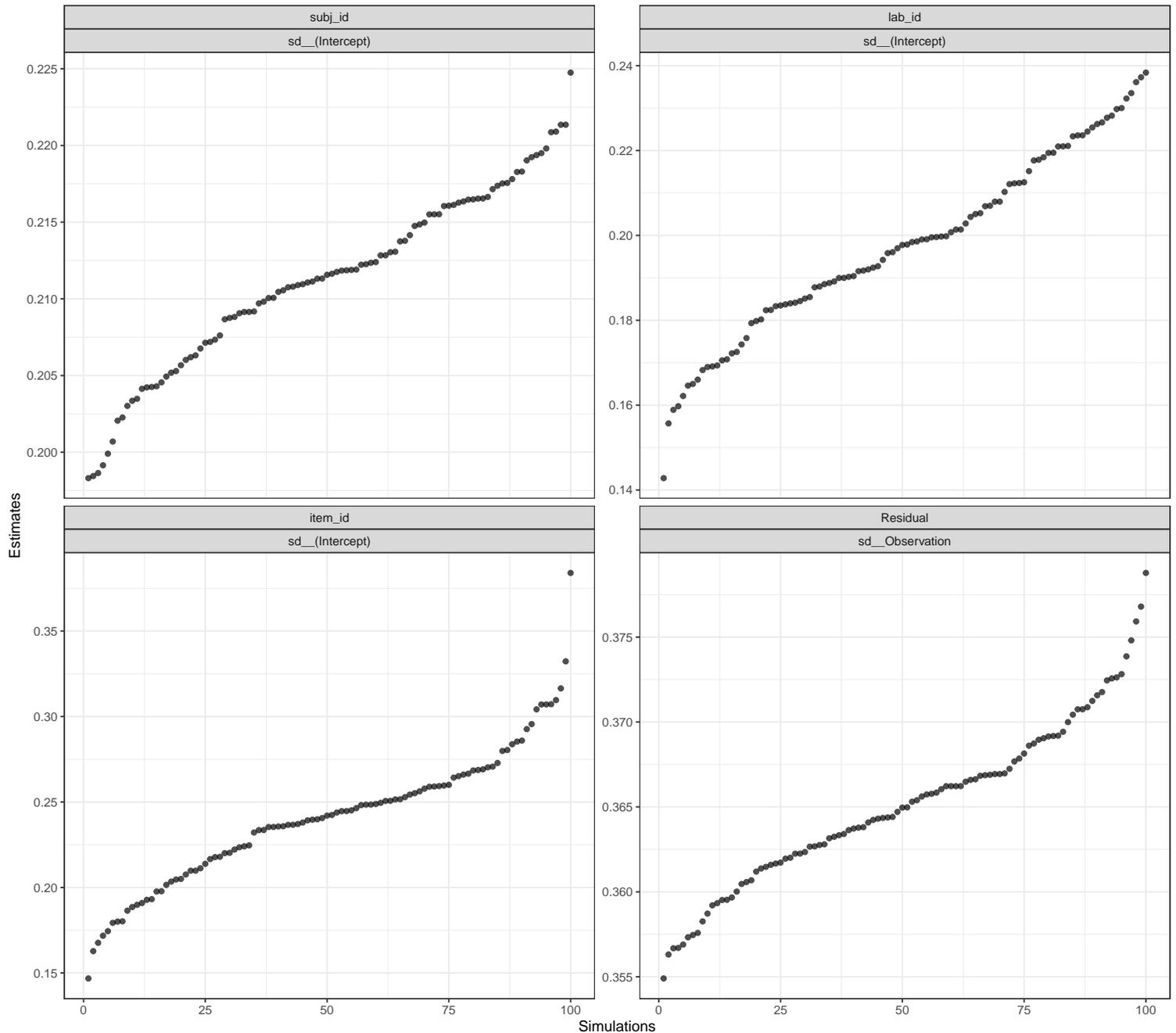
```
ran_missing_age_plot <- sims_50_missing_age_0.3 %>%  
  filter(effect == "ran_pars") %>%  
  ungroup() %>%  
  arrange(group, term, estimate) %>%  
  mutate(row = rep(seq(1:reps), 4)) %>%  
  ggplot(aes(x = row, y = estimate)) + geom_point(alpha = 0.7) +  
  facet_wrap(~group + term, scales = "free_y") + theme_bw() +  
  ylab("Estimates") + xlab("Simulations") + ggtitle("Estimates of Random Effects for Missing Data with Age, ef
```

```

theme_bw()
ran_missing_age_plot <- ran_missing_age_plot + theme(plot.title = element_text(hjust = 0.5,
  size = 20))
ran_missing_age_plot

```

## Estimates of Random Effects for Missing Data with Age, $ef = 0.3$



## 10.2 Simulation of missing data according to increasing trial number

```

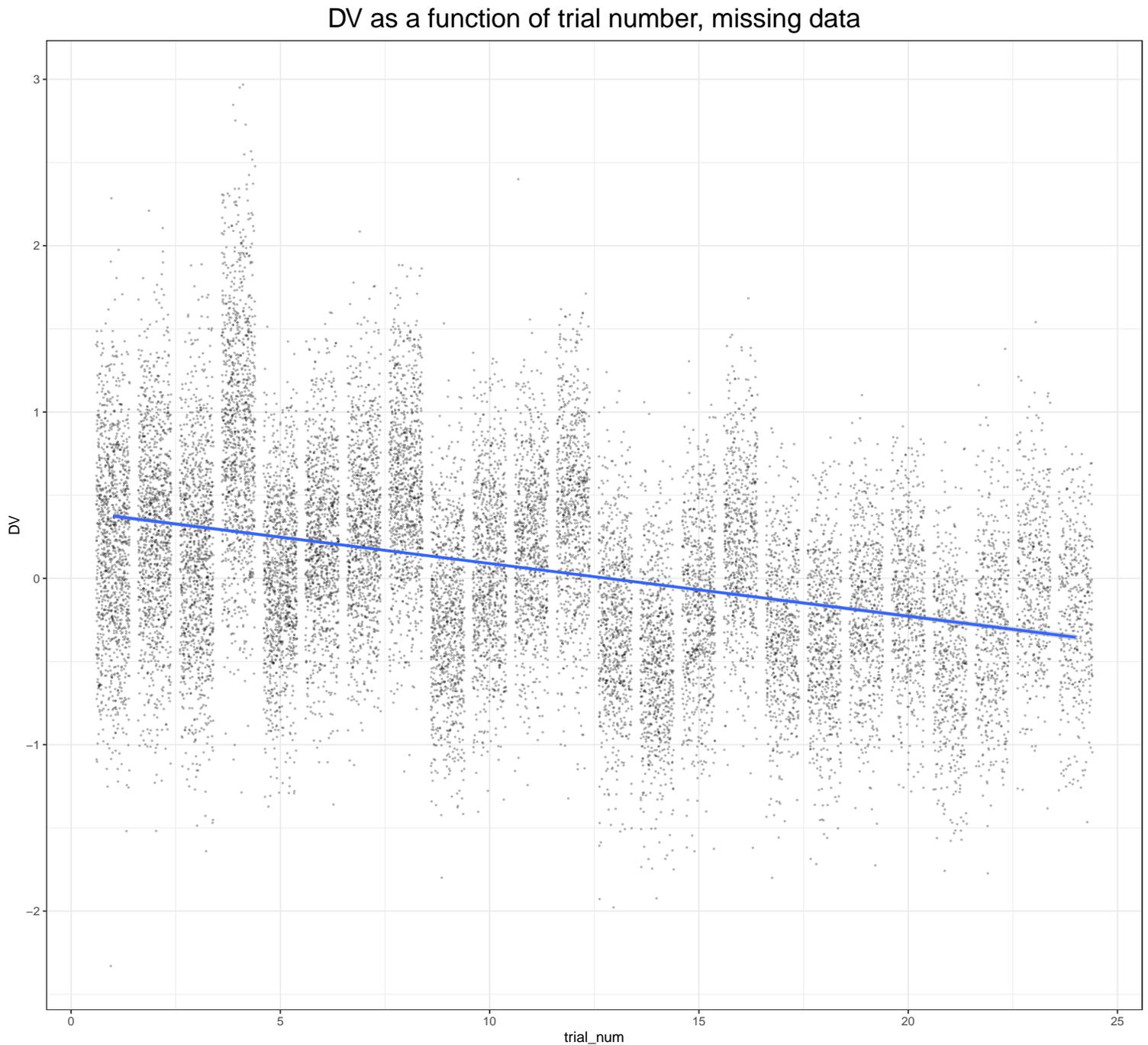
# Proportion of missing data increases with trial number:
missing_samples_trial <- dat_sim %>%
  mutate(nas = rbinom(n(), 1, 0.95 - ifelse(trial_num > 0,
    trial_num * 0.025, 0))) %>%
  mutate(DV = ifelse(nas == 1, DV, NA))

missing_trialnum_plot <- missing_samples_trial %>%
  drop_na() %>%

```

```
ggplot() + geom_point(aes(x = trial_num, y = DV), alpha = 0.3,  
size = 0.2, position = "jitter") + geom_smooth(aes(x = trial_num,  
y = DV), method = "lm", se = TRUE, formula = y ~ x) + ggtitle("DV as a function of trial number, missing data")  
theme_bw()
```

```
missing_trialnum_plot <- missing_trialnum_plot + theme(plot.title = element_text(hjust = 0.5,  
size = 20))  
missing_trialnum_plot
```



### 10.2.1 Simulation of models

```
# Number of simulations:  
reps <- 100  
  
# Simulation function:
```

```

run_sims <- function(filename_full, ef) {

  dat_sim <- my_sim_data(beta_c = ef,
                        beta_f = ef,
                        beta_a = ef,

                        beta_ca = ef,
                        beta_af = ef,
                        beta_cf = ef,

                        beta_cfa = ef)

  missing_samples_trial <- dat_sim %>%
    mutate(nas = rbinom(n(), 1, 0.95 - ifelse(trial_num > 0, trial_num*0.025, 0))) %>%
    mutate(DV = ifelse(nas == 1, DV, NA))

  mod_sim <- lmer(DV ~ 1 + X_a * X_c * X_f +
                 (1 | subj_id) +
                 (1 | lab_id) +
                 (1 | item_id),
                 data=missing_samples_trial)

  sim_results <- broom.mixed::tidy(mod_sim)

  # append the results to a file
  append <- file.exists(filename_full)
  write_csv(sim_results, filename_full, append = append)

  # return the tidy table
  sim_results
}

filename_full_0.3_missing_trial = 'sims/run_sims_0.3_trial_missing.csv'
start_time <- Sys.time()
sims <- purrr::map_df(1:reps, ~run_sims(filename_full = filename_full_0.3_missing_trial, ef = 0.3))
end_time <- Sys.time()
end_time - start_time

```

## 10.2.2 Visualise Estimates for Fixed Effects:

```

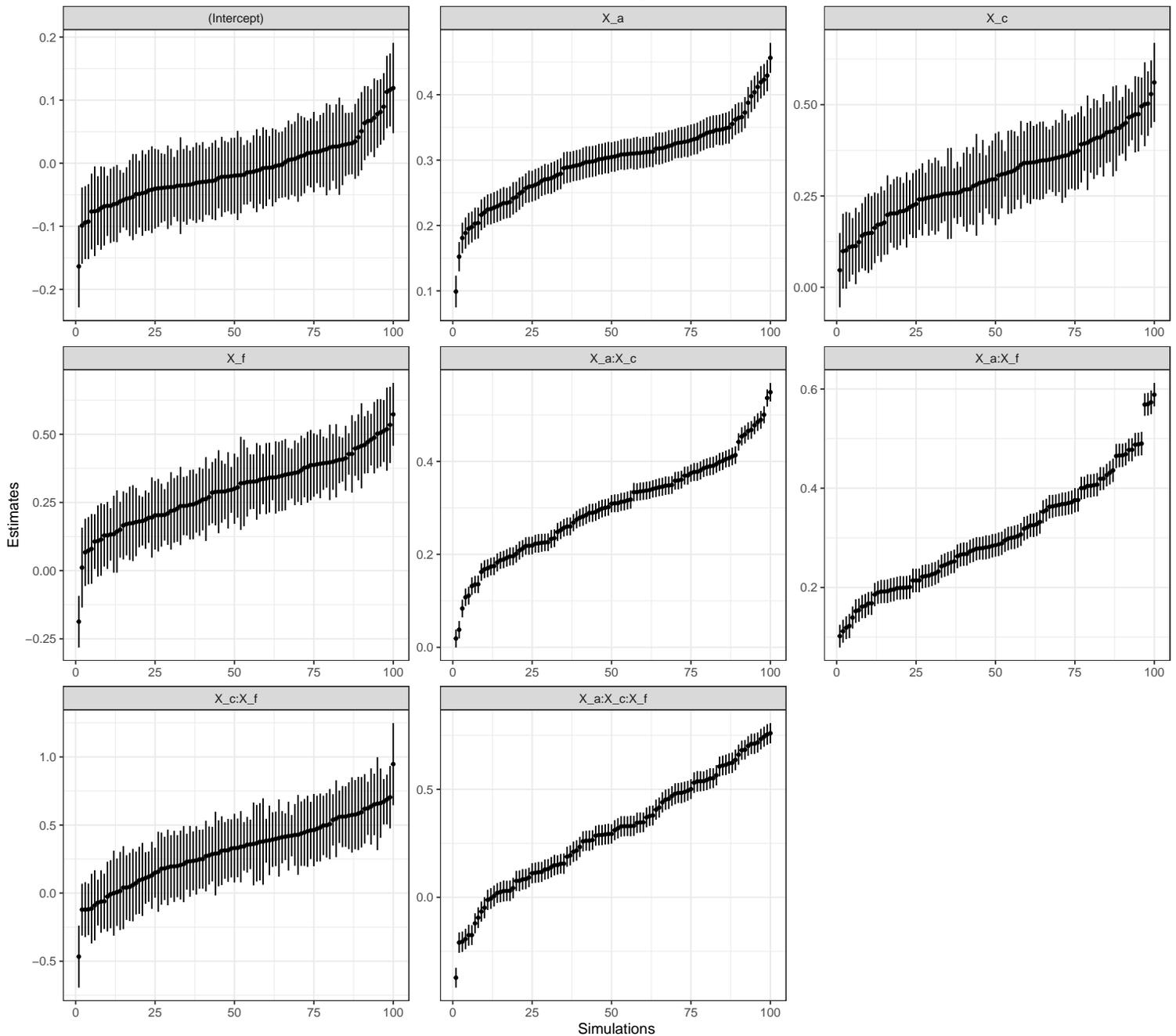
# read saved simulation data
sims_50_missing_trial_0.3 <- read_csv(filename_full_0.3_missing_trial, col_types = cols(
  # makes sure plots display in this order
  group = col_factor(ordered = TRUE),
  term = col_factor(ordered = TRUE)
))

fixed_missing_trial_plot <- sims_50_missing_trial_0.3 %>%
  filter(effect == "fixed") %>%
  ungroup() %>%
  arrange(term, estimate) %>%
  mutate(row = rep(seq(1:reps), 8)) %>%
  ggplot(aes(x = row, y = estimate, ymin = estimate-std.error, ymax = estimate+std.error)) +
  facet_wrap(~term, scales = "free") +
  geom_pointrange(fatten = 1/2) +
  ylab("Estimates") +
  xlab("Simulations") +
  ggtitle('Estimates of Fixed Effects for Missing Data with Trial Number, ef = 0.3') +
  theme_bw()

```

```
fixed_missing_trial_plot <- fixed_missing_trial_plot + theme(plot.title = element_text(hjust = 0.5, size=20))
fixed_missing_trial_plot
```

## Estimates of Fixed Effects for Missing Data with Trial Number, $ef = 0.3$

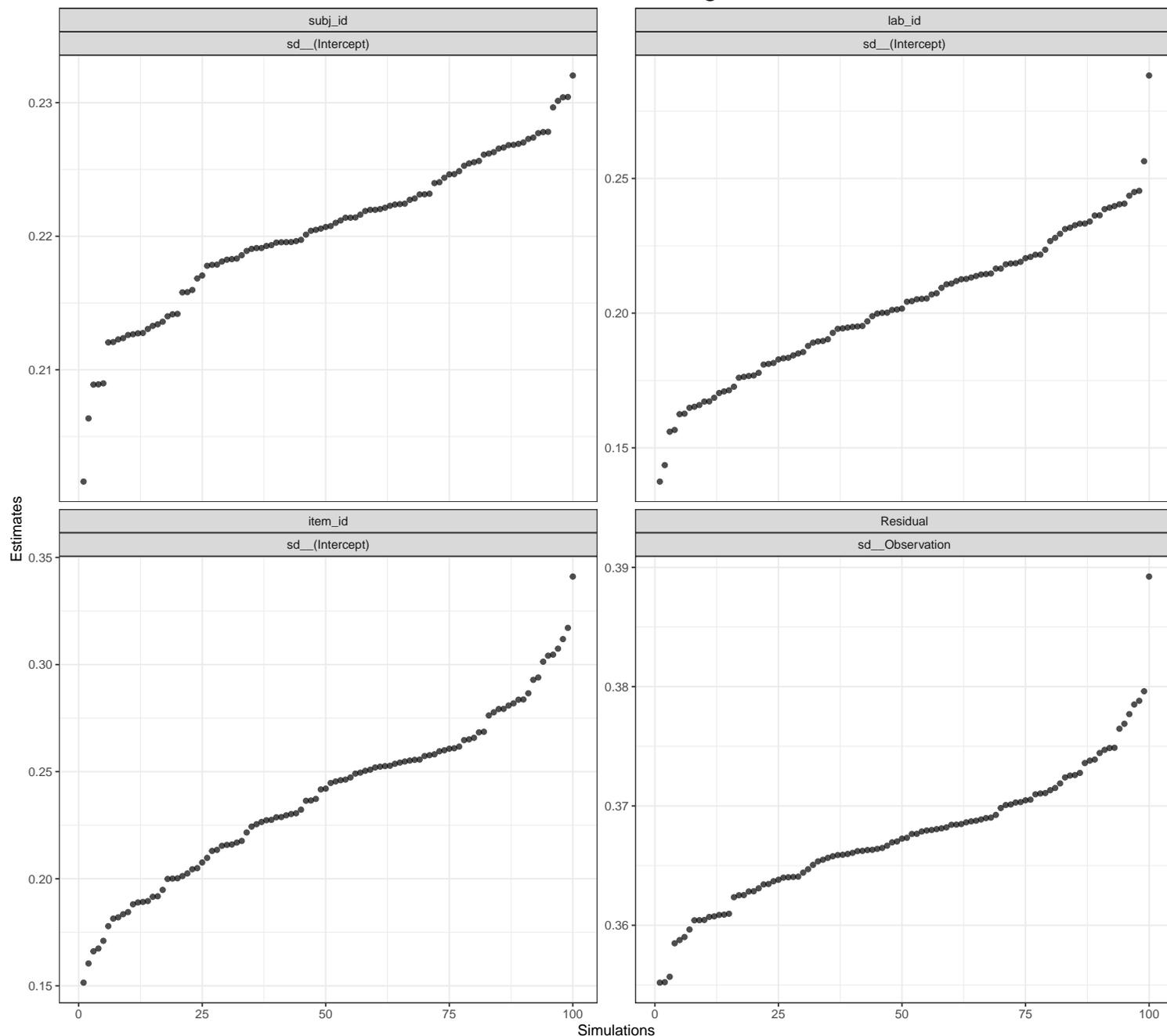


### 10.2.3 Visualise Estimates for Random Effects:

```
ran_missing_trial_plot <- sims_50_missing_trial_0.3 %>%
  filter(effect == "ran_pars") %>%
  ungroup() %>%
  arrange(group, term, estimate) %>%
  mutate(row = rep(seq(1:reps), 4)) %>%
  ggplot(aes(x = row, y = estimate)) + geom_point(alpha = 0.7) +
  facet_wrap(~group + term, scales = "free_y") + theme_bw() +
  ylab("Estimates") + xlab("Simulations") + ggtitle("Estimates of Random Effects for Missing Data with Trial, e
  theme_bw()
ran_missing_trial_plot <- ran_missing_trial_plot + theme(plot.title = element_text(hjust = 0.5,
```

```
size = 20))
ran_missing_trial_plot
```

### Estimates of Random Effects for Missing Data with Trial, $ef = 0.3$

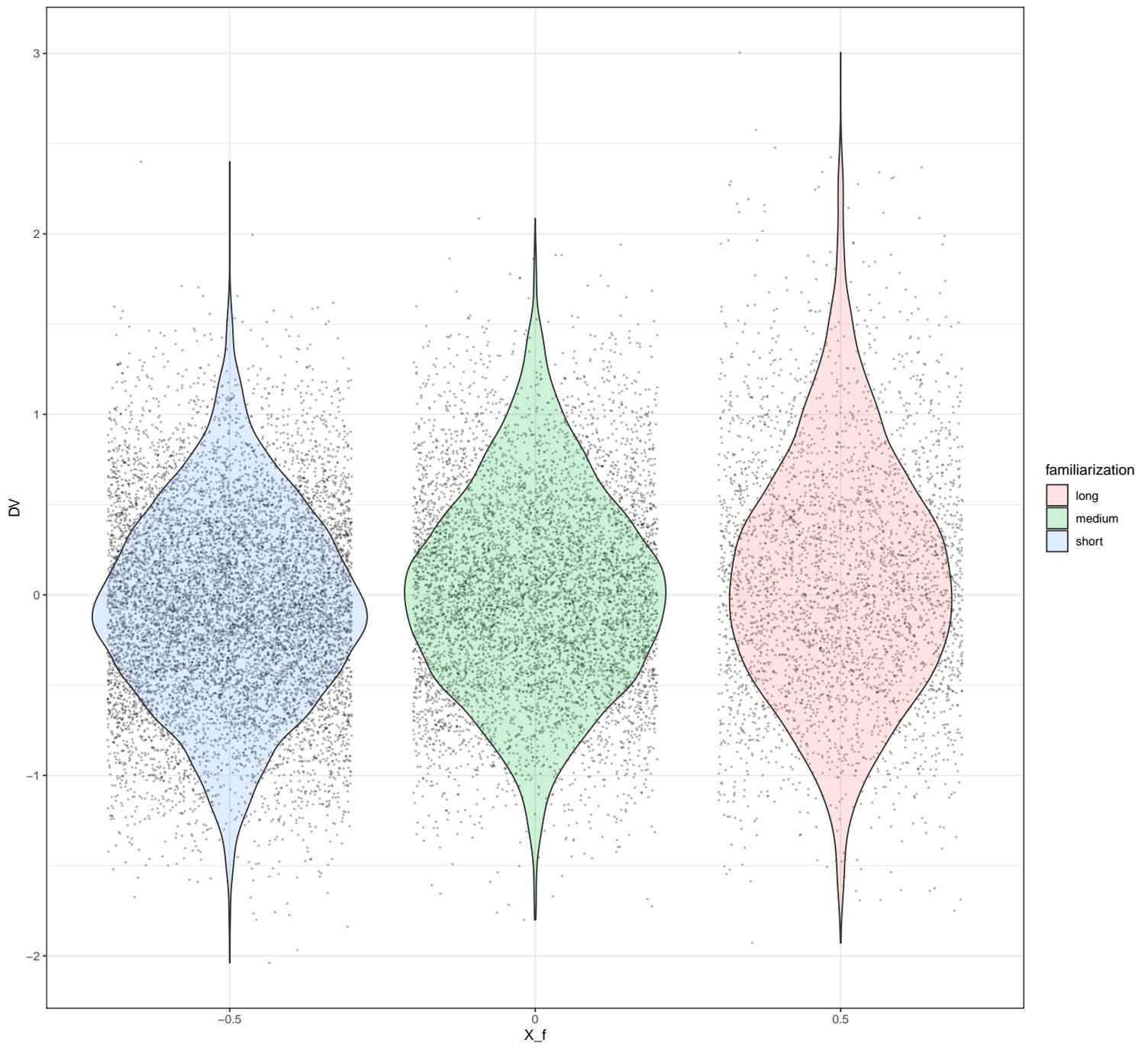


### 10.3 Simulation of missing data according to increasing familiarisation

```
# Proportion of missing data increases with
# familiarisation:
missing_samples_fam <- dat_sim %>%
  mutate(nas = rbinom(n(), 1, 0.95 - ifelse(X_f > -0.5, (X_f +
    0.5) * 0.6, 0))) %>%
  mutate(DV = ifelse(nas == 1, DV, NA))

missing_samples_fam %>%
  mutate(X_f = as.factor(X_f)) %>%
  drop_na() %>%
```

```
ggplot() + geom_point(aes(x = X_f, y = DV), alpha = 0.3,  
size = 0.2, position = "jitter") + geom_violin(aes(y = DV,  
x = X_f, fill = familiarization), alpha = 0.2) + theme_bw()
```



### 10.3.1 Simulation of models

```
# Number of simulations:  
reps <- 100  
  
# Simulation function:  
run_sims <- function(filename_full, ef) {  
  
  dat_sim <- my_sim_data(beta_c = ef,  
                        beta_f = ef,  
                        beta_a = ef,
```

```

        beta_ca = ef,
        beta_af = ef,
        beta_cf = ef,

        beta_cfa = ef)

missing_samples_fam <- dat_sim %>%
  mutate(nas = rbinom(n(), 1, 0.95 - ifelse(X_f > -0.5, (X_f+0.5)*0.6, 0))) %>%
  mutate(DV = ifelse(nas == 1, DV, NA))

mod_sim <- lmer(DV ~ 1 + X_a * X_c * X_f +
              (1 | subj_id) +
              (1 | lab_id) +
              (1 | item_id),
              data=missing_samples_fam)

sim_results <- broom.mixed::tidy(mod_sim)

# append the results to a file
append <- file.exists(filename_full)
write_csv(sim_results, filename_full, append = append)

# return the tidy table
sim_results
}

filename_full_0.3_missing_fam = 'sims/run_sims_0.3_fam_missing.csv'
start_time <- Sys.time()
sims <- purrr::map_df(1:reps, ~run_sims(filename_full = filename_full_0.3_missing_fam, ef = 0.3))
end_time <- Sys.time()
end_time - start_time

```

### 10.3.2 Visualise Estimates for Fixed Effects:

```

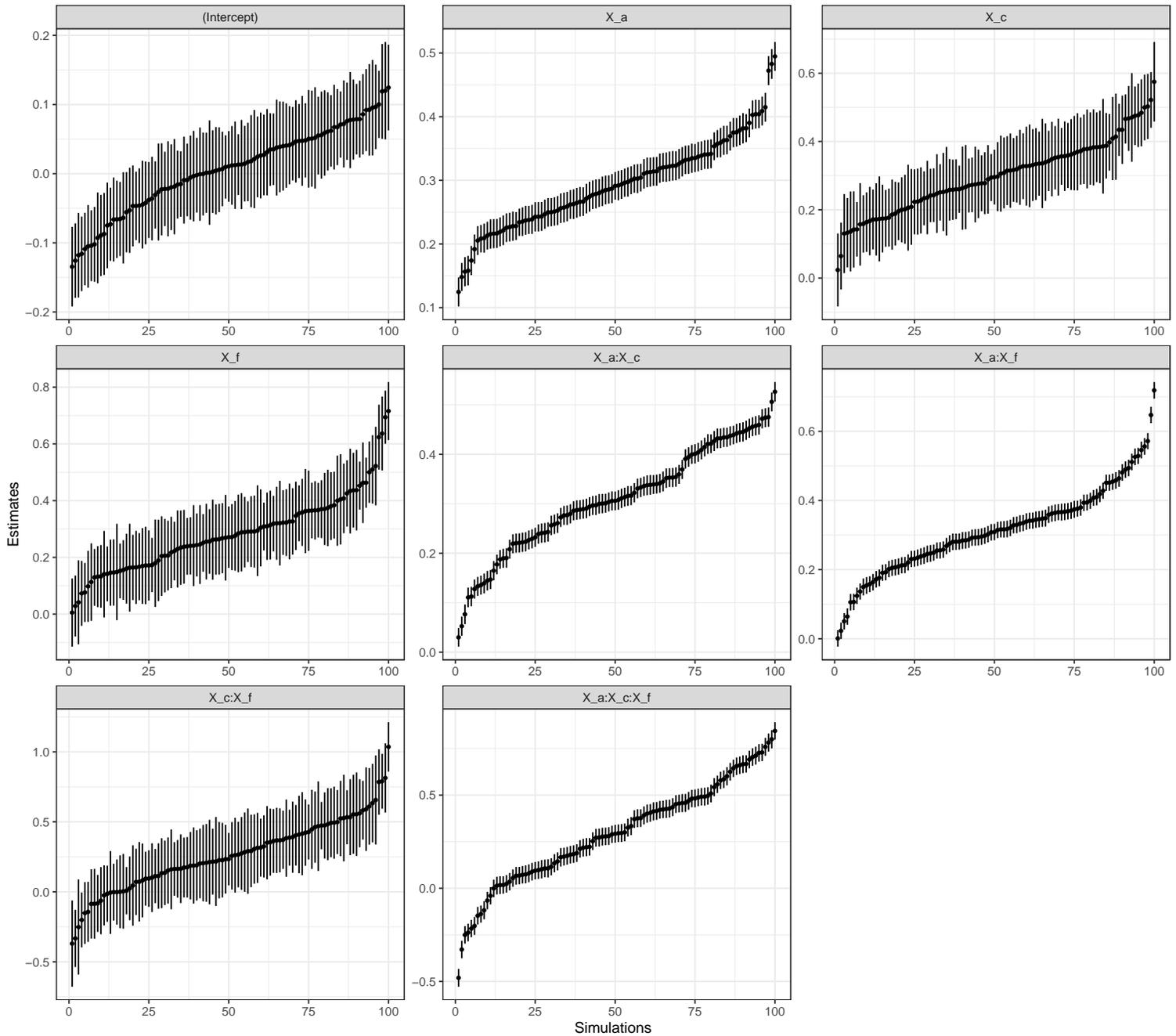
# read saved simulation data
sims_50_missing_fam_0.3 <- read_csv(filename_full_0.3_missing_fam, col_types = cols(
  # makes sure plots display in this order
  group = col_factor(ordered = TRUE),
  term = col_factor(ordered = TRUE)
))

fixed_missing_fam_plot <- sims_50_missing_fam_0.3 %>%
  filter(effect == "fixed") %>%
  ungroup() %>%
  arrange(term, estimate) %>%
  mutate(row = rep(seq(1:reps), 8)) %>%
  ggplot(aes(x = row, y = estimate, ymin = estimate-std.error, ymax = estimate+std.error)) +
  facet_wrap(~term, scales = "free") +
  geom_pointrange(fatten = 1/2) +
  ylab("Estimates") +
  xlab("Simulations") +
  ggtitle('Estimates of Fixed Effects for Missing Data with Familiarisation, ef = 0.3') +
  theme_bw()

fixed_missing_fam_plot <- fixed_missing_fam_plot + theme(plot.title = element_text(hjust = 0.5, size=20))
fixed_missing_fam_plot

```

## Estimates of Fixed Effects for Missing Data with Familiarisation, $ef = 0.3$



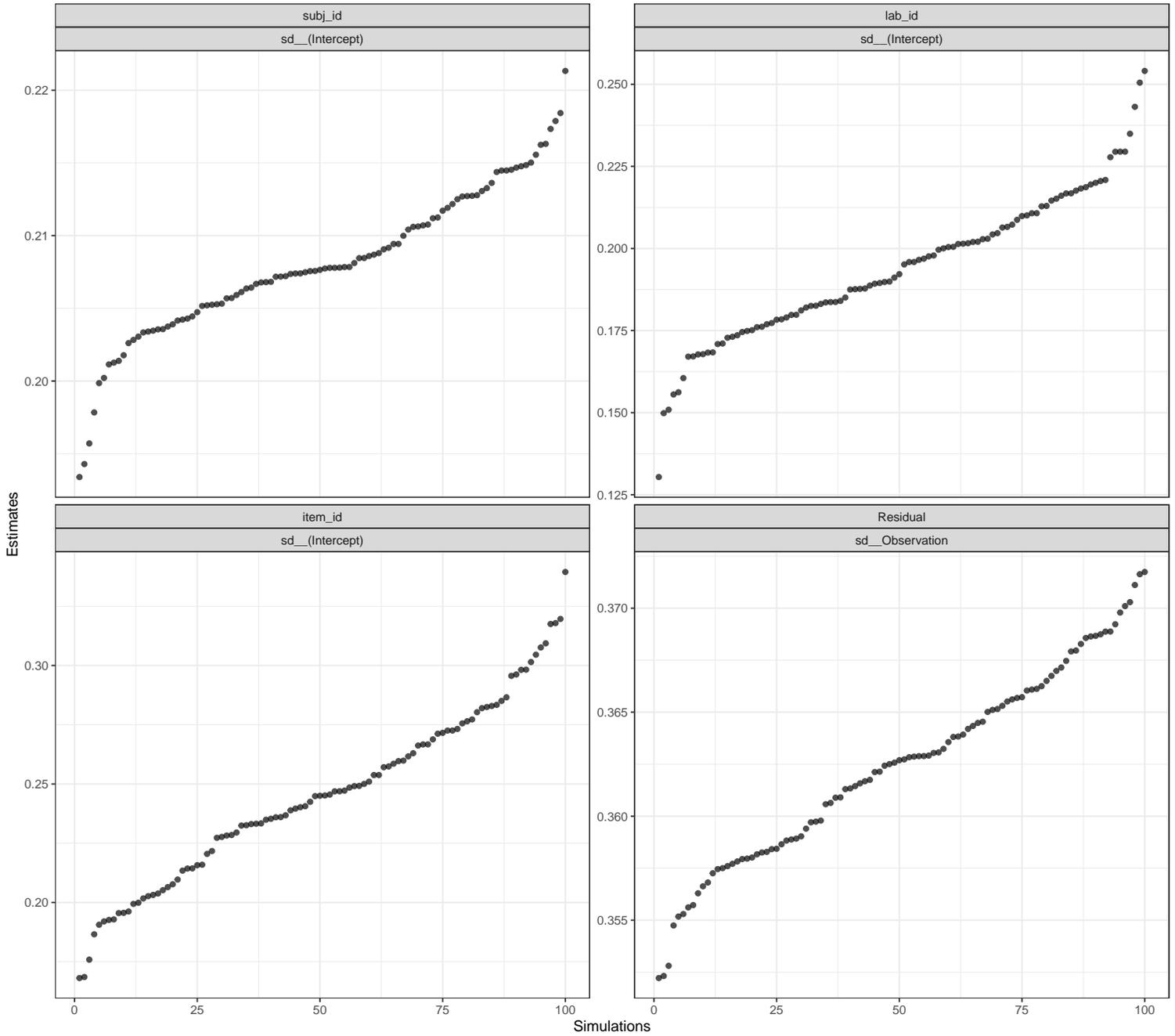
### Visualise Estimates for Random Effects:

```

ran_missing_fam_plot <- sims_50_missing_fam_0.3 %>%
  filter(effect == "ran_pars") %>%
  ungroup() %>%
  arrange(group, term, estimate) %>%
  mutate(row = rep(seq(1:reps), 4)) %>%
  ggplot(aes(x = row, y = estimate)) + geom_point(alpha = 0.7) +
  facet_wrap(~group + term, scales = "free_y") + theme_bw() +
  ylab("Estimates") + xlab("Simulations") + ggtitle("Estimates of Random Effects for Missing Data with Familiarisation") +
  theme_bw()
ran_missing_fam_plot <- ran_missing_fam_plot + theme(plot.title = element_text(hjust = 0.5,
  size = 20))
ran_missing_fam_plot

```

## Estimates of Random Effects for Missing Data with Familiarisation, $ef = 0.3$



## 11 Summary Statistics for Power Calculation with Data missing not completely at random

Table 9: Power for Simulations with Non-Random Missing Data and Varying Intercepts

term	missing with age	missing with trial	missing with fam
(Intercept)	0.07	0.01	0.06
X_a	1.00	1.00	1.00
X_c	0.79	0.79	0.78
X_f	0.66	0.62	0.58
X_a:X_c	0.98	0.99	0.99
X_a:X_f	0.94	1.00	0.98
X_c:X_f	0.22	0.24	0.12
X_a:X_c:X_f	0.81	0.85	0.85

Table 10: Bias for Simulations with Non-Random Missing Data and Varying Intercepts

term	age bias, ef = 0.3	trial bias, ef = 0.3	fam bias, ef = 0.3
(Intercept)	-0.010	0.020	-0.011
X_a	-0.008	-0.005	0.008
X_c	-0.003	-0.001	0.004
X_f	-0.019	-0.003	0.029
X_a:X_c	-0.001	-0.009	-0.007
X_a:X_f	0.005	0.014	-0.012
X_c:X_f	0.019	-0.031	0.055
X_a:X_c:X_f	0.022	-0.002	0.006

## 12 Logistic Regression Models

```

m_missing_age <- glmer(nas ~ X_f + X_a + X_c + trial_num + (1 |
  subj_id) + (1 | lab_id) + (1 | item_id), data = missing_samples_age,
  family = binomial)

summary(m_missing_age)

m_missing_fam <- glmer(nas ~ X_f + X_a + X_c + trial_num + (1 |
  subj_id) + (1 | lab_id) + (1 | item_id), data = missing_samples_fam,
  family = binomial)

summary(m_missing_fam)

m_missing_trial <- glmer(nas ~ X_f + X_a + X_c + trial_num +
  (1 | subj_id) + (1 | lab_id) + (1 | item_id), data = missing_samples_trial,
  family = binomial)

summary(m_missing_trial)

```