Same data, different analysts: variation in effect sizes due to analytical decisions in ecology and evolutionary biology.

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411 Abstract

412 Although variation in effect sizes and predicted values among studies of similar phenomena is 413 inevitable, such variation far exceeds what might be produced by sampling error alone. One possible 414 explanation for variation among results is differences among researchers in the decisions they make 415 regarding statistical analyses. A growing array of studies has explored this analytical variability in 416 different fields and has found substantial variability among results despite analysts having the same 417 data and research question. Many of these studies have been in the social sciences, but one small 418 'many analyst' study found similar variability in ecology. We expanded the scope of this prior work by implementing a large-scale empirical exploration of the variation in effect sizes and model 419 420 predictions generated by the analytical decisions of different researchers in ecology and evolutionary 421 biology. We used two unpublished datasets, one from evolutionary ecology (blue tit, Cyanistes 422 caeruleus, to compare sibling number and nestling growth) and one from conservation ecology 423 (Eucalyptus, to compare grass cover and tree seedling recruitment). The project leaders recruited 424 174 analyst teams, comprising 246 analysts, to investigate the answers to prespecified research 425 questions. Analyses conducted by these teams yielded 141 usable effects (compatible with our meta-426 analyses and with all necessary information provided) for the blue tit dataset, and 85 usable effects 427 for the Eucalyptus dataset. We found substantial heterogeneity among results for both datasets, 428 although the patterns of variation differed between them. For the blue tit analyses, the average 429 effect was convincingly negative, with less growth for nestlings living with more siblings, but there 430 was near continuous variation in effect size from large negative effects to effects near zero, and even 431 effects crossing the traditional threshold of statistical significance in the opposite direction. In 432 contrast, the average relationship between grass cover and *Eucalyptus* seedling number was only 433 slightly negative and not convincingly different from zero, and most effects ranged from weakly 434 negative to weakly positive, with about a third of effects crossing the traditional threshold of 435 significance in one direction or the other. However, there were also several striking outliers in 436 the Eucalyptus dataset, with effects far from zero. For both datasets, we found substantial variation in the variable selection and random effects structures among analyses, as well as in the ratings of 437 438 the analytical methods by peer reviewers, but we found no strong relationship between any of these 439 and deviation from the meta-analytic mean. In other words, analyses with results that were far from 440 the mean were no more or less likely to have dissimilar variable sets, use random effects in their 441 models, or receive poor peer reviews than those analyses that found results that were close to the 442 mean. The existence of substantial variability among analysis outcomes raises important questions 443 about how ecologists and evolutionary biologists should interpret published results, and how they 444 should conduct analyses in the future.

445 Introduction

446 One value of science derives from its production of replicable, and thus reliable, results. When we 447 repeat a study using the original methods, we should be able to expect a similar result. However, 448 perfect replicability is not a reasonable goal. Effect sizes will vary, and even reverse in sign, by chance 449 alone (Gelman and Weakliem 2009). Observed patterns can differ for other reasons as well. It could 450 be that we do not sufficiently understand the conditions that led to the original result so when we 451 seek to replicate it, the conditions differ due to some 'hidden moderator'. This hidden moderator 452 hypothesis is described by meta-analysts in ecology and evolutionary biology as 'true biological 453 heterogeneity' (Senior et al. 2016). This idea of true heterogeneity is popular in ecology and 454 evolutionary biology, and there are good reasons to expect it in the complex systems in which we

work (Shavit and Ellison 2017). However, despite similar expectations in psychology, recent evidence 455 456 in that discipline contradicts the hypothesis that moderators are common obstacles to replicability, 457 as variability in results in a large 'many labs' collaboration was mostly unrelated to commonly 458 hypothesized moderators such as the conditions under which the studies were administered (Klein et 459 al. 2018). Another possible explanation for variation in effect sizes is that researchers often present 460 biased samples of results, thus reducing the likelihood that later studies will produce similar effect sizes (Open Science Collaboration 2015; Parker et al. 2016; Forstmeier, Wagenmakers, and Parker 461 462 2017; Fraser et al. 2018; Parker and Yang 2023). It also may be that although researchers did 463 successfully replicate the conditions, the experiment, and measured variables, analytical decisions 464 differed sufficiently among studies to create divergent results (Simonsohn, Simmons, and Nelson 465 2015; Silberzahn et al. 2018).

466 Analytical decisions vary among studies because researchers have many options. Researchers need 467 to decide how to exclude possibly anomalous or unreliable data, how to construct variables, which 468 variables to include in their models, and which statistical methods to use. Depending on the dataset, 469 this short list of choices could encompass thousands or millions of possible alternative 470 specifications (Simonsohn, Simmons, and Nelson 2015). However, researchers making these 471 decisions presumably do so with the goal of doing the best possible analysis, or at least the best 472 analysis within their current skill set. Thus, it seems likely that some specification options are more 473 probable than others, possibly because they have previously been shown (or claimed) to be better, 474 or because they are more well known. Of course, some of these different analyses (maybe many of 475 them) may be equally valid alternatives. Regardless, on probably any topic in ecology and 476 evolutionary biology, we can encounter differences in choices of data analysis. The extent of these 477 differences in analyses and the degree to which these differences influence the outcomes of analyses 478 and therefore studies' conclusions are important empirical questions. These questions are especially 479 important given that many papers draw conclusions after applying a single method, or even a single 480 statistical model, to analyze a dataset.

481 The possibility that different analytical choices could lead to different outcomes has long been 482 recognized (Gelman and Loken 2013), and various efforts to address this possibility have been 483 pursued in the literature. For instance, one common method in ecology and evolutionary biology 484 involves creating a set of candidate models, each consisting of a different (though often similar) set 485 of predictor variables, and then, for the predictor variable of interest, averaging the slope across all 486 models (i.e. model averaging) (Burnham and Anderson 2002; Grueber et al. 2011). This method 487 reduces the chance that a conclusion is contingent upon a single model specification, though use and 488 interpretation of this method is not without challenges (Grueber et al. 2011). Further, the models 489 compared to each other typically differ only in the inclusion or exclusion of certain predictor 490 variables and not in other important ways, such as methods of parameter estimation. More explicit 491 examination of outcomes of differences in model structure, model type, data exclusion, or other 492 analytical choices can be implemented through sensitivity analyses (e.g., Noble et al. 2017). 493 Sensitivity analyses, however, are typically rather narrow in scope, and are designed to assess the 494 sensitivity of analytical outcomes to a particular analytical choice rather than to a large universe of 495 choices. Recently, however, analysts in the social sciences have proposed extremely thorough 496 sensitivity analysis, including 'multiverse analysis' (Steegen et al. 2016) and the 'specification 497 curve' (Simonsohn, Simmons, and Nelson 2015), as a means of increasing the reliability of results. 498 With these methods, researchers identify relevant decision points encountered during analysis and 499 conduct the analysis many times to incorporate many plausible decisions made at each of these 500 points. The study's conclusions are then based on a broad set of the possible analyses and so allow 501 the analyst to distinguish between robust conclusions and those that are highly contingent on

502 particular model specifications. These are useful outcomes, but specifying a universe of possible 503 modelling decisions is not a trivial undertaking. Further, the analyst's knowledge and biases will 504 influence decisions about the boundaries of that universe, and so there will always be room for 505 disagreement among analysts about what to include. Including more specifications is not necessarily 506 better. Some analytical decisions are better justified than others, and including biologically 507 implausible specifications may undermine this process. Regardless, these powerful methods have yet 508 to be adopted, and even the more limited forms of sensitivity analyses are not particularly 509 widespread. Most studies publish a small set of analyses and so the existing literature does not 510 provide much insight into the degree to which published results are contingent on analytical 511 decisions.

Despite the potential major impacts of analytical decisions on variance in results, the outcomes of 512 513 different individuals' data analysis choices have only recently begun to receive much empirical 514 attention. The only formal exploration of this that we were aware of when we submitted our Stage 1 515 manuscript were (1) an analysis in social science that asked whether male professional football 516 (soccer) players with darker skin tone were more likely to be issued red cards (ejection from the 517 game for rule violation) than players with lighter skin tone (Silberzahn et al. 2018) and (2) an analysis 518 in neuroimaging which evaluated nine separate hypotheses involving the neurological responses 519 detected with fMRI in 108 participants divided between two treatments in a decision making 520 task (Botvinik-Nezer et al. 2020). Several others have been published since (e.g., Huntington-Klein et 521 al. 2021; Schweinsberg et al. 2021; Breznau et al. 2022; Coretta et al. 2023), and we recently learned 522 of an earlier small study in ecology (Stanton-Geddes, Freitas, and Sales Dambros 2014). In the red 523 card study, 29 teams designed and implemented analyses of a dataset provided by the study 524 coordinators (Silberzahn et al. 2018). Analyses were peer reviewed (results blind) by at least two 525 other participating analysts; a level of scrutiny consistent with standard pre-publication peer review. 526 Among the final 29 analyses, odds-ratios varied from 0.89 to 2.93, meaning point estimates varied from having players with lighter skin tones receive more red cards (odds ratio < 1) to a strong effect 527 528 of players with darker skin tones receiving more red cards (odds ratio > 1). Twenty of the 29 teams 529 found a statistically-significant effect in the predicted direction of players with darker skin tones 530 being issued more red cards. This degree of variation in peer-reviewed analyses from identical data is 531 striking, but the generality of this finding has only just begun to be formally 532 investigated (e.g., Huntington-Klein et al. 2021; Schweinsberg et al. 2021; Breznau et al.

533 2022; Coretta et al. 2023).

534 In the neuroimaging study, 70 teams evaluated each of the nine different hypotheses with the 535 available fMRI data (Botvinik-Nezer et al. 2020). These 70 teams followed a divergent set of 536 workflows that produced a wide range of results. The rate of reporting of statistically significant 537 support for the nine hypotheses ranged from 21% to 84%, and for each hypothesis on average, 20% 538 of research teams observed effects that differed substantially from the majority of other teams. 539 Some of the variability in results among studies could be explained by analytical decisions such as 540 choice of software package, smoothing function, and parametric versus non-parametric corrections 541 for multiple comparisons. However, substantial variability among analyses remained unexplained, 542 and presumably emerged from the many different decisions each analyst made in their long 543 workflows. Such variability in results among analyses from this dataset and from the very different 544 red-card dataset suggests that sensitivity of analytical outcome to analytical choices may characterize 545 many distinct fields, as several more recent many-analyst studies also suggest (Huntington-Klein et al. 546 2021; Schweinsberg et al. 2021; Breznau et al. 2022).

547 To further develop the empirical understanding of the effects of analytical decisions on study 548 outcomes, we chose to estimate the extent to which researchers' data analysis choices drive 549 differences in effect sizes, model predictions, and qualitative conclusions in ecology and evolutionary 550 biology. This is an important extension of the meta-research agenda of evaluating factors influencing 551 replicability in ecology, evolutionary biology, and beyond (Fidler et al. 2017). To examine the effects 552 of analytical decisions, we used two different datasets and recruited researchers to analyze one or the other of these datasets to answer a question we defined. The first question was "To what extent 553 554 is the growth of nestling blue tits (Cyanistes caeruleus) influenced by competition with siblings?" To 555 answer this question, we provided a dataset that includes brood size manipulations from 332 broods 556 conducted over three years at Wytham Wood, UK. The second question was "How does grass cover 557 influence Eucalyptus spp. seedling recruitment?" For this question, analysts used a dataset that 558 includes, among other variables, number of seedlings in different size classes, percentage cover of 559 different life forms, tree canopy cover, and distance from canopy edge from 351 guadrats spread 560 among 18 sites in Victoria, Australia.

561 We explored the impacts of data analysts' choices with descriptive statistics and with a series of tests 562 to attempt to explain the variation among effect sizes and predicted values of the dependent variable 563 produced by the different analysis teams for both datasets separately. To describe the variability, we 564 present forest plots of the standardized effect sizes and predicted values produced by each of the analysis teams, estimate heterogeneity (both absolute, τ^2 , and proportional, l^2) in effect size and 565 566 predicted values among the results produced by these different teams, and calculate a similarity index that quantifies variability among the predictor variables selected for the different statistical 567 568 models constructed by the different analysis teams. These descriptive statistics provide the first 569 estimates of the extent to which explanatory statistical models and their outcomes in ecology and 570 evolutionary biology vary based on the decisions of different data analysts. We then quantified the 571 degree to which the variability in effect size and predicted values could be explained by (1) variation 572 in the quality of analyses as rated by peer reviewers and (2) the similarity of the choices of predictor 573 variables between individual analyses.

574 Methods

575 This project involved a series of steps (1-6) that began with identifying datasets for analyses and 576 continued through recruiting independent groups of scientists to analyze the data, allowing the 577 scientists to analyze the data as they saw fit, generating peer review ratings of the analyses (based 578 on methods, not results), evaluating the variation in effects among the different analyses, and 579 producing the final manuscript.

580 Step 1: Select datasets

581 We used two previously unpublished datasets, one from evolutionary ecology and the other from582 ecology and conservation.

583 Evolutionary ecology

584 Our evolutionary ecology dataset is relevant to a sub-discipline of life-history research which focuses 585 on identifying costs and trade-offs associated with different phenotypic conditions. These data were 586 derived from a brood-size manipulation experiment imposed on wild birds nesting in boxes provided 587 by researchers in an intensively studied population. Understanding how the growth of nestlings is 588 influenced by the numbers of siblings in the nest can give researchers insights into factors such as the evolution of clutch size, determination of provisioning rates by parents, and optimal levels of sibling
competition (Vander Werf 1992; DeKogel 1997; Royle et al. 1999; Verhulst, Holveck, and Riebel
2006; Nicolaus et al. 2009). Data analysts were provided this dataset and instructed to answer the
following question: "To what extent is the growth of nestling blue tits (*Cyanistes caeruleus*)
influenced by competition with siblings?"

594 Researchers conducted brood size manipulations and population monitoring of blue tits at Wytham 595 Wood, a 380 ha woodland in Oxfordshire, U.K (1º 20'W, 51º 47'N). Researchers regularly checked 596 approximately 1100 artificial nest boxes at the site and monitored the 330 to 450 blue tit pairs 597 occupying those boxes in 2001-2003 during the experiment. Nearly all birds made only one breeding 598 attempt during the April to June study period in a given year. At each blue tit nest, researchers 599 recorded the date the first egg appeared, clutch size, and hatching date. For all chicks alive at age 14 600 days, researchers measured mass and tarsus length and fitted a uniquely numbered, British Trust for 601 Ornithology (BTO) aluminium leg ring. Researchers attempted to capture all adults at their nests 602 between day 6 and day 14 of the chick-rearing period. For these captured adults, researchers 603 measured mass, tarsus length, and wing length and fitted a uniquely numbered BTO leg ring. During 604 the 2001-2003 breeding seasons, researchers manipulated brood sizes using cross fostering. They 605 matched broods for hatching date and brood size and moved chicks between these paired nests one 606 or two days after hatching. They sought to either enlarge or reduce all manipulated broods by 607 approximately one fourth. To control for effects of being moved, each reduced brood had a portion 608 of its brood replaced by chicks from the paired increased brood, and vice versa. Net manipulations 609 varied from plus or minus four chicks in broods of 12 to 16 to plus or minus one chick in broods of 4 610 or 5. Researchers left approximately one third of all broods unmanipulated. These unmanipulated 611 broods were not selected systematically to match manipulated broods in clutch size or laying date. 612 We have mass and tarsus length data from 3720 individual chicks divided among 167 experimentally 613 enlarged broods, 165 experimentally reduced broods, and 120 unmanipulated broods. The full list of 614 variables included in the dataset is publicly available (https://osf.io/hdv8m), along with the data 615 (https://osf.io/qjzby).

616 Ecology and conservation

Additional Explanation:

Shortly after beginning to recruit analysts, several analysts noted a small set of related errors in the blue tit dataset. We corrected the errors, replaced the dataset on our OSF site, and emailed the analysts on 19 April 2020 to instruct them to use the revised data. The email to analysts is available here (<u>https://osf.io/4h53z</u>). The errors are explained in that email.

617 Our ecology and conservation dataset is relevant to a sub-discipline of conservation research which 618 focuses on investigating how best to revegetate private land in agricultural landscapes. These data 619 were collected on private land under the Bush Returns program, an incentive system where 620 participants entered into a contract with the Goulburn Broken Catchment Management Authority 621 and received annual payments if they executed predetermined restoration activities. This particular 622 dataset is based on a passive regeneration initiative, where livestock grazing was removed from the 623 property in the hopes that the *Eucalyptus* spp. overstorey would regenerate without active (and 624 expensive) planting. Analyses of some related data have been published (Miles 2008; Vesk et al. 625 2016) but those analyses do not address the question analysts answered in our study. Data analysts 626 were provided this dataset and instructed to answer the following question: "How does grass cover 627 influence Eucalyptus spp. seedling recruitment?".

628 Researchers conducted three rounds of surveys at 18 sites across the Goulburn Broken catchment in 629 northern Victoria, Australia in winter and spring 2006 and autumn 2007. In each survey period, a 630 different set of 15 x 15 m quadrats were randomly allocated across each site within 60 m of existing 631 tree canopies. The number of quadrats at each site depended on the size of the site, ranging from 632 four at smaller sites to 11 at larger sites. The total number of quadrats surveyed across all sites and 633 seasons was 351. The number of *Eucalyptus* spp. seedlings was recorded in each quadrat along with 634 information on the GPS location, aspect, tree canopy cover, distance to tree canopy, and position in 635 the landscape. Ground layer plant species composition was recorded in three 0.5 x 0.5 m sub-636 quadrats within each quadrat. Subjective cover estimates of each species as well as bare ground, 637 litter, rock and moss/lichen/soil crusts were recorded. Subsequently, this was augmented with 638 information about the precipitation and solar radiation at each GPS location. The full list of variables 639 included in the dataset is publicly available (https://osf.io/r5gbn), along with the data 640 (https://osf.io/qz5cu).

641

642 Step 2: Recruitment and initial survey of analysts

643 The lead team (TP, HF, SN, EG, SG, PV, DH, FF) created a publicly available document providing a

Preregistration Deviation:

Due to the large number of recruited analysts and reviewers and the anticipated challenges of receiving and integrating feedback from so many authors, we limited analyst and reviewer participation in the production of the final manuscript to an invitation to call attention to serious problems with the manuscript draft.

644 general description of the project (https://osf.io/mn5aj/). The project was advertised at conferences, 645 via Twitter, using mailing lists for ecological societies (including Ecolog, Evoldir, and lists for the 646 Environmental Decisions Group, and Transparency in Ecology and Evolution), and via word of mouth. 647 The target population was active ecology, conservation, or evolutionary biology researchers with a 648 graduate degree (or currently studying for a graduate degree) in a relevant discipline. Researchers 649 could choose to work independently or in a small team. For the sake of simplicity, we refer to these 650 as 'analysis teams' though some comprised one individual. We aimed for a minimum of 12 analysis 651 teams independently evaluating each dataset (see sample size justification below). We 652 simultaneously recruited volunteers to peer review the analyses conducted by the other volunteers 653 through the same channels. Our goal was to recruit a similar number of peer reviewers and analysts, 654 and to ask each peer reviewer to review a minimum of four analyses. If we were unable to recruit at 655 least half the number of reviewers as analysis teams, we planned to ask analysts to serve also as 656 reviewers (after they had completed their analyses), but this was unnecessary. Therefore, no data 657 analysts peer reviewed analyses of the dataset they had analyzed. All analysts and reviewers were 658 offered the opportunity to share co-authorship on this manuscript and we planned to invite them to 659 participate in the collaborative process of producing the final manuscript. All analysts signed 660 [digitally] a consent (ethics) document (https://osf.io/xyp68/) approved by the Whitman College 661 Institutional Review Board prior to being allowed to participate.

- 662 We identified our minimum number of analysts per dataset by considering the number of effects
- 663 needed in a meta-analysis to generate an estimate of heterogeneity (τ^2) with a 95% confidence 664 interval that does not encompass zero. This minimum sample size is invariant regardless of τ^2 . This is
- 665 because the same t-statistic value will be obtained by the same sample size regardless of variance

666 (τ^2). We see this by first examining the formula for the standard error, SE for variance, (τ^2) or (SE τ^2)

667 assuming normality in an underlying distribution of effect sizes (Knight 2000):

$$\mathsf{568}\qquad\qquad \mathsf{SE}(\tau^2) = \sqrt{\frac{2\tau^4}{n-1}}$$

and then rearranging the above formula to show how the t-statistic is independent of τ^2 , as seen below.

$$t = \frac{\tau^2}{\mathsf{SE}(\tau^2)} = \sqrt{\frac{n-1}{2}}$$

672 We then find a minimum n = 12 according to this formula.

673 Step 3: Primary data analyses

674 Analysis teams registered and answered a demographic and expertise survey (https://osf.io/seqzy/). 675 We then provided them with the dataset of their choice and requested that they answer a specific 676 research question. For the evolutionary ecology dataset that question was "To what extent is the 677 growth of nestling blue tits (Cyanistes caeruleus) influenced by competition with siblings?" and for 678 the conservation ecology dataset it was "How does grass cover influence Eucalyptus spp. seedling 679 recruitment?" Once their analysis was complete, they answered a structured survey 680 (https://osf.io/neyc7/), providing analysis technique, explanations of their analytical choices, 681 quantitative results, and a statement describing their conclusions. They also were asked to upload 682 their analysis files (including the dataset as they formatted it for analysis and their analysis code [if

683 applicable]) and a detailed journal-ready statistical methods section.

684

Additional Information:

As is common in many studies in ecology and evolutionary biology, the datasets we provided contained many variables, and the research questions we provided could be addressed by our datasets in many different ways. For instance, volunteer analysts had to choose the dependent (response) variable and the independent variable, and make numerous other decisions about which variables and data to use and how to structure their model.

685

Preregistration Deviation:

We originally planned to have analysts complete a single survey (<u>https://osf.io/neyc7/</u>), but after we evaluated the results of that survey, we realized we would need a second survey (<u>https://osf.io/8w3v5/</u>) to adequately collect the information we needed to evaluate heterogeneity of results (step 5). We provided a set of detailed instructions with the follow-up survey, and these instructions are publicly available and can be found within the following files (blue tit: <u>https://osf.io/kr2g9</u>, *Eucalyptus*: <u>https://osf.io/dfvym</u>).

686 Step 4: Peer reviews of analyses

687 At minimum, each analysis was evaluated by four different reviewers, and each volunteer peer 688 reviewer was randomly assigned methods sections from at least four analyst teams (the exact 689 number varied). Each peer reviewer registered and answered a demographic and expertise survey 690 identical to that asked of the analysts, except we did not ask about 'team name' since reviewers did 691 not work in teams. Reviewers evaluated the methods of each of their assigned analyses one at a time 692 in a sequence determined by the project leaders. We systematically assigned the sequence so that, if 693 possible, each analysis was allocated to each position in the sequence for at least one reviewer. For 694 instance, if each reviewer were assigned four analyses to review, then each analysis would be the 695 first analysis assigned to at least one reviewer, the second analysis assigned to another reviewer, the 696 third analysis assigned to yet another reviewer, and the fourth analysis assigned to a fourth reviewer. 697 Balancing the order in which reviewers saw the analyses controls for order effects, e.g. a reviewer 698 might be less critical of the first methods section they read than the last.

699The process for a single reviewer was as follows. First, the reviewer received a description of the700methods of a single analysis. This included the narrative methods section, the analysis team's701answers to our survey questions regarding their methods, including analysis code, and the dataset.702The reviewer was then asked, in an online survey (https://osf.io/4t36u/), to rate that analysis on a703scale of 0-100 based on this prompt: "Rate the overall appropriateness of this analysis to answer the704research question (*one of the two research questions inserted here*) with the available data. To help705you calibrate your rating, please consider the following guidelines:

- 100. A perfect analysis with no conceivable improvements from the reviewer
- 75. An imperfect analysis but the needed changes are unlikely to dramatically alter outcomes
- 50. A flawed analysis likely to produce either an unreliable estimate of the relationship or an
 over-precise estimate of uncertainty
- 25. A flawed analysis likely to produce an unreliable estimate of the relationship and an over precise estimate of uncertainty
- 0. A dangerously misleading analysis, certain to produce both an estimate that is wrong and
 a substantially over-precise estimate of uncertainty that places undue confidence in the
 incorrect estimate.
- *Please note that these values are meant to calibrate your ratings. We welcome ratings of anynumber between 0 and 100.

717 After providing this rating, the reviewer was presented with this prompt, in multiple-choice format: 718 "Would the analytical methods presented produce an analysis that is (a) publishable as is, (b) 719 publishable with minor revision, (c) publishable with major revision, (d) deeply flawed and 720 unpublishable?" The reviewer was then provided with a series of text boxes and the following 721 prompts: "Please explain your ratings of this analysis. Please evaluate the choice of statistical analysis 722 type. Please evaluate the process of choosing variables for and structuring the statistical model. 723 Please evaluate the suitability of the variables included in (or excluded from) the statistical model. Please evaluate the suitability of the structure of the statistical model. Please evaluate choices to 724 725 exclude or not exclude subsets of the data. Please evaluate any choices to transform data (or, if there 726 were no transformations, but you think there should have been, please discuss that choice)." After 727 submitting this review, a methods section from a second analysis was then made available to the 728 reviewer. This same sequence was followed until all analyses allocated to a given reviewer were 729 provided and reviewed. After providing the final review, the reviewer was simultaneously provided 730 with all four (or more) methods sections the reviewer had just completed reviewing, the option to

- revise their original ratings, and a text box to provide an explanation. The invitation to revise the
- original ratings was as follows: "If, now that you have seen all the analyses you are reviewing, you
- vish to revise your ratings of any of these analyses, you may do so now." The text box was prefaced
- 734 with this prompt: "Please explain your choice to revise (or not to revise) your ratings."
- 735

Additional Information: unregistered analysis

To determine how consistent peer reviewers were in their ratings, we assessed inter-rater reliability among reviewers for both the categorical and quantitative ratings combining blue tit and *Eucalyptus* data using Krippendorff's alpha for ordinal and continuous data respectively. This provides a value that is between -1 (total disagreement between reviewers) and 1 (total agreement between reviewers).

736 Step 5: Evaluate variation

737



Figure 1: Schematic of research process showing recruited analyst and reviewer contributions in orange and core team contributions in blue. Items that are crossed out were preregistered but could not be conducted. Items with a greyed background were added as exploratory analyses after preregistration.

739 The lead team conducted the analyses outlined in this section. We described the variation in model 740 specification in several ways. We calculated summary statistics describing variation among analyses, 741 including mean, SD, and range of number of variables per model included as fixed effects, the 742 number of interaction terms, the number of random effects, and the mean, SD, and range of sample 743 sizes. We also present the number of analyses in which each variable was included. We summarized 744 the variability in standardized effect sizes and predicted values of dependent variables among the 745 individual analyses using standard random effects meta-analytic techniques. First, we derived 746 standardized effect sizes from each individual analysis. We did this for all linear models or 747 generalized linear models by converting the t value and the degree of freedom (df) associated with 748 regression coefficients (e.g. the effect of the number of siblings [predictor] on growth [response] or 749 the effect of grass cover [predictor] on seedling recruitment [response]) to the correlation 750 coefficient, r, using the following:

$$r = \frac{t^2}{(t^2 + df)}$$

752 This formula can only be applied if t and df values originate from linear or generalized linear models 753 [GLMs; Nakagawa and Cuthill (2007)]. If, instead, linear mixed-effects models (LMMs) or generalized 754 linear mixed-effects models (GLMMs) were used by a given analysis, the exact df cannot be 755 estimated. However, adjusted df can be estimated, for example, using the Satterthwaite 756 approximation of df, df_s , [note that SAS uses this approximation to obtain df for LMMs and 757 GLMMs; Luke (2017)]. For analyses using either LMMs or GLMMs that do not produce df_s we 758 planned to obtain df_s by rerunning the same (G)LMMs using the lmer() or glmer() function in 759 the ImerTest package in R (Kuznetsova, Brockhoff, and Christensen 2017; R Core Team 2024).

760

Preregistration Deviation:

Rather than re-run these analyses ourselves, we sent a follow-up survey (referenced above under "Primary data analyses") to analysts and asked them to follow our instructions for producing this information. The instructions are publicly available and can be found within the following files (blue tit: https://osf.io/kr2g9, Eucalyptus: https://osf.io/dfvym).

761 We then used the *t* values and df_s from the models to obtain *r* as per the formula above. All *r* and 762 accompanying df (or df_s) were converted to Fisher's Z_r .

$$Z_r = \frac{1}{2} \ln\left(\frac{1+r}{1-r}\right)$$

764 and its sampling variance; 1/(n-3) where n=df+1. Any analyses from which we could not derive a 765 signed Z_{r_i} for instance one with a quadratic function in which the slope changed sign, were 766 considered unusable for analyses of Z_r . We expected such analyses would be rare. In fact, most 767 submitted analyses excluded from our meta-analysis of Z_r were excluded because of a lack of 768 sufficient information provided by the analyst team rather than due to the use of effects that could 769 not be converted to Z_r . Regardless, as we describe below, we generated a second set of standardized 770 effects (predicted values) that could (in principle) be derived from any explanatory model produced 771 by these data.

Besides Z_r, which describes the strength of a relationship based on the amount of variation in a
 dependent variable explained by variation in an independent variable, we also examined differences

in the shape of the relationship between the independent and dependent variables. To accomplish
this, we derived a point estimate (out-of-sample predicted value) for the dependent variable of
interest for each of three values of our primary independent variable. We originally described these
three values as associated with the 25th percentile, median, and 75th percentile of the independent
variable and any covariates.

779

Preregistration Deviation:

The original description of the out-of-sample specifications did not account for the facts that (a) some variables are not distributed in a way that allowed division in percentiles and that (b) variables could be either positively or negatively correlated with the dependent variable. We provide a more thorough description here:

We derived three point-estimates (out-of-sample predicted values) for the dependent variable of interest; one for each of three values of our primary independent variable that we specified. We also specified values for all other variables that could have been included as independent variables in analysts' models so that we could derive the predicted values from a fully specified version of any model produced by analysts. For all potential independent variables, we selected three values or categories. Of the three we selected, one was associated with small, one with intermediate, and one with large values of one typical dependent variable (day 14 chick weight for the blue tit data and total number of seedlings for the Eucalyptus data; analysts could select other variables as their dependent variable, but the others typically correlated with the two identified here). For continuous variables, this means we identified the 25th percentile, median, and 75th percentile and, if the slope of the linear relationship between this variable and the typical dependent variable was positive, we left the quartiles ordered as is. If, instead, the slope was negative, we reversed the order of the independent variable quartiles so that the 'lower' guartile value was the one associated with the lower value for the dependent variable. In the case of categorical variables, we identified categories associated with the 25th percentile, median, and 75th percentile values of the typical dependent variable after averaging the values for each category. However, for some continuous and categorical predictors, we also made selections based on the principle of internal consistency between certain related variables, and we fixed a few categorical variables as identical across all three levels where doing so would simplify the modelling process (specification tables available: blue tit: https://osf.io/86akx; Eucalyptus: https://osf.io/jh7g5).

We used the 25th and 75th percentiles rather than minimum and maximum values to reduce the chance of occupying unrealistic parameter space. We planned to derive these predicted values from the model information provided by the individual analysts. All values (predictions) were first transformed to the original scale along with their standard errors (SE); we used the delta method (Ver Hoef 2012) for the transformation of SE. We used the square of the SE associated with predicted values as the sampling variance in the meta-analyses described below, and we planned to analyze these predicted values in exactly the same ways as we analyzed *Z_r* in the following analyses.

787

788 We plotted individual effect size estimates (Z_r) and predicted values of the dependent variable (y_i) 789 and their corresponding 95% confidence / credible intervals in forest plots to allow visualization of

Preregistration Deviation:

1. Standardizing blue tit out-of-sample predictions (y_i)

Because analysts of blue tit data chose different dependent variables on different scales, after transforming out-of-sample values to the original scales, we standardized all values as *z* scores ('standard scores') to put all dependent variables on the same scale and make them comparable. This involved taking each relevant value on the original scale (whether a predicted point estimate or a SE associated with that estimate) and subtracting the value in question from the mean value of that dependent variable derived from the full dataset and then dividing this difference by the standard deviation, SD, corresponding to the mean from the full dataset (<u>Supplementary</u> <u>Material B, Equation B.1</u>).

Note that we were unable to standardise some analyst-constructed variables, so these analyses were excluded from the final out-of-sample estimates meta-analysis, see <u>Supplementary Material</u> <u>B, section B.1.2.1</u> for details and explanation.

2. Log-transforming *Eucalyptus* out-of-sample predictions yi

All analyses of the *Eucalyptus* data chose dependent variables that were on the same scale, that is, *Eucalyptus* seedling counts. Although analysts may have used different size-classes of *Eucalyptus* seedlings for their dependent variable, we considered these choices to be akin to subsetting, rather than as different response variables, since changing the size-class of the dependent variable ultimately results in observations being omitted or included. Consequently, we did not standardise *Eucalyptus* out-of-sample predictions.

We were unable to fit quasi-Poisson or Poisson meta-regressions, as desired (O'Hara and Kotze 2010), because available meta-analysis packages (e.g. metafor:: and metainc::) do not provide implementation for outcomes as estimates-only, methods are only provided for outcomes as ratios or rate-differences between two groups. Consequently, we log-transformed the out-of-sample predictions for the *Eucalyptus* data and use the mean estimate for each prediction scenario as the dependent variable in our meta-analysis with the associated SE as the sampling variance in the meta-analysis (Nakagawa et al. 2023, Table 2).

the range and precision of effect size and predicted values. Further, we included these estimates in
random effects meta-analyses (Higgins et al. 2003; Borenstein et al. 2017) using the *metafor* package
in R (Viechtbauer 2010; R Core Team 2024):

- 793 $Z_r \sim 1 + (1|Effect ID)$
- 794 $y_i \sim 1 + (1|Effect ID)$

795 where y_i is the predicted value for the dependent variable at the 25th percentile, median, or 75th 796 percentile of the independent variables. The individual Z_r effect sizes were weighted with the inverse 797 of sampling variance for Z_r . The individual predicted values for dependent variable (y_i) were weighted 798 by the inverse of the associated SE² (original registration omitted "inverse of the" in error). These analyses provided an average Z_r score (\overline{Z}_r) or an average $y_i(\overline{y}_i)$ with corresponding 95% confidence 799 interval and allowed us to estimate two heterogeneity indices, τ^2 and I^2 . The former, τ^2 , is the 800 801 absolute measure of heterogeneity or the between-study variance (in our case, between-effect 802 variance) whereas I^2 is a relative measure of heterogeneity. We obtained the estimate of relative 803 heterogeneity (I^2) by dividing the between-effect variance by the sum of between-effect and withineffect variance (sampling error variance). l^2 is thus, in a standard meta-analysis, the proportion of variance that is due to heterogeneity as opposed to sampling error. When calculating l^2 , within-study variance is amalgamated across studies to create a "typical" within-study variance which serves as the sampling error variance (Higgins et al. 2003; Borenstein et al. 2017). Our goal here was to visualize and quantify the degree of variation among analyses in effect size estimates (Nakagawa and Cuthill 2007). We did not test for statistical significance.

810

Additional explanation:

Our use of I^2 to quantify heterogeneity violates an important assumption, but this violation does not invalidate our use of I² as a metric of how much heterogeneity can derive from analytical decisions. In standard meta-analysis, the statistic I² quantifies the proportion of variance that is greater than we would expect if differences among estimates were due to sampling error alone (Rosenberg 2013). However, it is clear that this interpretation does not apply to our value of l^2 because l^2 assumes that each estimate is based on an independent sample (although these analyses can account for non-independence via hierarchical modelling), whereas all our effects were derived from largely or entirely overlapping subsets of the same dataset. Despite this, we believe that I^2 remains a useful statistic for our purposes. This is because, in calculating I^2 , we are still setting a benchmark of expected variation due to sampling error based on the variance associated with each separate effect size estimate, and we are assessing how much (if at all) the variability among our effect sizes exceeds what would be expected had our effect sizes been based on independent data. In other words, our estimates can tell us how much proportional heterogeneity is possible from analytical decisions alone when sample sizes (and therefore metaanalytic within-estimate variance) are similar to the ones in our analyses. Among other implications, our violation of the independent sample assumption means that we (dramatically) over-estimate the variance expected due to sampling error, and because l^2 is a proportional estimate, we thus underestimate the actual proportion of variance due to differences among analyses other than sampling error. However, correcting this underestimation would create a trivial value since we designed the study so that much of the variance would derive from analytic decisions as opposed to differences in sampled data. Instead, retaining the l^2 value as typically calculated provides a useful comparison to l^2 values from typical meta-analyses.

Interpretation of τ^2 also differs somewhat from traditional meta-analysis, and we discuss this further in the Results.

811

812 Finally, we assessed the extent to which deviations from the meta-analytic mean by individual effect 813 sizes (Z_r) or the predicted values of the dependent variable (y_i) were explained by the peer rating of 814 each analysis team's method section, by a measurement of the distinctiveness of the set of predictor 815 variables included in each analysis, and by the choice of whether or not to include random effects in 816 the model. The deviation score, which served as the dependent variable in these analyses, is the 817 absolute value of the difference between the meta-analytic mean \overline{Z}_r (or \overline{y}_i) and the individual Z_r (or y_i) estimate for each analysis. We used the Box-Cox transformation on the absolute 818 819 values of deviation scores to achieve an approximately normal distribution (c.f. Fanelli and Ioannidis 820 2013; Fanelli, Costas, and Ioannidis 2017). We described variation in this dependent variable with

821 both a series of univariate analyses and a multivariate analysis. All these analyses were general linear

822 (mixed) models. These analyses were secondary to our estimation of variation in effect sizes

- 823 described above. We wished to quantify relationships among variables, but we had no a
- 824 priori expectation of effect size and made no dichotomous decisions about statistical significance.
- 825 When examining the extent to which reviewer ratings (on a scale from 0 to 100) explained deviation
- from the average effect (or predicted value), each analysis had been rated by multiple peer
- 827 reviewers, so for each reviewer score to be included, we include each deviation score in the analysis
- 828 multiple times. To account for the non-independence of multiple ratings of the same analysis, we
- planned to include analysis identity as a random effect in our general linear mixed model in
- the *lme4* package in R (Bates et al. 2015; R Core Team 2024). To account for potential differences
- among reviewers in their scoring of analyses, we also planned to include reviewer identity as a

Additional explanation:

In our meta-analyses based on Box-Cox transformed deviation scores, we leave these deviation scores unweighted. This is consistent with our registration, which did not mention weighting these scores. However, the fact that we did not mention weighting the scores was actually an error: we had intended to weight them, as is standard in meta-analysis, using the inverse variance of the Box-Cox transformed deviation scores Supplementary Material C, equation C.1. Unfortunately, when we did conduct the weighted analyses, they produced results in which some weighted estimates differed radically from the unweighted estimate because the weights were invalid. Such invalid weights can sometimes occur when the variance (upon which the weights depend) is partly a function of the effect size, as in our Box-Cox transformed deviation scores (Nakagawa et al. 2022). In the case of the Eucalyptus analyses, the most extreme outlier was weighted much more heavily (by close to two orders of magnitude) than any other effect sizes because the effect size was, itself, so high. Therefore, we made the decision to avoid weighting by inverse variance in all analyses of the Box-Cox transformed deviation scores. This was further justified because (a) most analyses have at least some moderately unreliable weights, and (b) the sample sizes were mostly very similar to each other across submitted analyses, and so meta-analytic weights are not particularly important here (Buck et al. 2022). We systematically investigated the impact of different weighting schemes and random effects on model convergence and results, see Supplementary Material C, section C.8 for more details.

832 random effect:

- B33 DeviationScore_i = BoxCox(DeviationFromMean_i)
- 834 DeviationScore_{*ii*} ~ Rating_{*ii*} + ReviewerID_{*i*} + $EffectID_i$
- 835 ReviewerID_i~ $N(0, \sigma_i^2)$
- 836 $EffectID \sim N(0, \sigma_i^2)$

837 Where DeviationFromMean_j is the deviation from the meta-analytic mean for the *j*th

analysis, Reviewer ID_i is the random intercept assigned to each *i* reviewer, and Effect ID_j is the

random intercept assigned to each *j* analysis, both of which are assumed to be normally distributed

840 with a mean of 0 and a variance of σ^2 . Absolute deviation scores were Box-Cox transformed using

the step_box_cox() function from the *timetk* package in R (Dancho and Vaughan 2023; R Core Team2024).

- 843 We conducted a similar analysis with the four categories of reviewer ratings ((1) deeply flawed and
- unpublishable, (2) publishable with major revision, (3) publishable with minor revision, (4)
- publishable as is) set as ordinal predictors numbered as shown here. As with the analyses above, we
- planned for these analyses to also include random effects of analysis identity and reviewer identity.
- 847 Both of these analyses (1: 1-100 ratings as the fixed effect, 2: categorical ratings as the fixed effects)
- 848 were planned to be conducted eight times for each dataset. Each of the four responses
- 849 $(Z_r, y_{25}, y_{50}, y_{75})$ were to be compared once to the initial ratings provided by the peer reviewers, and
- again based on the revised ratings provided by the peer reviewers.
- 851

Preregistration deviation:

- 1. We planned to include random effects of both analysis identity and reviewer identity in these models comparing reviewer ratings with deviation scores. However, after we received the analyses, we discovered that a subset of analyst teams had either conducted multiple analyses and/or identified multiple effects per analysis as answering the target question. We therefore faced an even more complex potential set of random effects. We decided that including Team ID and Effect ID along with Reviewer ID as random effects in the same model would almost certainly lead to model fit problems, and so we started with simpler models including just Effect ID and Reviewer ID. However, even with this simpler structure, our dataset was sparse, with reviewers rating a small number of analyses, resulting in models with singular fit (Supplementary Material C, section C.2). Removing one of the random effects was necessary for the models to converge. For both models of deviation from the meta-analytic mean explained by categorical or continuous reviewer ratings, we removed the random effect ID, leaving Reviewer ID as the only random effect.
- 2. We conducted analyses only with the final peer ratings after the opportunity for revision, not with the initial ratings. This was because when we recorded the final ratings, the initial ratings were over-written, therefore we did not have access to those initial values.

852 The next set of univariate analyses sought to explain deviations from the mean effects based on a 853 measure of the distinctiveness of the set of variables included in each analysis. As a 'distinctiveness' 854 score, we used Sorensen's Similarity Index (an index typically used to compare species composition 855 across sites), treating variables as species and individual analyses as sites. To generate an individual 856 Sorensen's value for each analysis required calculating the pairwise Sorensen's value for all pairs of 857 analyses (of the same dataset), and then taking the average across these Sorensen's values for each 858 analysis. We calculated the Sorensen's index values using the *betapart* package (Baselga et al. 859 2023) in R:

$$\beta Sorensen = \frac{b+c}{2a+b+c}$$

860

- 861 where a is the number of variables common to both analyses, b is the number of variables that occur
- in the first analysis but not in the second and c is the number of variables that occur in the second
- analysis. We then used the per-model average Sorensen's index value as an independent variable to
- 864 predict the deviation score in a general linear model, and included no random effect since each
- analysis is included only once, in R (R Core Team 2024):

866 $DeviationScore_j \sim \beta Sorensen_j$

Additional explanation:

When we planned this analysis, we anticipated that analysts would identify a single primary effect from each model, so that each model would appear in the analysis only once. Our expectation was incorrect because some analysts identified >1 effect per analysis, but we still chose to specify our model as registered and not use a random effect. This is because most models produced only one effect and so we expected that specifying a random effect to account for the few cases where >1 effect was included for a given model would prevent model convergence.

Note that this analysis contrasts with the analyses in which we used reviewer ratings as predictors because in the analyses with reviewer ratings, each effect appeared in the analysis approximately four times due to multiple reviews of each analysis, and so it was much more important to account for that variance through a random effect.

867 Next, we assessed the relationship between the inclusion of random effects in the analysis and the

- 868 deviation from the mean effect size. We anticipated that most analysts would use random effects in a
- 869 mixed model framework, but if we were wrong, we wanted to evaluate the differences in outcomes
- 870 when using random effects versus not using random effects. Thus, if there were at least 5 analyses
- that did and 5 analyses that did not include random effects, we would add a binary predictor variable
- 872 "random effects included (yes/no)" to our set of univariate analyses and would add this predictor
- 873 variable to our multivariate model described below. This standard was only met for

the *Eucalyptus* analyses, and so we only examined inclusion of random effects as a predictor variable
 in meta-analysis of this set to analyses.

- 876 Finally, we conducted a multivariate analysis with the five predictors described above (peer ratings 0-
- 877 100 and peer ratings of publishability 1-4; both original and revised and Sorensen's index, plus a sixth
- for *Eucalyptus*, presence / absence of random effects) with random effects of analysis identity and

879 reviewer identity in the *lme4* package in R (Bates et al. 2015; R Core Team 2024). We had stated here

- in the text that we would use only the revised (final) peer ratings in this analysis, so the absence of
- the initial ratings is not a deviation from our plan:
- 882 DeviationScore_{*j*} = BoxCox(DeviationFromMean_{*j*})
- 883 DeviationScore_{*ij*}~RatingContinuous_{*ij*} + RatingCategorical_{*ij*} + β Sorensen_{*j*} + ReviewerID_{*i*} 884 + Effect ID_{*j*}
- 885 ReviewerID_i~ $N(0, \sigma_i^2)$
- 886 EffectID_{*i*}~ $N(0, \sigma_i^2)$
- 887

- 888 We conducted all the analyses described above eight times; for each of the four responses
- 889 $(Z_r, y_{25}, y_{50}, y_{75})$ one time for each of the two datasets.
- 890 We have publicly archived all relevant data, code, and materials on the Open Science Framework 891 (https://osf.io/mn5ai/). Archived data includes the original datasets distributed to all analysts, any
- (<u>https://osf.io/mn5aj/</u>). Archived data includes the original datasets distributed to all analysts, any
 edited versions of the data analyzed by individual groups, and the data we analyzed with our meta-
- analyses, which include the effect sizes derived from separate analyses, the statistics describing
- variation in model structure among analyst groups, and the anonymized answers to our surveys of
- analysts and peer reviewers. Similarly, we have archived both the analysis code used for each
- individual analysis (where available) and the code from our meta-analyses. We have also archived
- 897 copies of our survey instruments from analysts and peer reviewers.
- 898Our rules for excluding data from our study were as follows. We excluded from our synthesis any899individual analysis submitted after we had completed peer review or those unaccompanied by
- analysis files that allow us to understand what the analysts did. We also excluded any individual
- analysis that did not produce an outcome that could be interpreted as an answer to our primary
- 902 question (as posed above) for the respective dataset. For instance, this means that in the case of the
- data on blue tit chick growth, we excluded any analysis that did not include something that can be
- 904 interpreted as growth or size as a dependent (response) variable, and in the case of
- 905 the *Eucalyptus* establishment data, we excluded any analysis that did not include a measure of grass
- 906 cover among the independent (predictor) variables. Also, as described above, any analysis that could
- not produce an effect that could be converted to a signed Z_r was excluded from analyses of Z_r .

Preregistration Deviation:

Some analysts had difficulty implementing our instructions to derive the out-of-sample predictions, and in some cases (especially for the *Eucalyptus* data), they submitted predictions with implausibly extreme values. We believed these values were incorrect and thus made the conservative decision to exclude out-of-sample predictions where the estimates were > 3 standard deviations from the mean value from the full dataset provided to teams for analysis.

Additional explanation: unregistered analyses

1. Evaluating model fit.

We evaluated all fitted models using the <u>performance::performance()</u> function from the *performance* package (Lüdecke, Ben-Shachar, et al. 2021) and the glance() function from the *broom.mixed* package (Bolker et al. 2024). For all models, we calculated the square root of the residual variance (Sigma) and the root mean squared error (RMSE). For

GLMMs <u>performance::performance()</u> calculates the marginal and conditional R² values as well as the contribution of random effects (ICC), based on Nakagawa et al. (2017). The

conditional R^2 accounts for both the fixed and random effects, while the marginal R^2 considers only the variance of the fixed effects. The contribution of random effects is obtained by subtracting the marginal R^2 from the conditional R^2 .

2. Exploring outliers and analysis quality.

After seeing the forest plots of *Zr* values and noticing the existence of a small number of extreme outliers, especially from the *Eucalyptus* analyses, we wanted to understand the degree to which our heterogeneity estimates were influenced by these outliers. To explore this question, we removed the highest two and lowest two values of *Zr* in each dataset and re-calculated our heterogeneity estimates.

To help understand the possible role of the quality of analyses in driving the heterogeneity we observed among estimates of Zr, we created forest plots and recalculated our heterogeneity estimates after removing all effects from analysis teams that had received at least one rating of "deeply flawed and unpublishable" and then again after removing all effects from analysis teams with at least one rating of either "deeply flawed and unpublishable" or "publishable with major revisions". We also used self-identified levels of statistical expertise to examine heterogeneity when we retained analyses only from analysis teams that contained at least one member who rated themselves as "highly proficient" or "expert" (rather than "novice" or "moderately proficient") in conducting statistical analyses in their research area in our intake survey. Additionally, to assess potential impacts of highly collinear predictor variables on estimates of Zr in blue tit analyses, we created forest plots (Supplementary Material B, Figure B.5) and recalculated our heterogeneity estimates after we removed analyses that contained the brood count after manipulation and the highly correlated (correlation of 0.89, Supplementary Material D, Figure D.2) brood count at day 14. This removal included the one effect based on a model that contained both these variables and a third highly correlated variable, the estimate of number of chicks fledged (the only model that included the estimate of number of chicks fledged). We did not conduct a similar analysis for the *Eucalyptus* dataset because there were no variables highly collinear with the primary predictors (grass cover variables) in that dataset (Supplementary Material D, Figure D.1).

3. Exploring possible impacts of lower quality estimates of degrees of freedom.

Our meta-analyses of variation in Z_r required variance estimates derived from estimates of the degrees of freedom in original analyses from which Z_r estimates were derived. While processing the estimates of degrees of freedom submitted by analysts, we identified a subset of these estimates in which we had lower confidence because two or more effects from the same analysis were submitted with identical degrees of freedom. We therefore conducted a second set of (more conservative) meta-analyses that excluded these Z_r estimates with identical estimates of degrees of freedom and we present these analyses in the supplement.

Additional explanation: Best practices in many-analysts research

After we initiated our project, a paper was published outlining best practices in many-analysts studies (Aczel et al. 2021). Although we did not have access to this document when we implemented our project, our study complies with these practices nearly completely. The one exception is that although we requested analysis code from analysts, we did not require submission of code.

910

911 Step 6: Facilitated Discussion and Collaborative Write-Up of

912 Manuscript

- 913 We planned for analysts and initiating authors to discuss the limitations, results, and implications of
- 914 the study and collaborate on writing the final manuscript for review as a stage-2 Registered Report.

Preregistration deviation:

As described above, due to the large number of recruited analysts and reviewers and the anticipated challenges of receiving and integrating feedback from so many authors, we limited analyst and reviewer participation in the production of the final manuscript to an invitation to call attention to serious problems with the manuscript draft.

- 915 We built an R package, ManyEcoEvo:: to conduct the analyses described in this study (Gould et al.
- 916 2023), which can be downloaded from https://github.com/egouldo/ManyEcoEvo/ to reproduce our
- 917 analyses or replicate the analyses described here using alternate datasets. Data cleaning and
- 918 preparation of analysis-data, as well as the analysis, is conducted in R (R Core Team
- 2024) reproducibly using the targets package (Landau 2021). This data and analysis pipeline is stored
- 920 in the ManyEcoEvo:: package repository and its outputs are made available to users of the package
- 921 when the library is loaded.
- 922 The full manuscript, including further analysis and presentation of results is written in Quarto (Allaire
- 923 et al. 2024). The source code to reproduce the manuscript is hosted at
- 924 <u>https://github.com/egouldo/ManyAnalysts/ (Gould et al. 2024)</u>, and the rendered version of the
- 925 source code may be viewed at https://egouldo.github.io/ManyAnalysts/. All R packages and their
- 926 versions used in the production of the manuscript are listed in Table 7 at the end of this paper.

927 Results

928 Summary Statistics

929 In total, 173 analyst teams, comprising 246 analysts, contributed 182 usable analyses (compatible

- 930 with our meta-analyses and provided with all information needed for inclusion) of the two datasets
- examined in this study which yielded 215 effects. Analysts produced 134 distinct effects that met our
- 932 criteria for inclusion in at least one of our meta-analyses for the blue tit dataset. Analysts produced
- 933 81 distinct effects meeting our criteria for inclusion for the *Eucalyptus* dataset. Excluded analyses and
- 934 effects either did not answer our specified biological questions, were submitted with insufficient
- information for inclusion in our meta-analyses, or were incompatible with production of our effect
- size(s). We expected cases of this final scenario (incompatible analyses), for instance we cannot
- 937 extract a Z_r from random forest models, which is why we analyzed two distinct types of effects, Z_r and

938 out-of-sample predictions. Some effects only provided sufficient information for a subset of analyses
 939 and were only included in that subset. For both datasets, most submitted analyses incorporated
 940 mixed effects. Submitted analyses of the blue tit dataset typically specified normal error and analyses
 941 of the *Eucalyptus* dataset typically specified a non-normal error distribution (Supplementary Material
 942 A, Table A.1).

943 For both datasets, the composition of models varied substantially in regards to the number of fixed 944 and random effects, interaction terms, and the number of data points used, and these patterns 945 differed somewhat between the blue tit and Eucalyptus analyses (See Supplementary Material A, 946 <u>Table A.2</u>). Focusing on the models included in the Z_r analyses (because this is the larger sample), 947 blue tit models included a similar number of fixed effects on average (mean 5.2 \pm 2.92 SD, range: 1 to 948 19) as Eucalyptus models (mean 5.01 ± 3.83 SD, range: 1 to 13), but the standard deviation in 949 number of fixed effects was somewhat larger in the Eucalyptus models. The average number of 950 interaction terms was much larger for the blue tit models (mean 0.44 ± 1.11 SD, range: 0 to 10) than 951 for the Eucalyptus models (mean 0.16 ± 0.65 SD, range: 0 to 5), but still under 0.5 for both, indicating 952 that most models did not contain interaction terms. Blue tit models also contained more random 953 effects (mean 3.53 ± 2.08 SD, range: 0 to 10) than Eucalyptus models (mean 1.41 ± 1.09 SD, range: 0 954 to 4). The maximum possible sample size in the blue tit dataset (3720 nestlings) was an order of 955 magnitude larger than the maximum possible in the *Eucalyptus* dataset (351 plots), and the means 956 and standard deviations of the sample size used to derive the effects eligible for our study were also 957 an order of magnitude greater for the blue tit dataset (mean 2611.09 ± 937.48 SD, range: 76 to 76) 958 relative to the Eucalyptus models (mean 298.43 ± 106.25 SD, range: 18 to 351). However, the 959 standard deviation in sample size from the Eucalyptus models was heavily influenced by a few cases 960 of dramatic sub-setting (described below). Approximately three quarters of Eucalyptus models used 961 sample sizes within 3% of the maximum. In contrast, fewer than 20% of blue tit models relied on 962 sample sizes within 3% of the maximum, and approximately 50% of blue tit models relied on sample 963 sizes 29% or more below the maximum.

Analysts provided qualitative descriptions of the conclusions of their analyses. Each analysis team
 provided one conclusion per dataset. These conclusions could take into account the results of any
 formal analyses completed by the team as well as exploratory and visual analyses of the data. Here
 we summarize all qualitative responses, regardless of whether we had sufficient information to use
 the corresponding model results in our quantitative analyses below. We classified these conclusions
 into the categories summarized below (Table 1):

- *Mixed*: some evidence supporting a positive effect, some evidence supporting a negative effect
- 971 *Conclusive negative*: negative relationship described without caveat
- *Qualified negative*: negative relationship but only in certain circumstances or where analysts
 express uncertainty in their result
- Conclusive none: analysts interpret the results as conclusive of no effect
- *Qualified none*: analysts describe finding no evidence of a relationship but they describe the
 potential for an undetected effect
- *Qualified positive*: positive relationship described but only in certain circumstances or where
 analysts express uncertainty in their result
- 979 *Conclusive positive*: positive relationship described without caveat

For the blue tit dataset, most analysts concluded that there was negative relationship between
 measures of sibling competition and nestling growth, though half the teams expressed qualifications
 or described effects as mixed or absent. No analysts concluded that there was a positive relationship

- 983 even though some individual effect sizes were positive, apparently because all analysts who
- 984 produced effects indicating positive relationships also produced effects indicating negative
- 985 relationships and therefore described their results as qualified, mixed, or absent. For
- 986 the *Eucalyptus* dataset, there was a broader spread of conclusions with at least one analyst team
- 987 providing conclusions consistent with each conclusion category. The most common conclusion for
- 988 the Eucalyptus dataset was that there was no relationship between grass cover
- 989 and Eucalyptus recruitment (either conclusive or qualified description of no relationship), but more
- 990 than half the teams concluded that there were effects; negative, positive, or mixed.
- 991 Table 1: Tallies of analysts' qualitative answers to the research questions addressed by their analyses.

Dataset	Mixed	Negative	Negative	None	None	Positive	Positive
		Conclusive	Qualified	Conclusive	Qualified	Qualified	Conclusive
blue tit	5	37	27	4	1	0	0
Eucalyptus	8	6	12	19	12	4	2

992

993 Distribution of effects

994 Effect sizes (Z_r)

995 Although the majority (118 of 131) of the usable Zr effects from the blue tit dataset found nestling 996 growth decreased with sibling competition, and the meta-analytic mean \bar{Z}_r (Fisher's transformation of the correlation coefficient) was convincingly negative (-0.35 ± 0.06 95%CI), there was substantial 997 998 variability in the strength and the direction of this effect. Z_r ranged from -1.55 to 0.38, and 999 approximately continuously from -0.93 to 0.19 (Figure 2a and Table 4), and of the 118 effects with 1000 negative slopes, 93 had confidence intervals excluding 0. Of the 13 with positive slopes indicating 1001 increased nestling growth in the presence of more siblings, 2 had confidence intervals excluding zero 1002 (Figure 2a).

1003 Meta-analysis of the Eucalyptus dataset also showed substantial variability in the strength of effects 1004 as measured by Z_{r_i} and unlike with the blue tits, a notable lack of consistency in the direction of 1005 effects (Figure 2b, Table 4). Zr ranged from -4.47 (Supplementary Material A, Figure A.2), indicating a 1006 strong tendency for reduced Eucalyptus seedling success as grass cover increased, to 0.39, indicating 1007 the opposite. Although the range of reported effects skewed strongly negative, this was due to a 1008 small number of substantial outliers. Most values of Z_r were relatively small with values < |0.2| and 1009 the meta-analytic mean effect size was close to zero (-0.09 ± 0.12 95%Cl). Of the 79 effects, fifty-1010 three had confidence intervals overlapping zero, approximately a quarter (fifteen) crossed the 1011 traditional threshold of statistical significance indicating a negative relationship between grass cover 1012 and seedling success, and eleven crossed the significance threshold indicating a positive relationship 1013 between grass cover and seedling success (Figure 2b).

1014



1015

1016 Figure 2: Forest plots of meta-analytic estimated standardized effect sizes (Z_r , blue triangles) and 1017 their 95% confidence intervals for each effect size included in the meta-analysis model. (A) Blue tit 1018 analyses: Points where Z_r are less than 0 indicate analyses that found a negative relationship 1019 between sibling number and nestling growth. (B) Eucalyptus analyses: Points where Z_r are less than 0 1020 indicate a negative relationship between grass cover and Eucalyptus seedling success. The meta-1021 analytic mean effect size is denoted by a black circle and a dashed vertical line, with error bars also 1022 representing the 95% confidence interval. The solid black vertical line demarcates effect size of 0, 1023 indicating no relationship between the test variable and the response variable. Note that 1024 the Eucalyptus plot omits one extreme outlier with the value of -4.47 (Supplementary Material A, 1025 Figure A.2) in order to standardize the x-axes on these two panels.

1026 Out-of-sample predictions (y_i)

1027 As with the effect size Z_{r_i} we observed substantial variability in the size of out-of-sample predictions 1028 derived from the analysts' models. Blue tit predictions (Figure 3a), which were z-score-standardised 1029 to accommodate the use of different response variables, always ranged far in excess of one standard 1030 deviation. In the y_{25} scenario, model predictions ranged from -1.84 to 0.42 (a range of 2.68 standard 1031 deviations), in the y₅₀ they ranged from -0.52 to 1.08 (a range of 1.63 standard deviations), and in 1032 the y₇₅ scenario they ranged from -0.03 to 1.59 (a range of 1.9 standard deviations). As should be 1033 expected given the existence of both negative and positive Z, values, all three out-of-sample 1034 scenarios produced both negative and positive predictions, although as with the Z_r values, there is a 1035 clear trend for scenarios with more siblings to be associated with smaller nestlings. This is supported 1036 by the meta-analytic means of these three sets of predictions which were -0.66 (95%CI -0.82–0.5) for 1037 the y₂₅, 0.34 (95%CI 0.2-0.48) for the y₅₀, and 0.67 (95%CI 0.57-0.77) for the y₇₅.

Eucalyptus out-of-sample predictions also varied substantially (Figure 3b), but because they were not
 z-score-standardised and are instead on the original count scale, the types of interpretations we can

1040 make differ. The predicted *Eucalyptus* seedling counts per 15 x 15 m plot for the y₂₅ scenario ranged

- 1041 from 0.04 to 26.99, for the y_{50} scenario ranged from 0.04 to 44.34, and for the y_{75} scenario they
- 1042 ranged from 0.03 to 61.34. The meta-analytic mean predictions for these three scenarios were
- 1043 similar; 1.27 (95%Cl 0.59-2.3) for the y_{25} , 2.92 (95%Cl 0.98-3.89) for the y_{50} , and 2.92 (95%Cl 1.59-
- 1044 4.9) for the y₇₅ scenarios respectively.



1045

1046

1047 Figure 3: Forest plot of meta-analytic estimated out-of-sample predictions. A) Standardized (z-score) 1048 blue tit out-of-sample predictions, y_i. B) response-scale (stem counts) Eucalyptus out-of-sample 1049 predictions. Triangles represent individual estimates. Circles represent the meta-analytic mean for 1050 each prediction scenario. Dark-blue points correspond to y25 scenario, medium-blue points 1051 correspond to the y_{50} scenario, while light blue points correspond to the y_{75} scenario. Error bars are 1052 95% confidence intervals. Note that, for the Eucalyptus analysis, outliers (observations more than 3 1053 SD above the mean) have been removed prior to model fitting and do not appear on this figure. The 1054 x-axis is truncated to approximately 140, and thus some error bars are incomplete. 1055 See Supplementary Material B, Figure B.6 for full figure.

1056

1057 Quantifying heterogeneity

1058 Effect sizes (Z_r)

1059 We quantified both absolute (τ^2) and relative (l^2) heterogeneity resulting from analytical variation.

- 1060 Both measures suggest that substantial variability among effect sizes was attributable to the
- 1061 analytical decisions of analysts.
- 1062 The total absolute level of variance beyond what would typically be expected due to sampling 1063 error, τ^2 (Table 2), among all usable blue tit effects was 0.08 and for *Eucalyptus* effects was 0.27. This
- 1064 is similar to or exceeding the median value (0.105) of τ^2 found across 31 recent meta-
- analyses (calculated from the data in Yang et al. 2023). The similarity of our observed values to
- 1066 values from meta-analyses of different studies based on different data suggest the potential for a
- 1067 large portion of heterogeneity to arise from analytical decisions. For further discussion of
- 1068 interpretation of τ^2 in our study, please consult discussion of post hoc analyses below.
- 1069Table 2: Heterogeneity in the estimated effects Z_r for meta-analyses of: the full dataset, as well as1070from post hoc analyses wherein analyses with outliers are removed, analyses with effects from1071analysis teams with at least one "unpublishable" rating are excluded, analyses receiving at least one1072"major revisions" rating or worse excluded, analyses from teams with at least one analyst self-rated1073as "highly proficient" or "expert" in statistical analysis are included, and (blue tit only) analyses that1074did not included the pair of highly collinear predictors together. τ^2_{Team} is the absolute heterogeneity1075for the random effect Team. $\tau^2_{Effect ID}$ is the absolute heterogeneity for the random effect Effect
- 1076 ID nested under Team. Effect ID is the unique identifier assigned to each individual statistical effect
- 1077 submitted by an analysis team. We nested Effect ID within analysis team identity (Team) because
- 1078 analysis teams often submitted >1 statistical effect, either because they considered >1 model or
- 1079 because they derived >1 effect per model, especially when a model contained a factor with multiple
- 1080 levels that produced >1 contrast. τ^2_{Total} is the total absolute heterogeneity. I^2_{Total} is the proportional
- 1081 heterogeneity; the proportion of the variance among effects not attributable to sampling
- 1082 error, l^2_{Team} is the subset of the proportional heterogeneity due to differences
- among Teams and $l^2_{\text{Team, Effect ID}}$ is subset of the proportional heterogeneity attributable to among-

1084	Effect ID differences.
------	------------------------

Dataset	N _{Obs}	τ^2_{Total}	τ^2_{Team}	$\tau^2_{\text{Effect ID}}$	I ² _{Total}	I ² _{Team}	I ² _{Team, Effect ID}		
All Analyses									
Eucalyptus	79	0.27	0.02	0.25	98.59%	6.89%	91.70%		
blue tit	131	0.08	0.03	0.05	97.61%	36.71%	60.90%		
Blue tit analy	/ses contain	ing highly co	ollinear pred	ictors remov	red				
blue tit	117	0.07	0.04	0.03	96.92%	58.18%	38.75%		
All analyses,	outliers rem	noved							
Eucalyptus	75	0.01	0.00	0.01	66.19%	19.25%	46.94%		
blue tit	127	0.07	0.04	0.02	96.84%	64.63%	32.21%		
Analyses rec	eiving at lea	st one 'Unp	ublishable' r	ating remove	ed				
Eucalyptus	55	0.01	0.01	0.01	79.74%	28.31%	51.43%		
blue tit	109	0.08	0.03	0.05	97.52%	35.68%	61.84%		
Analyses rec	eiving at lea	st one 'Unp	ublishable' a	nd or 'Majoı	r Revisions' r	ating remov	ed		
Eucalyptus	13	0.03	0.03	0.00	88.91%	88.91%	0.00%		
blue tit	32	0.14	0.01	0.13	98.72%	5.17%	93.55%		
Analyses from	m teams wit	h highly pro:	ficient or ex	pert data an	alysts				
Eucalyptus	34	0.58	0.02	0.56	99.41%	3.47%	95.94%		
blue tit	89	0.09	0.03	0.06	97.91%	31.43%	66.49%		

1085

1086 In our analyses, *l*² is a plausible index of how much more variability among effect sizes we have 1087 observed, as a proportion, than we would have observed if sampling error were driving variability. 1088 We discuss our interpretation of *l*² further in the methods, but in short, it is a useful metric for 1089 comparison to values from published meta-analyses and provides a plausible value for how much 1090 heterogeneity could arise in a normal meta-analysis with similar sample sizes due to analytical 1091 variability alone. In our study, total l^2 for the blue tit Z_r estimates was extremely large, at 97.61%, as 1092 was the *Eucalyptus* estimate (98.59% Table 2).

- 1093 Although the overall *l*² values were similar for both *Eucalyptus* and blue tit analyses, the relative 1094 composition of that heterogeneity differed. For both datasets, the majority of heterogeneity
- in Z_r was driven by differences among effects as opposed to differences among teams, though this
- 1096 was more prominent for the *Eucalyptus* dataset, where nearly all of the total heterogeneity was
- driven by differences among effects (91.7%) as opposed to differences among teams (6.89%) (Table 2).

1099 Out-of-sample predictions (y_i)

1100 We observed substantial heterogeneity among out-of-sample estimates, but the pattern differed 1101 somewhat from the Z_r values (Table 3). Among the blue tit predictions, l^2 ranged from medium-high 1102 for the y_{25} scenario (68.54) to low (27.9) for the y_{75} scenario. Among

1103 the *Eucalyptus* predictions, l^2 values were uniformly high (>82%). For both datasets, most of the

1104 existing heterogeneity among predicted values was attributable to among-team differences, with the

exception of the y_{50} analysis of the *Eucalyptus* dataset. We are limited in our interpretation of τ^2 for

1106 these estimates because, unlike for the Z_r estimates, we have no benchmark for comparison with 1107 other meta-analyses.

1108 Table 3: Heterogeneity among the out-of-sample predictions *y_i* for both blue tit

- and *Eucalyptus* datasets. τ^{2}_{Team} is the absolute heterogeneity for the random effect Team. $T^{2}_{Effect ID}$ is
- 1110 the absolute heterogeneity for the random effect Effect ID nested under Team. Effect ID is the unique
- 1111 identifier assigned to each individual statistical effect submitted by an analysis team. We
- 1112 nested Effect ID within analysis team identity (Team) because analysis teams often submitted >1
- 1113 statistical effect, either because they considered >1 model or because they derived >1 effect per
- 1114 model, especially when a model contained a factor with multiple levels that produced >1
- 1115 contrast. τ^2_{Total} is the total absolute heterogeneity. I^2_{Total} is the proportional heterogeneity; the
- 1116 proportion of the variance among effects not attributable to sampling error, l^2_{Team} is the subset of the
- 1117 proportional heterogeneity due to differences among Teams and $I^2_{\text{Team,Effect ID}}$ is subset of the
- 1118 proportional heterogeneity attributable to among-Effect ID differences.

Prediction	N _{Obs}	T _{Total}	T ² _{Team}	T ² Effect ID	I ² _{Total}	I ² _{Team}	I ² _{Team, Effect ID}
Scenario							
blue tit							
y25	63	0.23	0.11	0.03	68.54%	53.43%	15.11%
y50	60	0.23	0.06	0.00	50%	46.29%	3.71%
y75	63	0.23	0.02	0.00	27.9%	27.89%	0.01%
Eucalyptus							
y25	38	5.75	1.48	0.68	86.93%	59.54%	27.39%
y50	38	5.75	1.32	0.83	89.63%	55%	34.64%
y75	38	5.75	1.03	0.41	80.19%	57.41%	22.78%

1119

Post-hoc analysis: Exploring outlier characteristics and the effect ofoutlier removal on heterogeneity

1122 Effect sizes (Z_r)

1123 The outlier *Eucalyptus Zr* values were striking and merited special examination. The three negative 1124 outliers had very low sample sizes that were based on either small subsets of the dataset or, in one 1125 case, extreme aggregation of data. The outliers associated with small subsets had sample sizes 1126 (n= 117, 90, 18) that were less than half of the total possible sample size of 351. The case of extreme 1127 aggregation involved averaging all values within each of the 351 sites in the dataset.

1128 Surprisingly, both the largest and smallest effect sizes in the blue tit analyses (Figure 2a) come from 1129 the same analyst (anonymous ID: 'Adelong'), with identical models in terms of the explanatory 1130 variable structure, but with different response variables. However, the radical change in effect was 1131 primarily due to collinearity with covariates. The primary predictor variable (brood count after 1132 manipulation) was accompanied by several collinear variables, including the highly collinear 1133 (correlation of 0.89 Supplementary Material D, Figure D.2) covariate (brood count at day 14) in both 1134 analyses. In the analysis of nestling weight, brood count after manipulation showed a strong positive 1135 partial correlation with weight after controlling for brood count at day 14 and treatment category 1136 (increased, decreased, unmanipulated). In that same analysis, the most collinear covariate (the day 1137 14 count) had a negative partial correlation with weight. In the analysis with tarsus length as the 1138 response variable, these partial correlations were almost identical in absolute magnitude, but 1139 reversed in sign and so brood count after manipulation was now the collinear predictor with the 1140 negative relationship. The two models were therefore very similar, but the two collinear predictors 1141 simply switched roles, presumably because a subtle difference in the distribution of weight and 1142 tarsus length data.

When we dropped the *Eucalyptus* outliers, l^2 decreased from high (98.59 %), using Higgins' (Higgins 1143 1144 et al. 2003) suggested benchmark, to between moderate and high (66.19 %, Table 2). However, more 1145 notably, τ^2 dropped from 0.27 to 0.01, indicating that, once outliers were excluded, the observed 1146 variation in effects was similar to what we would expect if sampling error were driving the differences among effects (since τ^2 is the variance beyond that driven by sampling error). The 1147 interpretation of this value of τ^2 in the context of our many-analyst study is somewhat different than 1148 1149 a typical meta-analysis, however, since in our study (especially for *Eucalyptus*, where most analyses 1150 used almost exactly the same data points), there is almost no role for sampling error in driving the 1151 observed differences among the estimates. Thus, rather than concluding that the variability we 1152 observed among estimates (after removing outliers) was due only to sampling error (because τ^2 became small: 10% of the median from Yang et al. 2023), we instead conclude that 1153 1154 the observed variability, which must be due to the divergent choices of analysts rather than sampling 1155 error, is approximately of the same magnitude as what we would have expected if, instead, sampling 1156 error, and not analytical heterogeneity, were at work. Conversely, dropping outliers from the set of 1157 blue tit effects did not meaningfully reduce l^2 , and only modestly reduced τ^2 (Table 2). Thus, effects

at the extremes of the distribution were much stronger contributors to total heterogeneity for effectsfrom analyses of the *Eucalyptus* than for the blue tit dataset.

1160Table 4: Estimated mean value of the standardised correlation coefficient, \bar{Z}_r , along with its standard1161error and 95% confidence intervals. We re-computed the meta-analysis for different post hoc subsets1162of the data: All eligible effects, removal of effects from blue tit analyses that contained a pair of1163highly collinear predictor variables, removal of effects from analysis teams that received at least one

1164 peer rating of "deeply flawed and unpublishable", removal of any effects from analysis teams that

1165 received at least one peer rating of either "deeply flawed and unpublishable" or "publishable with

1166 major revisions",, inclusion of only effects from analysis teams that included at least one member

1167 who rated themselves as "highly proficient" or "expert" at conducting statistical analyses in their

1168 research area.

Dataset	û	$SE[\widehat{\mu}]$	95% CI	statistic	р				
All analyses									
Eucalyptus	-0.09	0.06	[-0.22,0.03]	-1.47	0.14				
blue tit	-0.35	0.03	[-0.41,-0.29]	-11.02	<0.001				
Blue tit analyses	s containing high	ly collinear predic	ctors removed						
blue tit	-0.36	0.03	[-0.42,-0.29]	-10.97	<0.001				
All analyses, ou	tliers removed								
Eucalyptus	-0.03	0.01	[-0.06,0.00]	-2.23	0.026				
blue tit	-0.36	0.03	[-0.42,-0.30]	-11.48	<0.001				
Analyses receivi	ing at least one 'l	Jnpublishable' ra	ting removed						
Eucalyptus	-0.02	0.02	[-0.07,0.02]	-1.15	0.3				
blue tit	-0.36	0.03	[-0.43,-0.30]	-10.82	<0.001				
Analyses receivi	ing at least one 'l	Jnpublishable' an	d or 'Major Revis	sions' rating remo	oved				
Eucalyptus	-0.04	0.05	[-0.15,0.07]	-0.77	0.4				
blue tit	-0.37	0.07	[-0.51,-0.23]	-5.34	<0.001				
Analyses from t	eams with highly	proficient or exp	ert data analysts						
Eucalyptus	-0.17	0.13	[-0.43,0.10]	-1.24	0.2				
blue tit	-0.36	0.04	[-0.44,-0.28]	-8.93	<0.001				

1169

1170 Out-of-sample predictions (y_i)

1171 We did not conduct these post hoc analyses on the out-of-sample predictions as the number of 1172 eligible effects was smaller and the pattern of outliers differed.

¹¹⁷³ Post hoc analysis: Exploring the effect of removing analyses with poor

1174 peer ratings on heterogeneity

1175 Effect sizes (Z_r)

1176 Removing poorly rated analyses had limited impact on the meta-analytic means (Supplementary

1177 Material B, Figure B.3). For the *Eucalyptus* dataset, the meta-analytic mean shifted from -0.09 to -

1178 0.02 when effects from analyses rated as unpublishable were removed, and to -0.04 when effects

1179 from analyses rated, at least once, as unpublishable or requiring major revisions were removed.

1180 Further, the confidence intervals for all of these means overlapped each of the other means

1181 (Table 4). We saw similar patterns for the blue tit dataset, with only small shifts in the meta-analytic

- mean, and confidence intervals of all three means overlapping each other mean (Table 4). Refitting
- 1183 the meta-analysis with a fixed effect for categorical ratings also showed no indication of differences
- 1184 in group meta-analytic means due to peer ratings (<u>Supplementary Material B, Figure B.1</u>).
- 1185 For the blue tit dataset, removing poorly-rated analyses led to only negligible changes in l^2_{Total} and
- 1186 relatively minor impacts on τ^2 . However, for the *Eucalyptus* dataset, removing poorly-rated analyses
- 1187 led to notable reductions in l^2_{Total} and substantial reductions in τ^2 . When including all analyses,
- 1188 the *Eucalyptus I*²_{Total} was 98.59% and τ^2 was 0.27, but eliminating analyses with ratings of
- 1189 "unpublishable" reduced I_{Total}^2 to 79.74% and τ^2 to 0.01, and removing also those analyses "needing

- 1190 major revisions" left l_{Total}^2 at 88.91% and τ^2 at 0.03 (Table 2). Additionally, the allocations of l^2 to the
- 1191 team versus individual effect were altered for both blue tit and *Eucalyptus* meta-analyses by
- 1192 removing poorly-rated analyses, but in different ways. For blue tit meta-analysis, between a third and
- 1193 two-thirds of the total l^2 was attributable to among-team variance in most analyses until both
- analyses rated "unpublishable" and analyses rated in need of "major revision" were eliminated, in
- 1195 which case almost all remaining heterogeneity was attributable to among-effect differences. In
- 1196 contrast, for *Eucalyptus* meta-analysis, the among-team component of l^2 was less than third until
- 1197 both analyses rated "unpublishable" and analyses rated in need of "major revision" were eliminated,
- in which case almost 90% of heterogeneity was attributable to differences among teams.

1199 Out-of-sample predictions (y_i)

- We did not conduct these post hoc analyses on the out-of-sample predictions as the number of
 eligible effects was smaller and our ability to interpret heterogeneity values for these analyses was
 limited
- 1203 Post hoc analysis: Exploring the effect of including only analyses
- 1204 conducted by analysis teams with at least one member self-rated as
- ¹²⁰⁵ "highly proficient" or "expert" in conducting statistical analyses in
- 1206 their research area

1207 Effect sizes (Z_r)

1208 Including only analyses conducted by teams that contained at least one member who rated
1209 themselves as "highly proficient" or "expert" in conducting the relevant statistical methods had

- 1210 negligible impacts on the meta-analytic means (Table 4), the distribution of Z_r effects (Supplementary 1211 Natorial D. Sigure D. 4), or beta approximates (Table 2), which remained automatic high
- 1211 <u>Material B, Figure B.4</u>), or heterogeneity estimates (Table 2), which remained extremely high.

1212 Out-of-sample predictions (y_i)

We did not conduct these post hoc analyses on the out-of-sample predictions as the number ofeligible effects was smaller.

Post hoc analysis: Exploring the effect of excluding estimates of Z_r in

1216 which we had reduced confidence

1217 As described in our addendum to the methods, we identified a subset of estimates of Z_r in which we 1218 had less confidence because of features of the submitted degrees of freedom. Excluding these effects 1219 in which we had lower confidence had minimal impact on the meta-analytic mean and the estimates 1220 of total I^2 and τ^2 for both blue tit and *Eucalyptus* meta-analyses, regardless of whether outliers were 1221 also excluded (Supplementary Material B, Table B.1).

1222 Post hoc analysis: Exploring the effect of excluding effects from blue

1223 tit models that contained two highly collinear predictors

1224 Effect sizes (Z_r)

- 1225 Excluding effects from blue tit models that contained the two highly collinear predictors (brood count
- 1226 after manipulation and brood count at day 14) had negligible impacts on the meta-analytic means

- 1227 (Table 4), the distribution of Z_r effects (Supplementary Material B, Figure B.5), or heterogeneity
- 1228 estimates (Table 2), which remained high.

1229 Out-of-sample predictions

1230 Inclusion of collinear predictors does not harm model prediction, and so we did not conduct these1231 post hoc analyses.

1232 Explaining Variation in Deviation Scores

None of the pre-registered predictors explained substantial variation in deviation among submittedstatistical effects from the meta-analytic mean (Table 5, Table 6).

- 1235 Table 5: Summary metrics for registered models seeking to explain deviation (Box-Cox transformed
- absolute deviation scores) from \overline{Z}_r as a function of Sorensen's Index, categorical peer ratings, and
- 1237 continuous peer ratings for blue tit and *Eucalyptus* analyses, and as a function of the presence or
- absence of random effects (in the analyst's models) for *Eucalyptus* analyses. We report coefficient of
- determination, R², for our models including only fixed effects as predictors of deviation, and we
- 1240 report R²_{Conditional}, R²_{Marginal} and the intra-class correlation (ICC) from our models that included both
- 1241 fixed and random effects. For all our models, we calculated the residual standard deviation σ and
- 1242 root mean squared error (RMSE).

Dataset	NObs	R ²	R ² _{Conditional}	R ² _{Marginal}	ICC	σ	RMSE				
Deviation ex	Deviation explained by categorical ratings										
Eucalyptus	346		0.13	0.01	0.12	1.06	1.02				
blue tit	473		0.09	7.47 × 10 ⁻³	0.08	0.5	0.48				
Deviation ex	plained by c	ontinuous rati	ngs								
Eucalyptus	346		0.12	7.44 × 10 ⁻³	0.11	1.06	1.03				
blue tit	473		0.09	3.44 × 10 ⁻³	0.09	0.5	0.48				
Deviation ex	plained by S	Sorensen's inde	ex								
Eucalyptus	79	1.84×10^{-4}				1.12	1.1				
blue tit	131	6.32 × 10 ⁻³				0.51	0.51				
Deviation ex	Deviation explained by inclusion of random effects										
Eucalyptus	79	8.75 × 10 ⁻⁸				1.12	1.1				

1243

- 1244 Table 6: Parameter estimates from models of Box-Cox transformed deviation scores from \bar{Z}_r as a
- 1245 function of continuous and categorical peer ratings, Sorensen scores, and the inclusion of random
- 1246 effects. Standard Errors (SE), 95% confidence intervals (95% CI) are reported for all estimates, while t
- 1247 values, degrees of freedom and p-values are presented for fixed-effects. Note that positive

1248 parameter estimates mean that as the predictor variable increases, so does the absolute value of the

1249 deviation from the meta-analytic mean.

Parameter	Random	Coefficient	SE	95% CI	t	df	р		
	effect								
Deviation explained	by inclusion of ra	andom effects -	Eucalypti	IS					
(Intercept)		-2.53	0.27	[-3.06, -1.99]	-9.31	77	<0.001		
Mixed model		0.00	0.31	[-0.60, 0.60]	0.00	77	>0.9		
Deviation explained I	by Sorensen's in	dex - <i>Eucalyptus</i>	;						
(Intercept)		-2.65	1.05	[-4.70, -0.60]	-2.53	77	0.011		
Mean Sorensen's		0.18	1 5 1	[-2.78, 3.14]	0.12	77	>0.9		
index		0.10	1.51	[-2.76, 3.14]	0.12	//	20.9		
Deviation explained l	Deviation explained by Sorensen's index - blue tit								

(Intercept)		-1.53	0.28	[-2.08, -0.98]	-5.42	129	< 0.001
Mean Sorensen's index		0.42	0.47	[-0.49, 1.34]	0.91	129	0.4
Deviation explained	by continuous ra	atings - Eucalypt	us				•
(Intercept)		-2.23	0.23	[-2.69, -1.78]	-9.65	342	< 0.001
RateAnalysis		-0.004	0	[-0.011, 0]	-1.44	342	0.15
SD (Intercept)	Reviewer ID	0.37	0.09	[0.24, 0.60]			
SD (Observations)	Residual	1.06	0.04	[0.98, 1.15]			
Deviation explained	by continuous ra	atings - blue tit					
(Intercept)		-1.16	0.11	[-1.37, -0.94]	-10.60	469	< 0.001
RateAnalysis		-0.002	0	[-0.004, 0]	-1.22	469	0.2
SD (Intercept)	Reviewer ID	0.16	0.03	[0.10,0.24]			
SD (Observations)	Residual	0.5	0.02	[0.46,0.53]			
Deviation explained	by categorical ra	tings - Eucalypt	us				
(Intercept)		-2.66	0.27	[-3.18, -2.13]	-9.97	340	< 0.001
Publishable with		0.29	0.29	[-0.27, 0.85]	1.02	340	0.3
major revision		0.25	0.25	[0.27, 0.05]	1.02	540	0.5
Publishable with		0.01	0.28	[-0.54, 0.56]	0.04	340	>0.9
minor revision							
Publishable as is		0.05	0.31	[-0.55, 0.66]	0.17	340	0.9
SD (Intercept)	Reviewer ID	0.39	0.09	[0.25, 0.61]			
SD (Observations)	Residual	1.06	0.04	[0.98, 1.15]			
Deviation explained	by categorical ra	tings - blue tit			•	T	
(Intercept)		-1.11	0.11	[-1.33, -0.89]	-9.91	467	< 0.001
Publishable with		-0.19	0.12	[-0.42, 0.04]	-1.62	467	0.10
major revision		0.20	0.11	[0112) 010 1]			0.20
Publishable with		-0.19	0.12	[-0.42, 0.04]	-1.65	467	0.10
minor revision			_	. , .		_	
Publishable as is		-0.13	0.13	[-0.39, 0.12]	-1.02	467	0.3
SD (Intercept)	Reviewer ID	0.15	0.04	[0.10, 0.24]			
SD (Observations)	Residual	0.5	0.02	[0.46, 0.53]			

1250

1251 Deviation scores as explained by reviewer ratings

1252 Effect sizes (Z_r)

We obtained reviews from 153 reviewers who reviewed analyses for a mean of 3.27 (range 1 - 11)
analysis teams. Analyses of the blue tit dataset received a total of 240 reviews, each was reviewed by
a mean of 3.87 (SD 0.71, range 3-5) reviewers. Analyses of the *Eucalyptus* dataset received a total of
178 reviews, each was reviewed by a mean of 4.24 (SD 0.79, range 3-6) reviewers. We tested for
inter-rater-reliability (IRR) to examine how similarly reviewers reviewed each analysis and found
approximately no agreement among reviewers. When considering continuous ratings, IRR was 0.01,
and for categorical ratings, IRR was -0.14.

1260 Many of the models of deviation as a function of peer ratings faced issues of failure to converge or

singularity due to sparse design matrices with our pre-registered random effects (Effect ID and

1262 Reviewer ID) (see Supplementary Material C). These issues persisted after increasing the tolerance

and changing the optimizer. For both *Eucalyptus* and blue tit datasets, models with continuous

1264 ratings as a predictor were singular when both pre-registered random effects were included.

1265 When using both categorical and continuous ratings as predictors, only models converged and 1266 allowed 95% confidence intervals to be calculated when specifying Reviewer ID as a random effect. 1267 The categorical ratings model had a R_{C}^{2} of 0.09 and a R_{M}^{2} of 0.01, the continuous ratings model had 1268 a R_{C}^{2} of 0.09 and a R_{M}^{2} of 0.01 for the blue tit dataset and a R_{C}^{2} of 0.12 and a R_{M}^{2} of 0.01 for the

1269 *Eucalyptus* dataset. Neither continuous or categorical reviewer ratings of the analyses meaningfully

predicted deviance from the meta-analytic mean (Table 6, Figure 4). We re-ran the multi-level meta-analysis with a fixed effect for the categorical publishability ratings and found no difference in mean

1272 standardised effect sizes among publishability ratings (<u>Supplementary Material B, Figure B.1</u>).



1273

1274 Figure 4: Violin plot of Box-Cox transformed deviation from meta-analytic mean \overline{Z}_r as a function of

1275 categorical peer rating. Grey points for each rating group denote model-estimated marginal mean

1276 deviation, and error bars denote 95%CI of the estimate. **A** Blue tit dataset, **B** *Eucalyptus* dataset.

1277 Out-of-sample predictions (y_i)

1278 Some models of the influence of reviewer ratings on out-of-sample predictions (y_i) had issues with

1279 convergence and singularity of fit (see <u>Supplementary Material C, Table C.3</u>) and those models that

1280 converged and were not singular showed no strong relationship (Supplementary Material C,

1281 Figure C.2, Supplementary Material C, Figure C.3), as with the *Z_r* analyses.

1282 Deviation scores as explained by the distinctiveness of variables in

1283 each analysis

1284 Effect sizes (Z_r)

1285 We employed Sorensen's index to calculate the distinctiveness of the set of predictor variables used 1286 in each model (Figure 5). The mean Sorensen's score for blue tit analyses was 0.59 (SD: 0.1, range

1287 0.43-0.86), and for *Eucalyptus* analyses was 0.69 (SD: 0.08, range 0.55-0.98).

1288 We found no meaningful relationship between distinctiveness of variables selected and deviation

1289 from the meta-analytic mean (Table 6, Figure 5) for either blue tit (mean 0.42, 95%CI -0.49,1.34)

1290 or *Eucalyptus* effects (mean 0.18, 95%CI -2.78, 3.14).



1291

Figure 5: Fitted model of the Box-Cox-transformed deviation score (deviation in effect size from meta-analytic mean) as a function of the mean Sorensen's index showing distinctiveness of the set of predictor variables. Grey ribbons on predicted values are 95% Cl's. A) blue tit dataset, B) *Eucalyptus* dataset.

1296 Out-of-sample predictions (y_i)

1297 As with the Z_r estimates, we did not observe any convincing relationships between deviation scores

1298 of out-of-sample predictions and Sorensen's index values (see <u>Supplementary Material C4.1</u>).

1299 Deviation scores as explained by the inclusion of random effects

1300 Effect sizes (Z_r)

1301 There were only three blue tit analyses that did not include random effects, which is below the pre-

1302 registered threshold for fitting a model of the Box-Cox transformed deviation from the meta-analytic

- 1303 mean as a function of whether the analysis included random-effects. However,
- 1304 17 *Eucalyptus* analyses included only fixed effects, which crossed our pre-registered threshold.
- 1305 Consequently, we performed this analysis for the *Eucalyptus* dataset only. There was no relationship
- 1306 between random-effect inclusion and deviation from meta-analytic mean among
- 1307 the *Eucalyptus* analyses (Table 6, Figure 6).



Random Effects Included

1309 Figure 6: Violin plot of mean Box-Cox transformed deviation from meta-analytic mean as a function

1310 of random-effects inclusion in *Eucalyptus* analyses. White point for each group of analyses denotes

1311 model-estimated marginal mean deviation, and error bars denote 95% CI of the estimate.

1312 Out-of-sample predictions (y_i)

1308

1313 As with the Z_r estimates, we did not examine the possibility of a relationship between the inclusion

1314 of random effects and the deviation scores of the blue tit out-of-sample predictions. When we

examined the possibility of this relationship for the *Eucalyptus* effects, we found consistent evidence

1316 of somewhat higher Box-Cox-transformed deviation values for models including a random effect,

1317 meaning the models including random effects averaged slightly higher deviation from the meta-

1318 analytic means (Supplementary Material C, Figure C.5).

1319 Multivariate Analysis Effect size (Z_r) and out-of-sample predictions (y_i)

1320 Like the univariate models, the multivariate models did a poor job of explaining deviations from the

1321 meta-analytic mean. Because we pre-registered a multivariate model that contained collinear

- 1322 predictors that produce results which are not readily interpretable, we present these models in the
- supplement. We also had difficulty with convergence and singularity for multivariate models of out-
- 1324 of-sample (*y_i*) result, and had to adjust which random effects we included (<u>Supplementary Material</u>
- 1325 <u>C, Table C.8</u>). However, no multivariate analyses of *Eucalyptus* out-of-sample results avoided
- 1326 problems of convergence or singularity, no matter which random effects we included
- 1327 (Supplementary Material C, Table C.8). We therefore present no multivariate *Eucalyptus y_i* models.
- 1328 We present parameter estimates from multivariate *Z_r* models for both datasets (Supplementary
- 1329 Material C, <u>Table C.6</u>, <u>Table C.7</u>) and from *y_i* models from the blue tit dataset (Supplementary
- 1330 Material C, <u>Table C.10</u>, <u>Table C.9</u>). We include interpretation of the results from these models in the

1331 supplement, but the results do not change the interpretations we present above based on the1332 univariate analyses.

1333 Discussion

1334 When a large pool of ecologists and evolutionary biologists analyzed the same two datasets to 1335 answer the corresponding two research questions, they produced substantially heterogeneous sets 1336 of answers. Although the variability in analytical outcomes was high for both datasets, the patterns 1337 of this variability differed distinctly between them. For the blue tit dataset, there was nearly 1338 continuous variability across a wide range of Z_r values. In contrast, for the *Eucalyptus* dataset, there 1339 was less variability across most of the range, but more striking outliers at the tails. Among out-of-1340 sample predictions, there was again almost continuous variation across a wide range (2 SD) among 1341 blue tit estimates. For *Eucalyptus*, out-of-sample predictions were also notably variable, with about 1342 half the predicted stem count values at <2 but the other half being much larger, and ranging to 1343 nearly 40 stems per 15 m x 15 m plot. We investigated several hypotheses for drivers of this 1344 variability within datasets, but found little support for any of these. Most notably, even when we 1345 excluded analyses that had received one or more poor peer reviews, the heterogeneity in results 1346 largely persisted. Regardless of what drives the variability, the existence of such dramatically 1347 heterogeneous results when ecologists and evolutionary biologists seek to answer the same 1348 questions with the same data should trigger conversations about how ecologists and evolutionary 1349 biologists analyze data and interpret the results of their own analyses and those of others in the 1350 literature (e.g., Silberzahn et al. 2018; Simonsohn, Simmons, and Nelson 2020; Auspurg and Brüderl 1351 2021; Breznau et al. 2022).

1352 Our observation of substantial heterogeneity due to analytical decisions is consistent with a small 1353 earlier study in ecology (Stanton-Geddes, de Freitas and de Sales Dambros 2014) and a growing body 1354 of work from the quantitative social sciences (e.g., Silberzahn et al. 2018; Botvinik-Nezer et al. 1355 2020; Huntington-Klein et al. 2021; Schweinsberg et al. 2021; Breznau et al. 2022; Coretta et al. 1356 2023). In these studies, when volunteers from the discipline analyzed the same data, they produced 1357 a worryingly diverse set of answers to a pre-set question. This diversity included a wide range of 1358 effect sizes, and in most cases, even involved effects in opposite directions. Thus, our result should 1359 not be viewed as an anomalous outcome from two particular datasets, but instead as evidence from 1360 additional disciplines regarding the heterogeneity that can emerge from analyses of complex 1361 datasets to answer questions in probabilistic science. Not only is our major observation consistent 1362 with other studies, it is, itself, robust because it derived primarily from simple forest plots that we 1363 produced based on a small set of decisions that were mostly registered before data gathering and 1364 which conform to widely accepted meta-analytic practices.

1365 Unlike the strong pattern we observed in the forest plots, our other analyses, both registered 1366 and post hoc, produced either inconsistent patterns, weak patterns, or the absence of patterns. Our 1367 registered analyses found that deviations from the meta-analytic mean by individual effect sizes (Zr)1368 or the predicted values of the dependent variable (\bar{y}) were poorly explained by our hypothesized 1369 predictors: peer rating of each analysis team's method section, a measurement of the distinctiveness 1370 of the set of predictor variables included in each analysis, or whether the model included random 1371 effects. However, in our post hoc analyses, we found that dropping analyses identified as 1372 unpublishable or in need of major revision by at least one reviewer modestly reduced the observed 1373 heterogeneity among the Z_r outcomes, but only for *Eucalyptus* analyses, apparently because this led 1374 to the dropping of the major outlier. This limited role for peer review in explaining the variability in

1375 our results should be interpreted cautiously because the inter-rater reliability among peer reviewers 1376 was extremely low, and at least some analyses that appeared flawed to us were not marked as 1377 flawed by reviewers. Thus, it seems that the peer reviews we received were of mixed quality, possibly 1378 due to lack of expertise or lack of care on the part of some reviewers. However, the hypothesis that 1379 poor quality analyses drove a substantial portion of the heterogeneity we observed was also 1380 contradicted by our observation that analysts' self-declared statistical expertise appeared unrelated 1381 to heterogeneity. When we retained only analyses from teams including at least one member with 1382 high self-declared levels of expertise, heterogeneity among effect sizes remained high. Thus, our 1383 results suggest lack of statistical expertise is not the primary factor responsible for the heterogeneity 1384 we observed, although further work is merited before rejecting a role for statistical expertise. 1385 Besides variability in expertise, it is also possible that the volunteer analysts varied in the effort they 1386 invested, and low effort presumably drove at least some heterogeneity in results. However, analysts 1387 often submitted thoughtful and extensive code, tables, figures, and textual explanation and 1388 interpretations, which is evidence of substantial investment. Further, we are confident that low effort 1389 alone is an insufficient explanation for the heterogeneity we observed because we have worked with 1390 these datasets ourselves, and we know from experience that there are countless plausible modeling 1391 alternatives that can produce a diversity of effects. Additionally, heterogeneity in analytical outcomes 1392 differed notably between datasets, and there is no reason to expect that one set of analysts took this 1393 project less seriously than the other. Returning to our exploratory analyses, not surprisingly, simply 1394 dropping outlier values of Z_r for Eucalyptus analyses, which had more extreme outliers, led to less 1395 observable heterogeneity in the forest plots, and also reductions in our quantitative measures of 1396 heterogeneity. We did not observe a similar effect in the blue tit dataset because that dataset had 1397 outliers that were much less extreme and instead had more variability across the core of the 1398 distribution.

1399 Our major observations raise two broad questions; why was the variability among results so high, 1400 and why did the pattern of variability differ between our two datasets. One important and plausible 1401 answer to the first question is that much of the heterogeneity derives from the lack of a precise 1402 relationship between the two biological research questions we posed and the data we provided. This 1403 lack of a precise relationship between data and question creates many opportunities for different 1404 model specifications, and so may inevitably lead to varied analytical outcomes (Auspurg and Brüderl 1405 2021). However, we believe that the research questions we posed are consistent with the kinds of 1406 research question that ecologists and evolutionary biologists typically work from. When designing 1407 the two biological research questions, we deliberately sought to represent the level of specificity we 1408 typically see in these disciplines. This level of specificity is evident when we look at the research 1409 questions posed by some recent meta-analyses in these fields:

- "how [does] urbanisation impact mean phenotypic values and phenotypic variation ... [in]
 paired urban and non-urban comparisons of avian life-history traits" (Capilla-Lasheras et al.
 2022)
- "[what are] the effects of ocean acidification on the crustacean exoskeleton, assessing both
 exoskeletal ion content (calcium and magnesium) and functional properties (biomechanical
 resistance and cuticle thickness)" (Siegel et al. 2022)
- "[what is] the extent to which restoration affects both the mean and variability of
 biodiversity outcomes ... [in] terrestrial restoration" (Atkinson et al. 2022)
- 1418 "[does] drought stress [have] a negative, positive, or null effect on aphid fitness" (Leybourne et al. 2021)

"[what is] the influence of nitrogen-fixing trees on soil nitrous oxide emissions" (Kou Giesbrecht and Menge 2021)

1422 There is not a single precise answer to any of these questions, nor to the questions we posed to 1423 analysts in our study. And this lack of single clear answers will obviously continue to cause 1424 uncertainty since ecologists and evolutionary biologists conceive of the different answers from the 1425 different statistical models as all being answers to the same general question. A possible response 1426 would be a call to avoid these general questions in favor of much more precise alternatives (Auspurg 1427 and Brüderl 2021). However, the research community rewards researchers who pose broad 1428 questions (Simons, Shoda, and Lindsay 2017), and so researchers are unlikely to narrow their scope 1429 without a change in incentives. Further, we suspect that even if individual studies specified narrow 1430 research questions, other scientists would group these more narrow questions into broader 1431 categories, for instance in meta-analyses, because it is these broader and more general questions 1432 that often interest the research community.

1433 Although variability in statistical outcomes among analysts may be inevitable, our results raise 1434 questions about why this variability differed between our two datasets. We are particularly 1435 interested in the differences in the distribution of Z_r since the distributions of out-of-sample 1436 predictions were on different scales for the two datasets, thus limiting the value of comparisons. The 1437 forest plots of Z_r from our two datasets showed distinct patterns, and these differences are 1438 consistent with several alternative hypotheses. The results submitted by analysts of 1439 the *Eucalyptus* dataset showed a small average (close to zero) with most estimates also close to zero 1440 (± 0.2), though about a third far enough above or below zero to cross the traditional threshold of 1441 statistical significance. There were a small number of striking outliers that were very far from zero. In 1442 contrast, the results submitted by analysts of the blue tit dataset showed an average much further 1443 from zero (-0.35) and a much greater spread in the core distribution of estimates across the range 1444 of Z_r values (± 0.5 from the mean), with few modest outliers. So, why was there more spread in effect 1445 sizes (across the estimates that are not outliers) in the blue tit analyses relative to 1446 the Eucalyptus analyses?

1447 One possible explanation for the lower heterogeneity among most *Eucalyptus Z_r* effects is that weak 1448 relationships may limit the opportunities for heterogeneity in analytical outcome. Some evidence for 1449 this idea comes from two sets of "many labs" studies in psychology (Klein et al. 2014, 2018). In these 1450 studies, many independent lab groups each replicated a large set of studies, including, for each 1451 study, the experiment, data collection, and statistical analyses. These studies showed that, when the 1452 meta-analytic mean across the replications from different labs was small, there was much less 1453 heterogeneity among the outcomes than when the mean effect sizes were large (Klein et al. 1454 2014, 2018). Of course, a weak average effect size would not prevent divergent effects in all 1455 circumstances. As we saw with the *Eucalyptus* analyses, taking a radically smaller subset of the data 1456 can lead to dramatically divergent effect sizes even when the mean with the full dataset is close to 1457 zero.

1458 Our observation that dramatic sub-setting in the *Eucalyptus* dataset was associated with 1459 correspondingly dramatic divergence in effect sizes leads us towards another hypothesis to explain 1460 the differences in heterogeneity between the *Eucalyptus* and blue tit analysis sets. It may be that 1461 when analysts often divide a dataset into subsets, the result will be greater heterogeneity in 1462 analytical outcome for that dataset. Although we saw sub-setting associated with dramatic outliers in 1463 the Eucalyptus dataset, nearly all other analyses of Eucalyptus data used close to the same set of 351 1464 samples, and as we saw, these effects did not vary substantially. However, analysts often analyzed 1465 only a subset of the blue tit data, and as we observed, sample sizes were much more variable among

blue tit effects, and the effects themselves were also much more variable. Important to note here is
that subsets of data may differ from each other for biological reasons, but they may also differ due to
sampling error. Sampling error is a function of sample size, and sub-samples are, by definition,
smaller samples, and so more subject to variability in effects due to sampling error (Jennions et al.
2013).

1471 Other features of datasets are also plausible candidates for driving heterogeneity in analytical 1472 outcomes, including features of covariates. In particular, relationships between covariates and the 1473 response variable as well as relationships between covariates and the primary independent variable 1474 (collinearity) can strongly influence the modeled relationship between the independent variable of 1475 interest and the dependent variable (Morrissey and Ruxton 2018; Dormann et al. 2013). Therefore, 1476 inclusion or exclusion of these covariates can drive heterogeneity in effect sizes (Z_r). Also, as we saw 1477 with the two most extreme Z_r values from the blue tit analyses, in multivariate models with collinear 1478 predictors, extreme effects can emerge when estimating partial correlation coefficients due to high 1479 collinearity, and conclusions can differ dramatically depending on which relationship receives the 1480 researcher's attention. Therefore, differences between datasets in the presence of strong and/or 1481 collinear covariates could influence the differences in heterogeneity in results among those datasets.

Although it is too early in the many-analyst research program to conclude which analytical decisions
or which features of datasets are the most important drivers of heterogeneity in analytical outcomes,
we must still grapple with the possibility that analytical outcomes may vary substantially based on
the choices we make as analysts. If we assume that, at least sometimes, different analysts will
produce dramatically different statistical outcomes, what should we do as ecologists and
evolutionary biologists? We review some ideas below.

1488 The easiest path forward after learning about this analytical heterogeneity would be simply to 1489 continue with "business as usual", where researchers report results from a small number of statistical 1490 models. A case could be made for this path based on our results. For instance, among the blue tit 1491 analyses, the precise values of the estimated Z_r effects varied substantially, but the average effect 1492 was convincingly different from zero, and a majority of individual effects (84%) were in the same 1493 direction. Arguably, many ecologists and evolutionary biologists appear primarily interested in the 1494 direction of a given effect and the corresponding p-value (Fidler et al. 2006), and so the variability we 1495 observed when analyzing the blue tit dataset may not worry these researchers. Similarly, most 1496 effects from the Eucalyptus analyses were relatively close to zero, and about two-thirds of these 1497 effects did not cross the traditional threshold of statistical significance. Therefore, a large proportion 1498 of people analyzing these data would conclude that there was no effect, and this is consistent with what we might conclude from the meta-analysis. 1499

1500 However, we find the counter arguments to "business as usual" to be compelling. For blue tits, there 1501 were a substantial minority of calculated effects that would be interpreted by many biologists as 1502 indicating the absence of an effect (28%), and there were three traditionally 'significant' effects in 1503 the opposite direction to the average. The qualitative conclusions of analysts also reflected 1504 substantial variability, with fully half of teams drawing a conclusion distinct from the one we draw 1505 from the distribution as a whole. These teams with different conclusions were either uncertain about 1506 the negative relationship between competition and nestling growth, or they concluded that effects 1507 were mixed or absent. For the Eucalyptus analyses, this issue is more concerning. Around two-thirds 1508 of effects had confidence intervals overlapping zero, and of the third of analyses with confidence 1509 intervals excluding zero, almost half were positive, and the rest were negative. Accordingly, the 1510 qualitative conclusions of the Eucalyptus teams were spread across the full range of possibilities. But,

as we describe in the next paragraph, even this striking lack of consensus may be much less of aproblem than what could emerge as scientists select which results to publish.

1513 A potentially larger argument against "business as usual" is that it provides the raw material for 1514 biasing the literature. When different model specifications readily lead to different results, analysts 1515 may be tempted to report the result that appears most interesting, or that is most consistent with 1516 expectation (Gelman and Loken 2013; Forstmeier, Wagenmakers and Parker 2017). There is growing 1517 evidence that researchers in ecology and evolutionary biology often report a biased subset of the 1518 results they produce (Deressa et al. 2023; Kimmel, Avolio and Ferraro 2023), and that this bias 1519 exaggerates the average size of effects in the published literature between 30 and 150% (Yang et al. 1520 2023; Parker and Yang 2023). The bias then accumulates in meta-analyses, apparently more than 1521 doubling the rate of conclusions of "statistical significance" in published meta-analyses above what 1522 would have been found in the absence of bias (Yang et al. 2023). Thus, "business as usual" does not 1523 just create noisy results, it helps create systematically misleading results.

1524 If we move away from "business as usual", where do we go? Many obvious options involve multiple 1525 analyses per dataset. For instance, there is the traditional robustness or sensitivity check (e.g., Pei et 1526 al. 2020; Briga and Verhulst 2021), in which the researcher presents several alternative versions of an 1527 analysis to demonstrate that the result is 'robust' (Lu and White 2014). Unfortunately, robustness 1528 checks are at risk of the same potential biases of reporting found in other studies (Silberzahn et al. 1529 2018), especially given the relatively few models typically presented. However, these risks could be 1530 minimized by running more models and doing so with a pre-registration or registered report. 1531 Another option is model averaging. Averages across models often perform well (e.g. Taylor and Taylor 1532 2023), and in some forms this may be a relatively simple solution. Model averaging, as most often 1533 practiced in ecology and evolutionary biology, involves first identifying a small suite of candidate 1534 models (see Burnham and Anderson 2002), then using Akaike weights, based on Akaike's Information 1535 Criterion (AIC), to calculate weighted averages for parameter estimates from those models. As with 1536 typical robustness checks, the small number of models limits the exploration of specification space, 1537 but examining a larger number of models could become the norm. However, there are more 1538 concerning limitations. The largest of these limitations is that averaging regression coefficients is 1539 problematic when models differ in interaction terms or collinear variables (Cade 2015). Additionally, 1540 weighting by AIC may often be inconsistent with our modelling goals. AIC balances the trade-off 1541 between model complexity and predictive ability, but penalizing models for complexity may not be 1542 suited for testing hypotheses about causation (Arif and MacNeil 2022). So, AIC may often not offer 1543 the weight we want to use, and we may also not wish to just generate an average at all. Instead, if we 1544 hope to understand an extensive universe of possible modelling outcomes, we could conduct a 1545 multiverse analysis, possibly with a specification curve (Simonsohn, Simmons, and Nelson 1546 2015, 2020). This could mean running hundreds or thousands of models (or more!) to examine the 1547 distribution of possible effects, and to see how different model specification choices map onto these 1548 effects. However, exploring large areas of specification space may come at the cost of including 1549 biologically implausible specifications. Thus, we expect a trade-off, and attempts to limit models to 1550 the most biologically plausible may become increasingly difficult in proportion to the number of 1551 variables and modeling choices. To make selecting plausible models easier, one could recruit multiple 1552 analysts to design one or a few plausible specifications each as with our 'many analyst' 1553 study (Silberzahn et al. 2018). An alternative that may be more labor intensive for the primary 1554 analyst, but which may lead to a more plausible set of models, could involve hypothesizing about 1555 causal pathways with DAGs [directed acyclic graphs; Arif and MacNeil (2023)] to constrain the model 1556 set. As with other options outlined above, generating model specifications with DAGs could be 1557 partnered with pre-registration to hinder bias from undisclosed data dredging.

1558 Responses to heterogeneity in analysis outcomes need not be limited to simply conducting more 1559 analyses, especially if it turns out that analysis quality drives some of the observed heterogeneity. As 1560 we noted above, we cannot yet rule out the possibility that insufficient statistical expertise or poor-1561 quality analyses might drive some portion of the heterogeneity we observed. Improving the quality of analyses might be accomplished with a deliberate increase in investment in statistical education. 1562 1563 Many ecology and evolutionary biology students learn their statistical practice informally, with many 1564 ecology doctoral programs in the USA not requiring a statistics course (Touchon and McCoy 2016), 1565 and no formal courses of any kind included in doctoral degrees in most other countries. In cases 1566 where formal investment in statistical education is lacking, informal resources, such as guidelines and 1567 checklists, may help researchers avoid common mistakes. However, unless following guidelines or 1568 checklists is enforced for publication, the adherence to guidelines is patchy. For example, despite the 1569 publication of guidelines for conducting meta-analyses in ecology, the quality of meta-analyses did 1570 not improve substantially over time (Koricheva and Gurevitch 2014). Even in medical research where adherence to guidelines such as the PRISMA standards for systematic reviews and meta-analyses is 1571 1572 more highly valued, adherence is often poor (Page and Moher 2017).

1573 Although we have reviewed a variety of potential responses to the existence of variability in 1574 analytical outcomes, we certainly do not wish to imply that this is a comprehensive set of possible 1575 responses. Nor do we wish to imply that the opinions we have expressed about these options are 1576 correct. Determining how the disciplines of ecology and evolutionary biology should respond to 1577 knowledge of the variability in analytical outcome will benefit from the contribution and discussion 1578 of ideas from across these disciplines. We look forward to learning from these discussions and to 1579 seeing how these disciplines ultimately respond.

1580 Conclusions

1581 Overall, our results suggest to us that, where there is a diverse set of plausible analysis options, no 1582 single analysis should be considered a complete or reliable answer to a research question. Further, 1583 because of the evidence that ecologists and evolutionary biologists often present a biased subset of 1584 the analyses they conduct (Deressa et al. 2023; Yang et al. 2023; Kimmel, Avolio and Ferraro 2023), 1585 we do not expect that even a collection of different effect sizes from different studies will accurately 1586 represent the true distribution of effects (Yang et al. 2023). Therefore, we believe that an increased 1587 level of skepticism of the outcomes of single analyses, or even single meta-analyses, is warranted 1588 going forward. We recognize that some researchers have long maintained a healthy level of 1589 skepticism of individual studies as part of sound and practical scientific practice, and it is possible 1590 that those researchers will be neither surprised nor concerned by our results. However, we doubt 1591 that many researchers are sufficiently aware of the potential problems of analytical flexibility to be 1592 appropriately skeptical. We hope that our work leads to conversations in ecology, evolutionary 1593 biology, and other disciplines about how best to contend with heterogeneity in results that is 1594 attributable to analytical decisions.

Appendix 1595

1596 R Package References and Session Information. Table 7 shows all R packages and their versions used

1597 in the production of the manuscript.

1598 Table 7: R packages used to generate this manuscript. Please see the ManyEcoEvo:: package for a full

1599 list of packages used in the analysis pipeline.

Package	Version	Citation
base	4.4.0	R Core Team (2024)
betapart	1.6	Baselga et al. (2023)
broom.mixed	0.2.9.5	<u>Bolker et al. (2024)</u>
colorspace	2.1.0	Zeileis et al. (2020)
cowplot	1.1.3	<u>Wilke (2024)</u>
devtools	2.4.5	Wickham et al. (2022)
EnvStats	2.8.1	Millard (2013)
GGally	2.2.1	<u>Schloerke et al. (2024)</u>
ggforestplot	0.1.0	Scheinin et al. (2020)
ggh4x	0.2.8	van den Brand (2024)
ggpubr	0.6.0	Kassambara (2023)
ggrepel	0.9.5	<u>Slowikowski (2024)</u>
ggthemes	5.1.0	<u>Arnold (2024)</u>
glmmTMB	1.1.8	Brooks et al. (2017)
gt	0.10.1	lannone et al. (2024)
gtsummary	1.7.2	Sjoberg et al. (2021)
here	1.0.1	<u>Müller (2020)</u>
Hmisc	5.1.2	Harrell Jr (2024)
irr	0.84.1	Gamer, Lemon, and Singh (2019)
janitor	2.2.0	<u>Firke (2023)</u>
knitr	1.46	<u>Xie (2024a)</u>
latex2exp	0.9.6	Meschiari (2022)
lme4	1.1.35.3	Bates et al. (2015)
ManyEcoEvo	2.7.6	<u>Gould et al. (2023)</u>
metafor	4.6.0	Viechtbauer (2010)
modelbased	0.8.7	Makowski et al. (2020)
multilevelmod	1.0.0	Kuhn and Frick (2022)
MuMIn	1.47.5	Bartoń (2023)
naniar	1.1.0	Tierney and Cook (2023)
NatParksPalettes	0.2.0	<u>Blake (2022)</u>
orchaRd	2	Nakagawa, Lagisz, et al. (2023)
parameters	0.21.7	Lüdecke et al. (2020)
patchwork	1.2.0	Pedersen (2024)
performance	0.11.0	Lüdecke, Ben-Shachar, et al. (2021)
renv	1.0.2	Ushey and Wickham (2023)

rmarkdown	2.27	Allaire et al. (2024)
sae	1.3	Molina and Marhuenda (2015)
scales	1.3.0	Wickham, Pedersen, and Seidel (2023)
see	0.8.4	Lüdecke, Patil, et al. (2021)
showtext	0.9.7	<u>Qiu (2024)</u>
specr	1.0.0	Masur and Scharkow (2020)
targets	1.7.0	Landau (2021)
tidymodels	1.1.1	Kuhn and Wickham (2020)
tidytext	0.4.2	Silge and Robinson (2016)
tidyverse	2.0.0	Wickham et al. (2019)
withr	3.0.0	Hester et al. (2024)
xfun	0.44	<u>Xie (2024b)</u>

1601	
1602	– Session info —————————————————————
1603	setting value
1604	version R version 4.4.0 (2024-04-24)
1605	os macOS Ventura 13.6.9
1606	system aarch64, darwin20
1607	ui X11
1608	language (EN)
1609	collate en_US.UTF-8
1610	ctype en_US.UTF-8
1611	tz Australia/Melbourne
1612	date 2024-09-17
1613	pandoc 3.1.12.2 @ /opt/homebrew/bin/ (via rmarkdown)
1614	

Declarations 1615

Ethics, consent and permissions 1616

1617 We obtained permission to conduct this research from the Whitman College Institutional Review 1618 Board (IRB). As part of this permission, the IRB approved the consent form (https://osf.io/xyp68/) 1619 that all participants completed prior to joining the study. The authors declare that they have no 1620 competing interests.

Availability of data and materials 1621

1622 All materials and data are archived and hosted on the OSF at https://osf.io/mn5aj/, including survey

1623 instruments and analyst / reviewer consent forms. The Evolutionary Ecology Data and Ecology and

1624 Conservation Data provided to analysts are available

1625 at https://osf.io/34fzc/ and https://osf.io/t76uy/ respectively. Data has been anonymised, and the 1626 non-anonymised data is stored on the project OSF within private components accessible to the lead 1627 authors.

- 1628 We built an R package, ManyEcoEvo to conduct the analyses described in this study (Gould et al.
- 1629 2023), which can be downloaded from https://github.com/egouldo/ManyEcoEvo/ to reproduce our
- 1630 analyses or replicate the analyses described here using alternate datasets. Data cleaning and
- 1631 preparation of analysis-data, as well as the analysis, is conducted in R (R Core Team
- 1632 2024) reproducibly using the targets package (Landau 2021). This data and analysis pipeline is stored
- 1633 in the ManyEcoEvo package repository and its outputs are made available to users of the package
- 1634 when the library is loaded.

- 1635 The full manuscript, including further analysis and presentation of results is written in Quarto (J. J.
- 1636 Allaire et al. 2024). The source code to reproduce the manuscript is hosted
- at https://github.com/egouldo/ManyAnalysts/, and the rendered version of the source code may be 1637
- 1638 viewed at https://egouldo.github.io/ManyAnalysts/. All R packages and their versions used in the
- 1639 production of this manuscript are listed in the session info at Section 6.6.

Competing interests 1640

1641 The authors declare that they have no competing interests

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1651

Author's contributions 1652

1653 HF, THP and FF conceptualized the project. PV provided raw data for Eucalyptus analyses and SG and 1654 THP provided raw data for blue tit analyses. DGH, HF and THP prepared surveys for collecting 1655 participating analysts and reviewer's data. EG, HF, THP, PV, SN and FF planned the analyses of the 1656 data provided by our analysts and reviewers, EG, HF, DGH, and THP curated the data, EG and HF 1657 wrote the software code to implement the analyses and prepare data visualisations. EG ensured that 1658 analyses were documented and reproducible. THP and HF administered the project, including 1659 coordinating with analysts and reviewers. FF provided funding for the project. THP, HF, and EG wrote 1660 the manuscript. Authors listed alphabetically contributed analyses of the primary datasets or reviews 1661 of those analyses. All authors read and approved the final manuscript.

References 1662

- 1663 Aczel, Balazs, Barnabas Szaszi, Gustav Nilsonne, Olmo R van den Akker, Casper J Albers, Marcel ALM van Assen, Jojanneke A Bastiaansen, et al. 2021. "Consensus-Based Guidance for Conducting and 1664
- 1665 Reporting Multi-Analyst Studies." eLife 10 (November). https://doi.org/10.7554/elife.72185.
- 1666 Allaire, J. J., Charles Teague, Carlos Scheidegger, Yihui Xie, and Christophe Dervieux.
- 1667 2024. "Quarto." https://doi.org/10.5281/zenodo.5960048.
- 1668 Allaire, JJ, Yihui Xie, Christophe Dervieux, Jonathan McPherson, Javier Luraschi, Kevin Ushey, Aron
- 1669 Atkins, et al. 2024. rmarkdown: Dynamic Documents for r. https://github.com/rstudio/rmarkdown.
- 1670 Arif, S., & M. Aaron MacNeil. 2022. "Predictive models aren't for causal inference." Ecology Letters
- 1671 25(8), 1741–1745. https://doi.org/10.1111/ele.14033

- 1672 Arif, Suchinta, and M. Aaron MacNeil. 2023. "Applying the Structural Causal Model Framework for
- 1673 Observational Causal Inference in Ecology." *Ecological Monographs* 93 (1): e1554.
- 1674 https://doi.org/<u>https://doi.org/10.1002/ecm.1554</u>.
- 1675 Arnold, Jeffrey B. 2024. ggthemes: Extra Themes, Scales and Geoms
- 1676 for "ggplot2". https://jrnold.github.io/ggthemes/.
- 1677 Atkinson, Joe, Lars A. Brudvig, Max Mallen-Cooper, Shinichi Nakagawa, Angela T. Moles, and Stephen
- 1678 P. Bonser. 2022. "Terrestrial Ecosystem Restoration Increases Biodiversity and Reduces Its Variability,
- 1679 but Not to Reference Levels: A Global Meta-Analysis." *Ecology Letters* 25 (7): 1725–37.
- 1680 https://doi.org/<u>https://doi.org/10.1111/ele.14025</u>.
- 1681 Auspurg, Katrin, and Josef Brüderl. 2021. "Has the Credibility of the Social Sciences Been Credibly
- 1682 Destroyed? Reanalyzing the 'Many Analysts, One Data Set' Project." Socius 7:
- 1683 23780231211024421. https://doi.org/10.1177/23780231211024421.
- 1684 Bartoń, Kamil. 2023. *MuMIn: Multi-Model Inference*.
- 1685 Baselga, Andres, David Orme, Sebastien Villeger, Julien De Bortoli, Fabien Leprieur, Maxime Logez,
- 1686 Sara Martinez-Santalla, et al. 2023. *betapart: Partitioning Beta Diversity into Turnover and*
- 1687 Nestedness Components. <u>https://CRAN.R-project.org/package=betapart</u>.
- Bates, Douglas, Martin Mächler, Ben Bolker, and Steve Walker. 2015. "Fitting Linear Mixed-Effects
 Models Using Ime4." 2015 67 (1): 48. https://doi.org/10.18637/jss.v067.i01.
- 1690 Blake, Kevin. 2022. NatParksPalettes: Color Palettes Inspired by National
- 1691 Parks. https://github.com/kevinsblake/NatParksPalettes.
- 1692 Bolker, Ben, David Robinson, Dieter Menne, Jonah Gabry, Paul Buerkner, Chrisopher Hau, William
- 1693 Petry, et al. 2024. broom.mixed: Tidying Methods for Mixed
- 1694 *Models*. <u>https://github.com/bbolker/broom.mixed</u>.
- 1695 Borenstein, Michael, Julian P. T. Higgins, Larry Hedges, and Hannah Rothstein. 2017. "Basics of Meta-
- 1696 Analysis: *I*² Is Not an Absolute Measure of Heterogeneity." *Research Synthesis Methods* 8: 5–
- 1697 18. <u>https://doi.org/10.1002/jrsm.1230</u>.
- Botvinik-Nezer, Rotem, Felix Holzmeister, Colin F. Camerer, Anna Dreber, Juergen Huber, Magnus
 Johannesson, Michael Kirchler, et al. 2020. "Variability in the Analysis of a Single Neuroimaging
 Deteast by Many Teams" (Nature 592 (7010): 04, 99
- 1700 Dataset by Many Teams." *Nature* 582 (7810): 84–88.
- 1701 Breznau, Nate, Eike Mark Rinke, Alexander Wuttke, Hung H. V. Nguyen, Muna Adem, Jule Adriaans,
- 1702 Amalia Alvarez-Benjumea, et al. 2022. "Observing Many Researchers Using the Same Data and
- 1703 Hypothesis Reveals a Hidden Universe of Uncertainty." *Proceedings of the National Academy of*
- 1704 *Sciences* 119 (44): e2203150119. <u>https://doi.org/10.1073/pnas.2203150119</u>.
- Briga, Michael, and Simon Verhulst. 2021. "Mosaic Metabolic Ageing: Basal and Standard Metabolic
 Rates Age in Opposite Directions and Independent of Environmental Quality, Sex and Life Span in a
 Passerine." *Functional Ecology* 35 (5): 1055–68. https://doi.org/<u>https://doi.org/10.1111/1365-</u>
 2435.13785.
- Brooks, Mollie E., Kasper Kristensen, Koen J. van Benthem, Arni Magnusson, Casper W. Berg, Anders
 Nielsen, Hans J. Skaug, Martin Maechler, and Benjamin M. Bolker. 2017. "glmmTMB Balances Speed
 and Flexibility Among Packages for Zero-Inflated Generalized Linear Mixed Modeling." *The R Journal* 9
- 1712 (2): 378–400. <u>https://doi.org/10.32614/RJ-2017-066</u>.

- 1713 Buck, Robert J., John Fieberg, and Daniel J. Larkin. 2022. "The use of weighted averages of Hedges' d
- 1714 in meta-analysis: Is it worth it?" *Methods in Ecology and Evolution* 13 (5): 1093–1105.
- 1715 <u>https://doi.org/10.1111/2041-210X.13818</u>.
- 1716 Burnham, K. P., and D. R. Anderson. 2002. Model Selection and Multimodel Inference: A Practical
- 1717 Information-Theoretical Approach. Book. 2nd ed. New York: Springer-
- 1718 Verlag. <u>https://doi.org/10.1007/b97636</u>.
- 1719 Cade, Brian S. 2015. "Model Averaging and Muddled Multimodel Inferences." Ecology 96 (9): 2370–
- 1720 82. <u>http://www.jstor.org.ezproxy.whitman.edu/stable/24702343</u>.
- 1721 Capilla-Lasheras, Pablo, Megan J. Thompson, Alfredo Sánchez-Tójar, Yacob Haddou, Claire J. Branston,
- 1722 Denis Réale, Anne Charmantier, and Davide M. Dominoni. 2022. "A Global Meta-Analysis Reveals
- 1723 Higher Variation in Breeding Phenology in Urban Birds Than in Their Non-Urban Neighbours." *Ecology*
- 1724 Letters 25 (11): 2552–70. <u>https://doi.org/10.1111/ele.14099</u>.
- 1725 Coretta, Stefano, Joseph V. Casillas, Simon Roessig, Michael Franke, Byron Ahn, Ali H. Al-Hoorie, Jalal
- 1726 Al-Tamimi, et al. 2023. "Multidimensional Signals and Analytic Flexibility: Estimating Degrees of
- 1727 Freedom in Human-Speech Analyses." Advances in Methods and Practices in Psychological Science 6
- 1728 (3): 25152459231162567. <u>https://doi.org/10.1177/25152459231162567</u>.
- 1729 Dancho, Matt, and Davis Vaughan. 2023. *Timetk: A Tool Kit for Working with Time*
- 1730 Series. <u>https://CRAN.R-project.org/package=timetk</u>.
- 1731 DeKogel, C. H. 1997. "Long-Term Effects of Brood Size Manipulation on Morphological Development
- and Sex-Specific Mortality of Offspring." *Journal of Animal Ecology* 66 (2): 167–78. <<u>Go to</u>
 ISI>://WOS:A1997WQ19600003.
- 1734 Deressa, Teshome, David Stern, Jaco Vangronsveld, Jan Minx, Sebastien Lizin, Robert Malina, and
- 1735 Stephan Bruns. 2023. "More Than Half of Statistically Significant Research Findings in the
- 1736 Environmental Sciences Are Actually Not." *EcoEvoRxiv*.
- 1737 https://doi.org/<u>https://doi.org/10.32942/X24G6Z</u>.
- 1738 Dormann, Carsten F., Jane Elith, Sven Bacher, Carsten Buchmann, Gudrun Carl, Gabriel Carré, Jaime
- 1739 R. García Marquéz, et al. 2013. "Collinearity: A Review of Methods to Deal with It and a Simulation
- 1740 Study Evaluating Their Performance." *Ecography* 36 (1): 27–46.
- 1741 https://doi.org/<u>https://doi.org/10.1111/j.1600-0587.2012.07348.x</u>.
- 1742 Fanelli, Daniele, Rodrigo Costas, and John P. A. Ioannidis. 2017. "Meta-Assessment of Bias in
- 1743 Science." Proceedings of the National Academy of Sciences 114: 3714–
- 1744 19. <u>https://doi.org/10.1073/pnas.1618569114</u>.
- 1745 Fanelli, Daniele, and John P. A. Ioannidis. 2013. "US Studies May Overestimate Effect Sizes in Softer
- 1746 Research." Proceedings of the National Academy of Sciences 110 (37): 15031–
- 1747 36. <u>https://doi.org/10.1073/pnas.1302997110</u>.
- 1748 Fidler, Fiona, Mark A. Burgman, Geoff Cumming, Robert Buttrose, and Neil Thomason. 2006. "Impact
- 1749 of Criticism of Null-Hypothesis Significance Testing on Statistical Reporting Practices in Conservation
- 1750 Biology." *Conservation Biology* 20 (5): 1539–44. <u>https://doi.org/10.1111/j.1523-1739.2006.00525.x</u>.
- 1751 Fidler, Fiona, Yung En Chee, Bonnie C. Wintle, Mark A. Burgman, Michael A. McCarthy, and Ascelin
- 1752 Gordon. 2017. "Metaresearch for Evaluating Reproducibility in Ecology and Evolution." *BioScience* 67
- 1753 (3): 282–89. <u>https://doi.org/10.1093/biosci/biw159</u>.

- 1754 Firke, Sam. 2023. janitor: Simple Tools for Examining and Cleaning Dirty
- 1755 Data. <u>https://github.com/sfirke/janitor</u>.
- 1756 Forstmeier, Wolfgang, Eric-Jan Wagenmakers, and T. H. Parker. 2017. "Detecting and Avoiding Likely
- 1757 False-Positive Findings a Practical Guide." Biological Reviews 92: 1941–
- 1758 68. <u>https://doi.org/10.1111/brv.12315</u>.
- 1759 Fraser, Hannah, Tim Parker, Shinichi Nakagawa, Ashley Barnett, and Fiona Fidler. 2018. "Questionable
- 1760 Research Practices in Ecology and Evolution." *PLOS ONE* 13 (7):
- 1761 e0200303. <u>https://doi.org/10.1371/journal.pone.0200303</u>.
- Gamer, Matthias, Jim Lemon, and Ian Fellows Puspendra Singh. 2019. *irr: Various Coefficients of Interrater Reliability and Agreement*. <u>https://www.r-project.org</u>.
- 1764 Gelman, Andrew, and Eric Loken. 2013. "The Garden of Forking Paths: Why Multiple Comparisons
- 1765 Can Be a Problem, Even When There Is No 'Fishing Expedition' or 'p-Hacking' and the Research
- 1766 Hypothesis Was Posited Ahead of Time." *Department of Statistics, Columbia University*.
- Gelman, Andrew, and David Weakliem. 2009. "Of Beauty, Sex, and Power." *American Scientist* 97:310–16.
- 1769 Gould, Elliot, Hannah S. Fraser, Shinichi Nakagawa, and Timothy H. Parker. 2023. "ManyEcoEvo:
- 1770 Meta-Analyse Data from ManyAnalyst Style
- 1771 Studies." Zenodo. https://doi.org/10.5281/zenodo.10046153.
- 1772 Gould, Ellliot, Hannah S. Fraser, Shinichi Nakagawa, Timothy H. Parker. 2024. egouldo/ManyAnalysts:
- 1773 Manuscript Source Code for 'Same data, different analysts: variation in effect sizes due to analytical
- decisions in ecology and evolutionary biology.' Zenodo. <u>https://doi.org/10.5281/zenodo.13850927</u>.
- 1775 Version 2.0.2.
- 1776 Grueber, C. E., S. Nakagawa, R. J. Laws, and I. G. Jamieson. 2011. "Multimodel Inference in Ecology
- 1777 and Evolution: Challenges and Solutions." Journal of Evolutionary Biology 24 (4): 699–
- 1778 711. <u>https://doi.org/doi:10.1111/j.1420-9101.2010.02210.x</u>.
- 1779 Harrell Jr, Frank E. 2024. *Hmisc: Harrell Miscellaneous*. <u>https://hbiostat.org/R/Hmisc/</u>.
- 1780 Hester, Jim, Lionel Henry, Kirill Müller, Kevin Ushey, Hadley Wickham, and Winston Chang.
- 1781 2024. *withr: Run Code "With" Temporarily Modified Global State*. <u>https://withr.r-lib.org</u>.
- 1782 Higgins, Julian P T, Simon G Thompson, Jonathan J Deeks, and Douglas G Altman. 2003. "Measuring
- 1783 Inconsistency in Meta-Analyses." BMJ 327 (7414): 557–
- 1784 60. <u>https://doi.org/10.1136/bmj.327.7414.557</u>.
- 1785 Huntington-Klein, Nick, Andreu Arenas, Emily Beam, Marco Bertoni, Jeffrey R. Bloem, Pralhad Burli,
- 1786 Naibin Chen, et al. 2021. "The Influence of Hidden Researcher Decisions in Applied
- 1787 Microeconomics." *Economic Inquiry* 59 (3): 944–60.
- 1788 https://doi.org/<u>https://doi.org/10.1111/ecin.12992</u>.
- 1789 Iannone, Richard, Joe Cheng, Barret Schloerke, Ellis Hughes, Alexandra Lauer, and JooYoung Seo.
- 1790 2024. gt: Easily Create Presentation-Ready Display Tables. <u>https://gt.rstudio.com</u>.
- 1791 Jennions, M. D., C. J. Lortie, M. S. Rosenberg, and H. R. Rothstein. 2013. "Publication and Related
- 1792 Biases." Book Section. In Handbook of Meta-Analysis in Ecology and Evolution, edited by J. Koricheva,
- 1793 J. Gurevitch, and K. Mengersen, 207–36. Princeton, USA: Princeton University Press.

1794 Kassambara, Alboukadel. 2023. ggpubr: "ggplot2" Based Publication Ready

1795 Plots. https://rpkgs.datanovia.com/ggpubr/.

1796 Kimmel, Kaitlin, Meghan L. Avolio, and Paul J. Ferraro. 2023. "Empirical Evidence of Widespread

1797 Exaggeration Bias and Selective Reporting in Ecology." Nature Ecology &

1798 Evolution. https://doi.org/10.1038/s41559-023-02144-3.

1799 Klein, Richard A., Kate A. Ratliff, Michelangelo Vianello, Reginald B. Adams Jr., Štěpán Bahník, Michael

1800 J. Bernstein, Konrad Bocian, et al. 2014. "Investigating Variation in Replicability: A "Many Labs"

1801 Replication Project." Social Psychology 45 (3): 142–52. <u>https://doi.org/10.1027/1864-9335/a000178</u>.

1802 Klein, Richard A., Michelangelo Vianello, Fred Hasselman, Byron G. Adams, Reginald B. Adams, Sinan

1803 Alper, Mark Aveyard, et al. 2018. "Many Labs 2: Investigating Variation in Replicability Across Samples

and Settings." Advances in Methods and Practices in Psychological Science 1 (4): 443–

1805 90. <u>https://doi.org/10.1177/2515245918810225</u>.

1806 Knight, K. 2000. *Mathematical Statistics*. Book. New York: Chapman; Hall.

1807 Koricheva, Julia, and Jessica Gurevitch. 2014. "Uses and Misuses of Meta-Analysis in Plant

1808 Ecology." Journal of Ecology 102 (4): 828–44. https://doi.org/<u>https://doi.org/10.1111/1365-</u>
1809 2745.12224.

- 1810 Kou-Giesbrecht, Sian, and Duncan N. L. Menge. 2021. "Nitrogen-Fixing Trees Increase Soil Nitrous
- 1811 Oxide Emissions: A Meta-Analysis." *Ecology* 102 (8): e03415.
- 1812 https://doi.org/<u>https://doi.org/10.1002/ecy.3415</u>.

1813 Kuhn, Max, and Hannah Frick. 2022. *multilevelmod: Model Wrappers for Multi-Level* 1814 *Models*. https://github.com/tidymodels/multilevelmod.

- 1815 Kuhn, Max, and Hadley Wickham. 2020. *Tidymodels: A Collection of Packages for Modeling and* 1816 *Machine Learning Using Tidyverse Principles*. <u>https://www.tidymodels.org</u>.
- 1817 Kuznetsova, Alexandra, Per B. Brockhoff, and Rune H. B. Christensen. 2017. "ImerTest Package: Tests
- 1818 in Linear Mixed Effects Models." Journal of Statistical Software 82 (13): 1–
- 1819 26. https://doi.org/10.18637/jss.v082.i13.
- 1820 Landau, William Michael. 2021. "The Targets r Package: A Dynamic Make-Like Function-Oriented
- 1821 Pipeline Toolkit for Reproducibility and High-Performance Computing." *Journal of Open Source*1822 Software 6 (57): 2959. https://doi.org/10.21105/joss.02959.
- 1823 Leybourne, Daniel J., Katharine F. Preedy, Tracy A. Valentine, Jorunn I. B. Bos, and Alison J. Karley.

1824 2021. "Drought Has Negative Consequences on Aphid Fitness and Plant Vigor: Insights from a Meta-

- 1825 Analysis." *Ecology and Evolution* 11 (17): 11915–29.
- 1826 https://doi.org/<u>https://doi.org/10.1002/ece3.7957</u>.
- 1827 Lu, Xun, and Halbert White. 2014. "Robustness Checks and Robustness Tests in Applied
- 1828 Economics." Journal of Econometrics 178: 194–206.
- 1829 https://doi.org/<u>https://doi.org/10.1016/j.jeconom.2013.08.016</u>.
- 1830 Lüdecke, Daniel, Mattan S. Ben-Shachar, Indrajeet Patil, and Dominique Makowski. 2020. "Extracting,
- 1831 Computing and Exploring the Parameters of Statistical Models Using R." *Journal of Open Source*
- 1832 Software 5 (53): 2445. <u>https://doi.org/10.21105/joss.02445</u>.

- 1833 Lüdecke, Daniel, Mattan S. Ben-Shachar, Indrajeet Patil, Philip Waggoner, and Dominique Makowski.
- 1834 2021. "performance: An R Package for Assessment, Comparison and Testing of Statistical
- 1835 Models." Journal of Open Source Software 6 (60): 3139. <u>https://doi.org/10.21105/joss.03139</u>.
- 1836 Lüdecke, Daniel, Indrajeet Patil, Mattan S. Ben-Shachar, Brenton M. Wiernik, Philip Waggoner, and
- 1837 Dominique Makowski. 2021. "see: An R Package for Visualizing Statistical Models." *Journal of Open*
- 1838 Source Software 6 (64): 3393. <u>https://doi.org/10.21105/joss.03393</u>.
- Luke, S. G. 2017. "Evaluating Significance in Linear Mixed-Effects Models in r." *Behavior Research Methods* 49 (4): 1494–1502.
- 1841 Makowski, Dominique, Mattan S. Ben-Shachar, Indrajeet Patil, and Daniel Lüdecke. 2020. "Estimation
- 1842 of Model-Based Predictions, Contrasts and
- 1843 Means." CRAN. https://github.com/easystats/modelbased.
- 1844 Masur, Philipp K., and Michael Scharkow. 2020. "specr: Conducting and Visualizing Specification
- 1845 Curve Analyses (Version 1.0.0)." <u>https://CRAN.R-project.org/package=specr</u>.
- 1846 Meschiari, Stefano. 2022. Latex2exp: Use LaTeX Expressions in
- 1847 Plots. <u>https://www.stefanom.io/latex2exp/</u>.
- 1848 Miles, C. 2008. "Testing Market-Based Instruments for Conservation in Northern Victoria." Book
- 1849 Section. In *Biodiversity: Integrating Conservation and Production: Case Studies from Australian*
- 1850 *Farms, Forests and Fisheries*, edited by T. Norton, T. Lefroy, K. Bailey, and G. Unwin, 133–46.
- 1851 Melbourne, Australia: CSIRO Publishing.
- 1852 Millard, Steven P. 2013. *EnvStats: An r Package for Environmental Statistics*. New York:
 1853 Springer. <u>https://www.springer.com</u>.
- Molina, Isabel, and Yolanda Marhuenda. 2015. "sae: An R Package for Small Area Estimation." *The R Journal* 7 (1): 81–98. <u>https://journal.r-project.org/archive/2015/RJ-2015-007/RJ-2015-007.pdf</u>.
- 1856 Morrissey, Michael B., and Graeme D. Ruxton. 2018. "Multiple Regression Is Not Multiple
- 1857 Regressions: The Meaning of Multiple Regression and the Non-Problem of Collinearity." Philosophy,
- 1858 Theory, and Practice in Biology 10 (3). <u>https://doi.org/10.3998/ptpbio.16039257.0010.003</u>.
- 1859 Müller, Kirill. 2020. here: A Simpler Way to Find Your Files. https://here.r-lib.org/.
- 1860 Nakagawa, Shinichi, and Innes C. Cuthill. 2007. "Effect Size, Confidence Interval and Statistical
- 1861 Significance: A Practical Guide for Biologists." *Biological Reviews* 82 (4): 591–
- 1862 605. <u>https://doi.org/10.1111/j.1469-185X.2007.00027.x</u>.
- 1863 Nakagawa, Shinichi, Malgorzata Lagisz, Michael D. Jennions, Julia Koricheva, Daniel W. A. Noble,
- 1864 Timothy H. Parker, Alfredo Sánchez-Tójar, Yefeng Yang, and Rose E. O'Dea. 2022. "Methods for
- 1865 Testing Publication Bias in Ecological and Evolutionary Meta-Analyses." *Methods in Ecology and*
- 1866 Evolution 13 (1): 4–21. https://doi.org/<u>https://doi.org/10.1111/2041-210X.13724</u>.
- 1867 Nakagawa, Shinichi, Malgorzata Lagisz, Rose E. O'Dea, Patrice Pottier, Joanna Rutkowska, Alistair M.
- 1868 Senior, Yefeng Yang, and Daniel W. A. Noble. 2023. "orchaRd 2.0: An r Package for Visualizing Meta-
- 1869 Analyses with Orchard Plots." *EcoEvoRxiv* 12: 4–12.
- 1870 https://doi.org/<u>https://doi.org/10.32942/X2QC7K</u>.
- 1871 Nakagawa, Shinichi, Yefeng Yang, Erin L. Macartney, Rebecca Spake, and Malgorzata Lagisz.
- 1872 2023. "Quantitative Evidence Synthesis: A Practical Guide on Meta-Analysis, Meta-Regression, and

- 1873 Publication Bias Tests for Environmental Sciences." Environmental Evidence 12 (1):
- 1874 8. <u>https://doi.org/10.1186/s13750-023-00301-6</u>.
- Nakagawa, S., D. W. Noble, A. M. Senior, and M. Lagisz. 2017. "Meta-Evaluation of Meta-Analysis: Ten
 Appraisal Questions for Biologists." *BMC Biology* 15 (1): 18. <u>https://doi.org/10.1186/s12915-017-</u>
- 1877 <u>0357-7</u>.
- 1878 Nicolaus, M., S. P. M. Michler, R. Ubels, M. van der Velde, J. Komdeur, C. Both, and J. M. Tinbergen.
- 1879 2009. "Sex-Specific Effects of Altered Competition on Nestling Growth and Survival: An Experimental
- 1880 Manipulation of Brood Size and Sex Ratio." Journal of Animal Ecology 78 (2): 414–
- 1881 26. <u>https://doi.org/10.1111/j.1365-2656.2008.01505.x</u>.
- 1882 Noble, Daniel W. A., Malgorzata Lagisz, Rose E. O'Dea, and Shinichi Nakagawa.
- 1883 2017. "Nonindependence and Sensitivity Analyses in Ecological and Evolutionary Meta-
- 1884 Analyses." *Molecular Ecology* 26 (9): 2410–25. <u>https://doi.org/10.1111/mec.14031</u>.
- 1885 O'Hara, Robert B., and D. Johan Kotze. 2010. "Do Not Log-Transform Count Data." *Methods in* 1886 *Ecology and Evolution* 1 (2): 118–22. https://doi.org/10.1111/j.2041-210x.2010.00021.x.
- 1887 Open Science Collaboration. 2015. "Estimating the Reproducibility of Psychological
- 1888 Science." *Science* 349 (6251): aac4716. <u>https://doi.org/10.1126/science.aac4716</u>.
- 1889 Page, Matthew J., and David Moher. 2017. "Evaluations of the Uptake and Impact of the Preferred
- 1890 Reporting Items for Systematic Reviews and Meta-Analyses (PRISMA) Statement and Extensions: A
- 1891 Scoping Review." Systematic Reviews 6 (1): 263. <u>https://doi.org/10.1186/s13643-017-0663-8</u>.
- 1892 Parker, Timothy H., Wolfgang Forstmeier, Julia Koricheva, Fiona Fidler, Jarrod D. Hadfield, Yung En
- 1893 Chee, Clint D. Kelly, Jessica Gurevitch, and Shinichi Nakagawa. 2016. "Transparency in Ecology and
- 1894 Evolution: Real Problems, Real Solutions." Trends in Ecology & Evolution 31 (9): 711–
- 1895 19. <u>https://doi.org/10.1016/j.tree.2016.07.002</u>.
- Parker, Timothy H., and Yefeng Yang. 2023. "Exaggerated Effects in Ecology." *Nature Ecology & Evolution*. <u>https://doi.org/10.1038/s41559-023-02156-z</u>.
- Pedersen, Thomas Lin. 2024. *patchwork: The Composer of Plots*. <u>https://patchwork.data-</u>
 <u>imaginist.com</u>.
- 1900 Pei, Yifan, Wolfgang Forstmeier, Daiping Wang, Katrin Martin, Joanna Rutkowska, and Bart
- 1901 Kempenaers. 2020. "Proximate Causes of Infertility and Embryo Mortality in Captive Zebra
- 1902 Finches." The American Naturalist 196 (5): 577–96. <u>https://doi.org/10.1086/710956</u>.
- 1903 Qiu, Yixuan. 2024. showtext: Using Fonts More Easily in r
- 1904 Graphs. <u>https://github.com/yixuan/showtext</u>.
- 1905 R Core Team. 2024. *R: A Language and Environment for Statistical Computing*. Vienna, Austria: R
 1906 Foundation for Statistical Computing. <u>https://www.R-project.org/</u>.
- 1907 Rosenberg, M. S. 2013. "Moment and Least-Squares Based Approaches to Metaanalytic
- 1908 Inference." Book Section. In *Handbook of Meta-Analysis in Ecology and Evolution*, edited by J.
- 1909 Koricheva, J. Gurevitch, and K. Mengersen, 108–24. Princeton, USA: Princeton University Press.
- 1910 Royle, N. J., I. R. Hartley, I. P. F. Owens, and G. A. Parker. 1999. "Sibling Competition and the Evolution
- 1911 of Growth Rates in Birds." *Proceedings of the Royal Society B-Biological Sciences* 266 (1422): 923–
- 1912 32. <u>https://doi.org/10.1098/rspb.1999.0725</u>.

- 1913 Scheinin, Ilari, Maria Kalimeri, Vilma Jagerroos, Juuso Parkkinen, Emmi Tikkanen, Peter Würtz, and
- 1914 Antti Kangas. 2020. ggforestplot: Forestplots of Measures of Effects and Their Confidence
- 1915 Intervals. https://github.com/NightingaleHealth/ggforestplot.
- 1916 Schloerke, Barret, Di Cook, Joseph Larmarange, Francois Briatte, Moritz Marbach, Edwin Thoen,
- 1917 Amos Elberg, and Jason Crowley. 2024. GGally: Extension
- 1918 to "ggplot2". https://ggobi.github.io/ggally/.
- 1919 Schweinsberg, M., M. Feldman, N. Staub, O. R. van den Akker, R. C. M. van Aert, Malm van Assen, Y.
- 1920 Liu, et al. 2021. "Same Data, Different Conclusions: Radical Dispersion in Empirical Results When
- 1921 Independent Analysts Operationalize and Test the Same Hypothesis." Organizational Behavior and
- 1922 Human Decision Processes 165: 228–49. <u>https://doi.org/10.1016/j.obhdp.2021.02.003</u>.
- 1923 Senior, Alistair M., Catherine E. Grueber, Tsukushi Kamiya, Malgorzata Lagisz, Katie O'Dwyer, Eduardo
- 1924 S. A. Santos, and Shinichi Nakagawa. 2016. "Heterogeneity in Ecological and Evolutionary Meta-
- 1925 Analyses: Its Magnitude and Implications." Ecology 97 (12): 3293–
- 1926 99. <u>https://doi.org/10.1002/ecy.1591</u>.
- Shavit, A., and Aaron M. Ellison. 2017. *Stepping in the Same River Twice: Replication in Biological Research*. Edited Book. New Haven, Connecticut, USA: Yale University Press.
- 1929 Siegel, Kyle R., Muskanjot Kaur, A. Calvin Grigal, Rebecca A. Metzler, and Gary H. Dickinson.
- 1930 2022. "Meta-Analysis Suggests Negative, but pCO2-Specific, Effects of Ocean Acidification on the
- 1931 Structural and Functional Properties of Crustacean Biomaterials." *Ecology and Evolution* 12 (6):
- 1932 e8922. https://doi.org/<u>https://doi.org/10.1002/ece3.8922</u>.
- 1933 Silberzahn, R., E. L. Uhlmann, D. P. Martin, P. Anselmi, F. Aust, E. Awtrey, Š. Bahník, et al. 2018. "Many
- 1934 Analysts, One Data Set: Making Transparent How Variations in Analytic Choices Affect
- 1935 Results." Advances in Methods and Practices in Psychological Science 1 (3): 337–
- 1936 56. <u>https://doi.org/10.1177/2515245917747646</u>.
- Silge, Julia, and David Robinson. 2016. "tidytext: Text Mining and Analysis Using Tidy Data Principles
 in r." JOSS 1 (3). <u>https://doi.org/10.21105/joss.00037</u>.
- 1939 Simons, Daniel J., Yuichi Shoda, and D. Stephen Lindsay. 2017. "Constraints on Generality (COG): A
- 1940 Proposed Addition to All Empirical Papers." *Perspectives on Psychological*
- 1941 Science. https://doi.org/10.1177/174569161770863.
- 1942 Simonsohn, Uri, Joseph P. Simmons, and Leif D. Nelson. 2015. "Specification Curve: Descriptive and
- 1943 Inferential Statistics on All Reasonable Specifications." Manuscript. SSRN Electronic
- 1944 Journal. <u>https://doi.org/10.2139/ssrn.2694998</u>.
- 1945 ———. 2020. "Specification Curve Analysis." Nature Human Behaviour 4 (11): 1208–
- 1946 14. <u>https://doi.org/10.1038/s41562-020-0912-z</u>.
- 1947 Sjoberg, Daniel D., Karissa Whiting, Michael Curry, Jessica A. Lavery, and Joseph Larmarange.
- 1948 2021. "Reproducible Summary Tables with the Gtsummary Package." *The R Journal* 13: 570–
 1949 80. https://doi.org/10.32614/RJ-2021-053.
- 1950 Slowikowski, Kamil. 2024. ggrepel: Automatically Position Non-Overlapping Text Labels
- 1951 with "ggplot2". https://ggrepel.slowkow.com/.

- 1952 Stanton-Geddes, John, Cintia Gomes de Freitas, and Cristian de Sales Dambros. 2014. "In Defense of
- 1953 p Values: Comment on the Statistical Methods Actually Used by Ecologists." *Ecology* 95 (3): 637–42.
- 1954 https://doi.org/<u>https://doi.org/10.1890/13-1156.1</u>.
- 1955 Steegen, Sara, Francis Tuerlinckx, Andrew Gelman, and Wolf Vanpaemel. 2016. "Increasing
- 1956 Transparency Through a Multiverse Analysis." Perspectives on Psychological Science 11 (5): 702–
- **1957 12**. <u>https://doi.org/10.1177/1745691616658637</u>.
- 1958 Taylor, James W., and Kathryn S. Taylor. 2023. "Combining Probabilistic Forecasts of COVID-19
- 1959 Mortality in the United States." *European Journal of Operational Research* 304 (1): 25–41.
- 1960 https://doi.org/<u>https://doi.org/10.1016/j.ejor.2021.06.044</u>.
- 1961 Tierney, Nicholas, and Dianne Cook. 2023. "Expanding Tidy Data Principles to Facilitate Missing Data
- Exploration, Visualization and Assessment of Imputations." *Journal of Statistical Software* 105 (7): 1–
 31. https://doi.org/10.18637/jss.v105.i07.
- 1964 Touchon, Justin C., and Michael W. McCoy. 2016. "The Mismatch Between Current Statistical Practice
- and Doctoral Training in Ecology." *Ecosphere* 7 (8): e01394.
- 1966 https://doi.org/<u>https://doi.org/10.1002/ecs2.1394</u>.
- 1967 Ushey, Kevin, and Hadley Wickham. 2023. renv: Project
- 1968 Environments. <u>https://rstudio.github.io/renv/</u>.
- 1969 van den Brand, Teun. 2024. *Ggh4x: Hacks for "ggplot2"*. <u>https://github.com/teunbrand/ggh4x</u>.
- 1970 Vander Werf, Eric. 1992. "Lack's Clutch Size Hypothesis: An Examination of the Evidence Using Meta-
- 1971 Analysis." *Ecology* 73 (5): 1699–1705. <u>https://doi.org/10.2307/1940021</u>.
- 1972 Ver Hoef, Jay M. 2012. "Who Invented the Delta Method?" *The American Statistician* 66 (2): 124–
 1973 27. https://doi.org/10.1080/00031305.2012.687494.
- 1974 Verhulst, S., M. J. Holveck, and K. Riebel. 2006. "Long-Term Effects of Manipulated Natal Brood Size
- 1975 on Metabolic Rate in Zebra Finches." Biology Letters 2 (3): 478–
- 1976 80. <u>https://doi.org/10.1098/rsbl.2006.0496</u>.
- 1977 Vesk, P. A., W. K. Morris, W. McCallum, R. Apted, and C. Miles. 2016. "Processes of Woodland
- 1978 Eucalypt Regeneration: Lessons from the Bush Returns Trial." *Proceedings of the Royal Society of*1979 *Victoria* 128: 54–63.
- Viechtbauer, Wolfgang. 2010. "Conducting Meta-Analyses in R with the metafor Package." *Journal of Statistical Software* 36 (3): 1–48. <u>https://doi.org/10.18637/jss.v036.i03</u>.
- Wickham, Hadley, Mara Averick, Jennifer Bryan, Winston Chang, Lucy D'Agostino McGowan, Romain
 François, Garrett Grolemund, et al. 2019. "Welcome to the tidyverse." *Journal of Open Source*
- 1984 Software 4 (43): 1686. https://doi.org/10.21105/joss.01686.
- Wickham, Hadley, Jim Hester, Winston Chang, and Jennifer Bryan. 2022. *devtools: Tools to Make Developing r Packages Easier*. <u>https://devtools.r-lib.org/</u>.
- Wickham, Hadley, Thomas Lin Pedersen, and Dana Seidel. 2023. *scales: Scale Functions for Visualization*. <u>https://scales.r-lib.org</u>.
- 1989 Wilke, Claus O. 2024. *cowplot: Streamlined Plot Theme and Plot Annotations*
- 1990 for "ggplot2". <u>https://wilkelab.org/cowplot/</u>.

- 1991 Xie, Yihui. 2024a. *knitr: A General-Purpose Package for Dynamic Report Generation in* 1992 r. <u>https://yihui.org/knitr/</u>.
- 1993 ———. 2024b. *xfun: Supporting Functions for Packages Maintained by "Yihui* 1994 Xie". <u>https://github.com/yihui/xfun</u>.
- 1995 Yang, Yefeng, Alfredo Sánchez-Tójar, Rose E. O'Dea, Daniel W. A. Noble, Julia Koricheva, Michael D.
- 1996 Jennions, Timothy H. Parker, Malgorzata Lagisz, and Shinichi Nakagawa. 2023. "Publication Bias
- 1997 Impacts on Effect Size, Statistical Power, and Magnitude (Type m) and Sign (Type s) Errors in Ecology
- 1998 and Evolutionary Biology." BMC Biology 21 (1): 71. <u>https://doi.org/10.1186/s12915-022-01485-y</u>.
- 1999 Zeileis, Achim, Jason C. Fisher, Kurt Hornik, Ross Ihaka, Claire D. McWhite, Paul Murrell, Reto Stauffer,
- 2000 and Claus O. Wilke. 2020. "colorspace: A Toolbox for Manipulating and Assessing Colors and
- 2001 Palettes." Journal of Statistical Software 96 (1): 1–49. <u>https://doi.org/10.18637/jss.v096.i01</u>.

Page numbers in this document:

Supplement A (SM A) – Summarising Variation Among Analysis Specifications – page 2

Supplement B (SM B) – Effect Size Analysis – page 16

Supplement C (SM C) – Explaining Variation in Deviation Scores – page 43

Supplement D (SM D) – Correlation Matrices of Case Study Data – page 137

Links to HTML versions:

Supplement A (SM A) – https://egouldo.github.io/ManyAnalysts/supp_mat/SM1_summary.html

Supplement B (SM B) – https://egouldo.github.io/ManyAnalysts/supp_mat/SM2_EffectSizeAnalysis.html

Supplement C (SM C) – https://egouldo.github.io/ManyAnalysts/supp_mat/SM3_ExplainingDeviation.html

Supplement D (SM D) – https://egouldo.github.io/ManyAnalysts/supp_mat/SM4_case_study_datasets.html

A: Summarising Variation Among Analysis Specifications

```
library(tidyverse)
library(targets)
library(withr)
library(here)
library(metafor)
library(ManyEcoEvo)
library(tidyverse)
library(broom)
library(gt)
library(specr)
library(colorspace)
library(ggthemes)
library(ggh4x)
library(showtext)
```

```
set.seed(1234)
source(here::here("utils.R"))
# extrafont::font_install("Lato")
```

A.1 Summary Statistics

A.1.1 Number of analyses of different types

As described in the summary statistics section of the manuscript, 63 teams submitted 131 Z_r model estimates and 43 teams submitted 64 y_i predictions for the blue tit dataset. Similarly, 40 submitted 79 Z_r model estimates and 14 teams submitted 24 y_i predictions for the *Eucalytpus* dataset. The majority of the blue tit analyses specified normal error distributions and were non-Bayesian mixed effects models. Analyses of the *Eucalyptus* dataset rarely specified normal error distributions, likely because the response variable was in the form of counts. Mixed effects models were also common for *Eucalytpus* analyses (Table A.1).

```
Table1 %>%
  rename(subset = subset name) %>%
  rename_with(~ str_remove(., "sum_")) %>%
  group_by(dataset) %>%
  gt::gt(rowname_col = "subset") %>%
  gt::cols_label(dataset = "dataset",
                 subset = "Subset",
                 totalanalyses = "No. Analyses",
                 teams = "No. Teams",
                 linear = "Normal Distribution",
                 mixed = "Mixed Effect") %>%
  gt::sub values(columns = subset, values = c("effects"),
                 replacement = gt::md("$$Z_r$$")) %>%
  gt::sub values(columns = subset, values = c("predictions"),
                 replacement = gt::md("$$y_i$$")) %>%
  gt::sub_values(columns = subset, values = c("all"),
```

Table A.1: Summary of the number of analysis teams, total analyses, models with normal error distributions, mixed effects models, and models developed with Bayesian statistical methods for effect size analyses only (Z_r) and out-of-sample prediction only (y_i) .

	No. Analyses	No. Teams	Normal Distribution	Mixed Effect	Bayesian
blue tit					
Z_r	131	63	124	128	10
y_i	64	43	59	63	10
Eucalyptus					
Z_r	79	40	15	62	5
y_i	24	14	1	16	3

A.1.2 Model composition

The composition of models varied substantially (Table A.2) in regards to the number of fixed and random effects, interaction terms and the number of data points used. For the blue tit dataset, models used up to 19 fixed effects, 12 random effects, and 10 interaction terms and had sample sizes ranging from 76 to 3720. For the *Eucalyptus* dataset models had up to 13 fixed effects, 4 random effects, 5 interaction terms and sample sizes ranging from 18 to 351.

```
Table2 %>%
  rename(SD = sd, subset = subset name) %>%
  group_by(variable) %>%
  pivot_wider(
    names_from = dataset,
    names_sep = ".",
   values from = c(mean, SD, min, max)
  ) %>%
  mutate(variable = case_when(variable == "samplesize" ~ "N",
                              TRUE ~ variable)) %>%
  gt::gt(rowname col = "subset") %>%
  gt::row group order(groups = c("fixed", "random", "interactions", "N")) %>%
  gt::tab_spanner_delim(delim = ".") %>%
  gt::fmt_scientific(columns = "mean.blue tit",
                     rows = `mean.blue tit` < 0.01,</pre>
                     decimals = 2) %>%
    gt::fmt_scientific(columns = "SD.blue tit",
                     rows = `SD.blue tit` < 0.01,
                     decimals = 2) %>%
    gt::fmt_scientific(columns = "mean.eucalyptus",
                     rows = `mean.eucalyptus` < 0.01,</pre>
                     decimals = 2) %>%
    gt::fmt_scientific(columns = "SD.eucalyptus",
                     rows = `SD.eucalyptus` < 0.01,</pre>
                     decimals = 2) %>%
  gt::cols_label_with(fn = Hmisc::capitalize) %>%
  gt::tab style(
    style = gt::cell_text(transform = "capitalize"),
    locations = gt::cells_column_spanners()
  ) %>%
  gt::tab_style(style = gt::cell_text(transform = "capitalize"),locations = c
ells_row_groups()) %>%
gt::tab style(style = gt::cell text(style = "italic"), locations = cells row
groups(groups = "N")) %>%
  gt::cols label with(c(contains("Eucalyptus")),
                      fn = ~ gt::md(paste0("*",.x, "*"))) %>%
  gt::sub_values(columns = subset, values = c("effects"),
                 replacement = gt::md("$$Z_r$$")) %>%
  gt::sub_values(columns = subset, values = c("predictions"),
                 replacement = gt::md("$$y_i$$")) %>%
  gt::sub_values(columns = subset, values = c("all"),
                 replacement = gt::md("All analyses")) %>%
  gt::opt_stylize(style = 6, color = "gray") %>%
 gt::as_raw_html()
```

Table A.2: Mean, standard deviation and range of number of fixed and random variables, interaction terms used in models and analysis sample size (N). Repeated for effect-size analyses only (Z_r) and out-of-sample predictions only (y_i).

	Me	an		5D		<i>l</i> in	I	Мах		
	Blue tit	Eucalyptus	Blue tit	Eucalyptus	Blue tit	Eucalyptus	Blue tit	Eucalyptus		
Fixed										
Z_r	5.20	5.01	2.92	3.83	1	1	19	13		
y_i	4.78	4.67	2.35	3.45	1	1	10	13		
Random										
Z_r	3.53	1.41	2.08	1.09	0	0	10	4		
y_i	4.42	0.96	2.78	0.81	1	0	12	3		
Interactio	ons									
Z_r	0.44	0.16	1.11	0.65	0	0	10	5		
y_i	0.28	0.17	0.63	0.48	0	0	3	2		
N										
Z_r	2611.09	298.43	937.48	106.25	76	18	3720	351		
y_i	2816.71	325.55	773.21	64.17	396	90	3720	350		

A.1.3 Choice of variables

The choice of variables also differed substantially among analyses (Table A.3) and some analysts constructed new variables that transformed or aggregated one or more existing variables. The blue tit dataset had a total of 52 candidate variables. These variables were included in a mean of 20.5 Z_r analyses (range 0- 100). The *Eucalyptus* dataset had a total of 59 candidate variables. The variables in the *Eucalyptus* dataset were included in a mean of 8.92 Z_r analyses (range 0- 55).

```
#table 3 - summary of mean, sd and range for the number of analyses in which
each variable was used
Table3 %>%
    rename(SD = sd, subset = subset_name) %>%
    pivot wider(
```

```
names from = dataset,
    names_sep = ".",
    values_from = c(mean, SD, min, max)
  ) %>%
  ungroup %>%
  gt::gt(rowname_col = "subset") %>%
  gt::tab spanner delim(delim = ".") %>%
  gt::fmt_scientific(columns = "mean.blue tit",
                     rows = `mean.blue tit` < 0.01,</pre>
                     decimals = 2) %>%
    gt::fmt_scientific(columns = "SD.blue tit",
                     rows = `SD.blue tit` < 0.01,</pre>
                     decimals = 2) %>%
    gt::fmt_scientific(columns = "mean.eucalyptus",
                     rows = `mean.eucalyptus` < 0.01,</pre>
                     decimals = 2) %>%
    gt::fmt_scientific(columns = "SD.eucalyptus",
                     rows = `SD.eucalyptus` < 0.01,</pre>
                     decimals = 2) %>%
  gt::fmt number(decimals = 2,drop trailing zeros = T, drop trailing dec mark
= T) %>%
 gt::cols_label_with(fn = Hmisc::capitalize) %>%
  gt::cols_label_with(c(contains("Eucalyptus")), fn = ~ gt::md(paste0("*",.x,
"*"))) %>%
  gt::sub_values(columns = subset, values = c("effects"),
                 replacement = gt::md("$$Z_r$$")) %>%
  gt::sub values(columns = subset, values = c("predictions"),
                 replacement = gt::md("$$y_i$$")) %>%
  gt::sub_values(columns = subset, values = c("all"),
                 replacement = gt::md("All analyses")) %>%
  gt::opt_stylize(style = 6, color = "gray") %>%
  gt::tab_style(
    style = gt::cell text(transform = "capitalize"),
    locations = gt::cells_column_spanners()
  ) %>%
 gt::as raw html()
```

Table A.3: Mean, SE, minimum and maximum number of analyses in which each variable was used, for effect size analyses only (Z_r) , out-of-sample prediction only (y_i) , using the full dataset.

	Mean		SD		Min		Мах	
	Blue tit	Eucalyptus	Blue tit	Eucalyptus	Eu Blue tit	ucalyptus	Blue tit	Eucalyptus
Z_r	20.5	8.92	27	12.28	0	0	100	55
y_i	10.79	2.2	13.87	3.7	0	0	52	17

A.2 Effect Size Specification Analysis

We used a specification curve (Simonsohn, Simmons, and Nelson 2015) to look for relationships between Z_r values and several modeling decisions, including the choice of independent and dependent variable, transformation of the dependent variable, and other features of the models that produced those Z_r values (Figure A.1, Figure A.2). Each effect can be matched to the model features that produced it by following a vertical line down the figure.

A.2.1 Blue tit

We observed few clear trends in the blue tit specification curve (Figure A.1). The clearest trend was for the independent variable 'contrast: reduced broods vs. unmanipulated broods' to produce weak or even positive relationships, but never strongly negative relationships. The biological interpretation of this pattern is that nestlings in reduced broods averaged similar growth to nestlings in unmanipulated broods, and sometimes the nestlings in reduced broods even grew less than the nestlings in unmanipulated broods. Therefore, it may be that competition limits nestling growth primarily when the number of nestlings exceeds the clutch size produced by the parents, and not in unmanipulated broods. The other relatively clear trend was that the strongest negative relationships were never based on the independent variable 'contrast: unmanipulated broods vs. enlarged broods'. These observations demonstrate the potential value of specification curves.

```
coefs_MA_mod <- bind_rows( ManyEcoEvo_viz %>%
    filter(model_name == "MA_mod",
        exclusion_set == "complete",
        expertise_subset == "All"),
    ManyEcoEvo_viz %>%
    filter(model_name == "MA_mod",
        exclusion_set == "complete-rm_outliers",
        expertise_subset == "All")
) %>%
    hoist(tidy_mod_summary) %>%
    select(-starts_with("mod"), -ends_with("plot"), -estimate_type) %>%
    unnest(cols = c(tidy mod summary))
```
```
analytical choices bt <- ManyEcoEvo results$effects analysis[[1]] %>%
  select(study id,
         response_transformation_description, # Don't need constructed, as th
is is accounted for in y
         response_variable_name,
         test variable,
         Bayesian,
         linear model,
         model_subclass,
         sample size,
         starts_with("num"),
         link function reported,
         mixed model) %>%
  mutate(across(starts_with("num"), as.numeric),
         response_transformation_description = case_when(is.na(response_trans
formation_description) ~ "None",
                                                         TRUE ~ response tran
sformation description)) %>%
  rename(y = response variable name, x = test variable, model = linear model)
%>%
  select(study id,x,y,model, model subclass, response transformation descript
ion, link_function_reported, mixed_model, sample_size) %>%
  pivot_longer(-study_id, names_to = "variable_type", values_to = "variable_n
ame",values transform = as.character) %>%
  left join(forest plot new labels) %>%
  mutate(variable name = case when(is.na(user_friendly_name) ~ variable_name,
TRUE ~ user friendly name)) %>%
  select(-user_friendly_name) %>%
  pivot_wider(names_from = variable_type, values_from = variable_name) %>%
  mutate(sample size = as.numeric(sample size))
MA_mean_bt <- ManyEcoEvo_viz$model[[1]] %>%
  broom::tidy(conf.int = TRUE) %>%
  rename(study_id = term)
results_bt <- ManyEcoEvo_viz$model[[1]] %>%
  broom::tidy(conf.int = TRUE, include_studies = TRUE) %>%
  rename(study_id = term) %>%
  semi join(analytical choices bt) %>%
  left join(analytical choices bt)
samp_size_hist_bt <- specr::plot_samplesizes(results_bt %>%
                                               rename(fit nobs = sample size)
) +
  cowplot::theme_half_open() +
  theme(axis.ticks.x = element blank(),
        axis.text.x = element_blank())
```

```
curve bt <- specr::plot curve(results bt) +</pre>
  geom_hline(yintercept = 0,
             linetype = "dashed",
             color = "black") +
  geom_pointrange(mapping = aes(x = 0, y = estimate, ymin = conf.low, ymax =
conf.high ),
                  data = MA mean bt,
                  colour = "black", shape = "diamond") +
  labs(x = "", y = "Standardized Effect Size Zr") +
  cowplot::theme half open() +
  theme(axis.ticks.x = element_blank(),
        axis.text.x = element blank())
specs_bt <- specr::plot_choices(results_bt %>%
                                  rename("Independent\nVariable" = x,
                                          "Dependent\nVariable" = y,
                                         Model = model,
                                          "Model Subclass" = model_subclass,
                                          "Mixed Model" = mixed_model,
                                          "Response\nTransformation\nDescripti
on" = response_transformation_description,
                                          "Link Function" = link_function_repo
rted),
                                choices = c("Independent\nVariable",
                                             "Dependent\nVariable",
                                             "Model",
                                             "Model Subclass",
                                             "Mixed Model",
                                             "Response\nTransformation\nDescri
ption",
                                             "Link Function")) +
  labs(x = "specifications (ranked)") +
 theme(strip.text.x = element blank(),
        strip.text.y = element_text(size = 8, angle = 360, face = "bold"),
        axis.ticks.x = element blank(),
        axis.text.x = element_blank())
cowplot::plot_grid(curve_bt, specs_bt, samp_size_hist bt,
                   ncol = 1,
                   align = "v",
                   rel heights = c(1.5, 2.2, 0.8),
                   axis = "rbl",labels = "AUTO")
```



Figure A.1: **A.** Forest plot for blue tit analyses: standardized effect-sizes (circles) and their 95% confidence intervals are displayed for each analysis included in the meta-analysis model. The meta-analytic mean effect-size is denoted by a black diamond, with error bars also representing the 95% confidence interval. The dashed black line demarcates effect sizes of 0, whereby no effect of the test variable on the response variable is found. Blue points where Zr and its associated confidence intervals are greater than 0 indicate analyses that found a negative effect of sibling number on nestling growth. Gray coloured points have confidence

intervals crossing 0, indicating no relationship between the test and response variable. Red points indicate the analysis found a positive relationship between sibling number and nestling growth. **B.** Analysis specification plot: for each analysis plotted in A, the corresponding combination of analysis decisions is plotted. Each decision and its alternative choices is grouped into its own facet, with the decision point described on the right of the panel, and the option shown on the left. Lines indicate the option chosen used in the corresponding point in plot A. **C.** Sample sizes of each analysis. Note that empty bars indicate analyst did not report sample size and sample size could not be derived by lead team.

A.2.2 Eucalyptus

In the *Eucalyptus* specification curve, there are no strong trends (Figure A.2). It is, perhaps, the case that choosing the dependent variable 'count of seedlings 0-0.5m high' corresponds to more positive results and choosing 'count of all *Eucalytpus* seedlings' might find more negative results. Choosing the independent variable 'sum of all grass types (with or without non-grass graminoids)' might be associated with more results close to zero consistent with the absence of an effect.

```
analytical choices euc <- ManyEcoEvo results$effects analysis[[2]] %>%
  select(study_id,
         response transformation description, # Don't need constructed, as th
is is accounted for in y
         response_variable_name,
         test_variable,
         Bayesian,
         linear_model,
         model_subclass,
         sample_size,
         starts_with("num"),
         transformation,
         mixed model,
         link_function_reported) %>%
  mutate(across(starts_with("num"), as.numeric),
         response_transformation_description = case_when(is.na(response_trans
formation_description) ~ "None",
                                                         TRUE ~ response tran
sformation description),
         response variable name = case when(response variable name == "averag")
e.proportion.of.plots.containing.at.least.one.euc.seedling.of.any.size" ~ "me
an.prop.plots>=1seedling",
                                            TRUE ~ response_variable_name)) %
>%
  rename(y = response_variable_name, x = test_variable, model = linear_model)
%>%
  select(study_id,x,y,model, model_subclass, response_transformation_descript
ion, link_function_reported, mixed_model, sample_size) %>%
  pivot_longer(-study_id, names_to = "variable_type", values_to = "variable_n"
ame",values_transform = as.character) %>%
```

```
left join(forest plot new labels) %>%
  mutate(variable name = case when(is.na(user friendly name) ~ variable name,
TRUE ~ user_friendly_name)) %>%
  select(-user friendly name) %>%
  pivot_wider(names_from = variable_type, values_from = variable_name, values
_fn = list) %>%
  unnest(cols = everything()) %>%
  mutate(sample size = as.numeric(sample size))
MA mean euc <- ManyEcoEvo viz %>%
  filter(model_name == "MA_mod", publishable_subset == "All", dataset == "euc
alyptus", exclusion_set == "complete") %>%
  pluck("model", 1) %>%
  broom::tidy(conf.int = TRUE) %>%
  rename(study id = term)
results euc <- ManyEcoEvo viz %>%
  filter(model_name == "MA_mod", publishable_subset == "All", dataset == "euc
alyptus", exclusion set == "complete") %>%
  pluck("model", 1) %>%
  broom::tidy(conf.int = TRUE, include studies = TRUE) %>%
  rename(study id = term) %>%
  semi join(analytical choices euc) %>%
  left_join(analytical_choices_euc)
samp size hist euc <- specr::plot samplesizes(results euc %>% rename(fit nobs
= sample size)) +
  cowplot::theme_half_open() +
  theme(axis.ticks.x = element blank(), axis.text.x = element blank())
curve_euc <- specr::plot_curve(results_euc) +</pre>
  geom hline(yintercept = 0,
             linetype = "dashed",
             color = "black") +
  geom pointrange(mapping = aes(x = 0, y = estimate, ymin = conf.low, ymax =
conf.high ),
                  data = MA_mean_euc,
                  colour = "black", shape = "diamond") +
  labs(x = "", y = "Standardized Effect Size Zr") +
  cowplot::theme half open() +
  theme(axis.ticks.x = element blank(),
        axis.text.x = element_blank())
specs_euc <- specr::plot_choices(results_euc %>%
                                  rename("Independent\nVariable" = x,
                                         "Dependent\nVariable" = y,
                                         Model = model,
                                         "Model Subclass" = model_subclass,
                                         "Mixed Model" = mixed model,
```

```
"Response\nTransformation\nDescripti
on" = response_transformation_description,
                                         "Link Function" = link_function_repo
rted),
                                choices = c("Independent\nVariable",
                                            "Dependent\nVariable",
                                            "Model",
                                            "Model Subclass",
                                            "Mixed Model",
                                            "Response\nTransformation\nDescri
ption",
                                            "Link Function")) +
  labs(x = "specifications (ranked)") +
  theme(strip.text.x = element_blank(),
        strip.text.y = element_text(size = 8, angle = 360, face = "bold"),
        axis.ticks.x = element_blank(),
        axis.text.x = element_blank())
cowplot::plot_grid(curve_euc, specs_euc, samp_size_hist_euc,
          ncol = 1,
          align = "v",
          rel_heights = c(1.5, 2.2, 0.8),
          axis = "rbl",labels = "AUTO")
```



Figure A.2: **A.** Forest plot for Eucalyptus analyses: standardized effect-sizes (circles) and their 95% confidence intervals are displayed for each analysis included in the meta-analysis model. The meta-analytic mean effect-size is denoted by a black diamond, with error bars also representing the 95% confidence interval. The dashed black line demarcates effect sizes of 0, whereby no effect of the test variable on the response variable is found. Blue points where Z_r and its associated confidence intervals are greater than 0 indicate analyses that found a positive relationship of grass cover on Eucalyptus seedling success. Gray coloured points have

confidence intervals crossing 0, indicating no relationship between the test and response variable. Red points indicate the analysis found a negative relationship between grass cover and Eucalyptus seedling success. **B.** Analysis specification plot: for each analysis plotted in A, the corresponding combination of analysis decisions is plotted. Each decision and its alternative choices is grouped into its own facet, with the decision point described on the right of the panel, and the option shown on the left. Lines indicate the option chosen used in the corresponding point in plot A. **C.** Sample sizes of each analysis. Note that empty bars indicate analyst did not report sample size and sample size could not be derived by lead team.

SM B: Effect Size Analysis

```
library(withr)
library(here)
library(tidyverse)
library(performance)
library(broom.mixed)
library(gt)
library(gtExtras)
library(lme4)
library(MuMIn)
library(ManyEcoEvo)
library(ggrepel)
library(glue)
library(gluedown)
set.seed(1234)
source(here::here("utils.R"))
ManyEcoEvo results <-
  ManyEcoEvo results %>%
  mutate(effects analysis =
           map(effects_analysis,
               rename,
               id_col = study_id)) #%>%
  # mutate_at(c("data",
                "diversity_data",
  #
  #
                "diversity indices",
                "effects_analysis"),
  #
  #
              .funs = ~ map(.x, anonymise_teams,
                         TeamIdentifier lookup))
```

B.1 Meta-analysis

#

B.1.1 Effect Sizes Z_r

Effect of categorical review rating

The figures below (Figure B.1, Figure B.2) hows the fixed effect of categorical review rating on deviation from the meta-analytic mean. There is very little difference in deviation for analyses in any of the review categories. It is worth noting that each analysis features multiple times in these figures corresponding to the multiple reviewers that provided ratings.

```
orchard_publishability <- function(dat){</pre>
  rma mod rating <-</pre>
    metafor::rma.mv(yi = Zr,
                     V = VZr,
                     data = dat,
                     control = list(maxiter = 1000),mods = ~ PublishableAsIs,
                     sparse = TRUE,
                     random = list(~1|response_id, ~1|ReviewerId))
```

```
orchaRd::orchard_plot(rma_mod_rating,
                        mod = "PublishableAsIs",
                        group = "id_col",
                        xlab = "Standardised Correlation Coefficient (Zr)",
                        cb = TRUE, angle = 45)
}
ManyEcoEvo results$effects analysis[[2]] %>%
    filter(Zr > -4) %>%
    unnest(review data) %>%
    select(Zr, VZr, id_col, PublishableAsIs, ReviewerId, response_id) %>%
    mutate(PublishableAsIs = forcats::as factor(PublishableAsIs) %>%
               forcats::fct relevel(c("deeply flawed and unpublishable",
                                      "publishable with major revision",
                                      "publishable with minor revision",
                                      "publishable as is" ))) %>%
    orchard_publishability() +
    theme(text = element text(size = 20),axis.text.y = element text(size = 20)
)) +
    scale x discrete(labels=c("Deeply Flawed\n & Unpublishable", "Publishable
With\n Major Revision", "Publishable With\n Minor Revision", "Publishable\n A
```

```
s Is"))
```



Figure B.1: Orchard plot of meta-analytic model fitted to all Eucalyptus analyses with a fixed effect for categorical peer-review ratings, and random effects for analyst ID and reviewer ID. Black circles denote coefficient mean for each categorical publishability rating. Thick error

bars represent 95% confidence intervals and whiskers indicate 95% prediction intervals. Effect sizes are represented by circles and their size corresponds to the precision of the estimate.

```
With\n Major Revision", "Publishable With\n Minor Revision", "Publishable\n A s Is"))
```



Figure B.2: Orchard plot of meta-analytic model fitted to all blue tit analyses with a fixed effect for categorical peer-review ratings, and random effects for analyst ID and reviewer ID. Black circles denote coefficient mean for each categorical publishability rating. Thick error bars represent 95% confidence intervals and whiskers indicate 95% prediction intervals. Effect sizes are represented by circles and their size corresponds to the precision of the estimate.

Post-hoc analysis: Exploring the effect of removing analyses with poor peer-review ratings on heterogeneity

The forest plots in Figure B.3 compare the distributions of Z_r effects from our full set of analyses with the distributions of Z_r effects from our post-hoc analyses, which removed either analyses that were reviewed at least once as being 'unpublishable', and analyses that were reviewed at least once as being 'unpublishable' or requiring 'major revisions'. Removing these analyses from the blue tit data had little impact on the overall distribution of the results. When 'unpublishable' analyses of the *Eucalyptus* dataset were removed, the extreme outlier 'Brooklyn-2-2-1' was also removed, resulting in a substantial difference to the amount of observed deviation from the meta-analytic mean.

```
plot_forest <- function(data, intercept = TRUE, MA_mean = TRUE){</pre>
  if (MA_mean == FALSE){
    data <- filter(data, Parameter != "overall")</pre>
  }
  p <- ggplot(data, aes(y = estimate,</pre>
                        x = term,
                        ymin = conf.low,
                         ymax = conf.high,
                         shape = point_shape,
                         colour = parameter_type)) +
    geom_pointrange(fatten = 2) +
    ggforestplot::theme forest() +
    theme(axis.line = element_line(linewidth = 0.10,
                                    colour = "black"),
          axis.line.y = element_blank(),
          text = element_text(family = "Helvetica")#,
          # axis.text.y = element_blank()
    ) +
    guides(shape = "none", colour = "none") +
    coord flip() +
    ylab(bquote(Standardised~Effect~Size~Z[r])) +
    xlab(element_blank()) +
    # scale_y_continuous(breaks = c(-4,-3,-2,-1,0,1),
    # minor_breaks = seq(from = -4.5, to = 1.5, by = 0.5)) +
    NatParksPalettes::scale color natparks d("Glacier")
  if(intercept == TRUE){
    p <- p + geom hline(yintercept = 0)</pre>
  }
  if(MA mean == TRUE){
    p <- p + geom_hline(aes(yintercept = meta_analytic_mean),</pre>
                         data = data,
                         colour = "#01353D",
                         linetype = "dashed")
  }
```

```
return(p)
}
publishable_subsets_forest_data <-</pre>
  ManyEcoEvo viz %>%
  filter(model name == "MA mod",
         exclusion_set == "complete",
         expertise_subset == "All") %>%
  select(ends_with("set"), model, -expertise_subset) %>%
  mutate(plot data =
           map(model,
               .f =
                 ~broom::tidy(.x,
                              conf.int = TRUE,
                              include studies = TRUE) %>%
                 mutate(Parameter =
                          forcats::fct_reorder(term, estimate))),
         meta_analytic_mean =
           map_dbl(plot_data,
                   ~ filter(.x,
                            Parameter == "overall") %>%
                     pull(estimate))) %>%
  select(dataset,
         publishable subset,
         plot_data,
         meta analytic mean) %>%
  unnest(cols = c("plot data")) %>%
  mutate(parameter_type =
           case when(
             str detect(Parameter, "overall") ~ "mean",
             TRUE ~ "study")) %>%
  group_by(dataset, publishable_subset) %>%
  dplyr::mutate(point shape =
                  ifelse(stringr::str detect(term, "overall"),
                         "diamond",
                         "circle"))
# publishable subsets forest data <-</pre>
    publishable_subsets_forest_data %>%
#
   rename(id col = term) %>%
#
    group_by(type) %>%
#
    group_split() %>%
#
    set_names(., publishable_subsets_forest_data$type %>% unique) %>%
#
  # map if(.x = ., names(.) == "study",
#
#
   #
             .f = ~ anonymise_teams(.x, TeamIdentifier_lookup)) %>%
#
  bind rows() %>%
#
   rename(term = id_col)
```

library(tidytext)

```
tidy_overall_labeller <- . %>%
  str_split("_") %>%
  flatten_chr() %>%
  pluck(1)
tidy_forest_labels <- Vectorize(tidy_overall_labeller)</pre>
publishable_subsets_forest_data %>%
  group by(dataset, publishable subset) %>%
  mutate(term = case when(term == "overall" ~
                            paste(term,
                                   dataset,
                                   publishable_subset,
                                  sep = "_"),
                          TRUE ~ term),
         dataset = case_when(dataset == "blue tit" ~ "Blue tit",
                              dataset == "eucalyptus" ~ "Eucalyptus",
                              TRUE ~ NA)) %>%
  arrange(across(.cols = c(type, estimate)),
          .by_group = TRUE) %>%
  rowid_to_column() %>%
  mutate(term = reorder(term, rowid),
         publishable subset =
           case when(publishable subset == "All" ~
                       "All analyses",
                     publishable subset == "data flawed" ~
                       "'Unpublishable'\nremoved",
                     publishable_subset == "data_flawed_major" ~
                       "'Unpublishable' &\n'Major Revisions'\nremoved",
                     TRUE ~ "")) %>%
  plot_forest() +
  scale_x_reordered(labels = tidy_forest_labels) +
  ggh4x::facet nested(dataset ~ publishable subset,
                      independent = "y",
                      scales = "free")
```



Figure B.3: Forest plots of meta-analytic estimated standardized effect sizes (Z_r , blue circles) and their 95% confidence intervals for each effect size included in the meta-analysis model. The meta-analytic mean effect size is denoted by a black triangle and a dashed vertical line,

with error bars also representing the 95% confidence interval. The solid black vertical line demarcates effect size of 0, indicating no relationship between the test variable and the response variable. The left side of each panel shows the analysis team names (anonymous arbitrary names assigned by us), each followed by three numbers. The first number is the submission ID (some analyst teams submitted results to us on >1 submission form), the second number is the analysis ID (some analyst teams included results of >1 analysis in a given submission), and the third number is the effect ID (some analysts submitted values for >1 effect per analysis). Thus, each row in each forest plot is uniquely identified, but it is possible to determine which effects come from which analyses and which analysis teams. The plots in the top row depict effects from analyses of blue tit data, and the bottom row plots depict effects from analyses of Eucalyptus data. The right-most plots depict all usable effect sizes. The plots on the left exclude effects from analysis sets that received at least one rating of "unpublishable" from peer reviewers, and the plots in the middle exclude effects from analysis sets that received at least one rating of either "unpublishable" or "major revision" from peer reviewers.

Post-hoc analysis: Exploring the effect of excluding estimates in which we had reduced confidence

For each dataset (blue tit, Eucalyptus), we created a second, more conservative version, that excluded effects based on estimates of df that we considered less reliable (Table B.1). We compared the outcomes of analyses of the primary dataset (constituted according to our registered plan) with the outcomes of analyses of the more conservative version of the dataset. We also compared results from analyses of both of these versions of the dataset to versions with our post-hoc removal of outliers described in the main text. Our more conservative exclusions (based on unreliable estimates of df) had minimal impact on the meta-analytic mean for both blue tit and Eucalyptus analyses, regardless of whether outliers were excluded (Table B.1).

```
ManyEcoEvo viz %>%
    dplyr::filter(estimate_type == "Zr",
                  model_name == "MA_mod",
                  collinearity_subset != "collinearity_removed",
                  publishable_subset == "All",
                  expertise subset == "All") %>%
    select(dataset, exclusion set, tidy mod summary) %>%
    unnest(tidy_mod_summary) %>%
    filter(type == "summary") %>%
  select(-term, -type) %>%
  mutate(exclusion set =
           case when(exclusion set == "complete" ~
                       "Primary dataset",
                     exclusion_set == "complete-rm_outliers" ~
                       "Primary dataset, outliers removed",
                     exclusion_set == "partial" ~
                       "Conservative exclusions",
                     TRUE ~ "Conservative exclusions, outliers removed")) %>%
```

```
group_by(exclusion set) %>%
  gt::gt() %>%
  gt::opt_stylize(style = 6, color = "gray") %>%
  gt::fmt(columns = "p.value",
          fns = function(x) gtsummary::style_pvalue(x, prepend_p = FALSE)) %>
%
  gt::fmt number(columns = c(-p.value, -dataset)) %>%
  gt::cols_label(estimate = gt::md("$$\\hat\\mu$$"),
                 std.error = gt::md("$$\text{SE}[\\hat\\mu]$$"),
                 conf.low = gt::md("95\\%CI")) %>%
  gt::cols_merge(columns = starts_with("conf"),
                 pattern = "[{1},{2}]") %>%
  gt::cols_move(columns = conf.low, after = std.error) %>%
   gt::tab_style(
    style = list(gt::cell_text(transform = "capitalize"),
                 gt::cell_text(style = "italic")),
    locations = gt::cells_body(columns = "dataset", rows = dataset == "eucaly
ptus")
  )
```

Table B.1: Estimated meta-analytic mean, standard error, and 95% confidence intervals, from analyses of the primary data set, the more conservative version of the dataset which excluded effects based on less reliable estimates of df, and both of these datasets with outliers removed.

	$\hat{\mu}$	$extSE[\hat{\mu}]$			
dataset	,	L / J	95%CI	statistic	p.value
Primary dataset					
blue tit	-0.35	0.03	[-0.41,-0.29]	-11.02	<0.001
Eucalyptus	-0.09	0.06	[-0.22,0.03]	-1.47	0.14
Conservative exclusions					
blue tit	-0.36	0.03	[-0.42,-0.29]	-10.77	<0.001
Eucalyptus	-0.11	0.07	[-0.24,0.03]	-1.55	0.12
Primary dataset, outliers removed					
blue tit	-0.36	0.03	[-0.42,-0.30]	-11.48	<0.001
Eucalyptus	-0.03	0.01	[-0.06,0.00]	-2.23	0.026
Conservative exclusions, outliers removed					
blue tit	-0.36	0.03	[-0.43,-0.30]	-11.38	<0.001
Eucalyptus	-0.04	0.02	[-0.07,-0.01]	-2.52	0.012

```
plot forest <- function(data, intercept = TRUE, MA mean = TRUE ){</pre>
  if (MA mean == FALSE) {
    data <- filter(data, term != "Overall")</pre>
  }
    p <- ggplot(data, aes(y = term,</pre>
                         x = estimate,
                         ymin = conf.low,
                         ymax = conf.high,
                         # shape = point shape,
                         colour = parameter_type)) +
    geom pointrange() +
    ggforestplot::theme_forest() +
    theme(axis.line = element_line(size = 0.10, colour = "black"),
          axis.line.y = element_blank(),
          text = element_text(family = "Helvetica"),
          axis.text.y = element blank()) +
    guides(shape = "none", colour = "none") +
    coord_flip() +
    labs(y = "Standardised Effect Size, Zr",
         x = element blank()) +
    scale_x_continuous(breaks = c(-4, -3, -2, -1, 0, 1)),
                        minor breaks = seq(from = -4.5, to = 1.5, by = 0.5)) +
    NatParksPalettes::scale_color_natparks_d("Glacier")
    if (intercept == TRUE) {
      p <- p + geom_hline(yintercept = 0)</pre>
    }
    if (MA mean == TRUE) {
      p <- p + geom_hline(aes(yintercept = meta_analytic_mean),</pre>
                           data = data,
                           colour = "#01353D",
                           linetype = "dashed")
    }
  return(p)
}
complete_euc_data <-</pre>
  ManyEcoEvo viz %>%
  filter(exclusion set == "complete",
         estimate_type == "Zr",
         model_name == "MA_mod",
         dataset == "eucalyptus",
         publishable subset == "All",
         expertise subset == "All") %>%
  select(tidy mod summary) %>%
  mutate(plot_data = map(tidy_mod_summary,
                          .f = ~ dplyr::mutate(.x,
                                              point shape =
```

```
ifelse(stringr::str detect(term
, "overall"),
                                                      "diamond",
                                                      "circle"),
                                            Parameter =
                                              forcats::fct_reorder(term,
                                                                    estimate)
%>%
                                              forcats::fct_reorder(.,
                                                                    point_shap
e,
                                                                    .desc = TR
UE))
  ),
  meta_analytic_mean = map_dbl(plot_data,
                               ~ filter(.x, Parameter == "overall") %>%
                                 pull(estimate))) %>%
  select(plot_data, meta_analytic_mean) %>%
  unnest(cols = c("plot_data")) %>%
  mutate(parameter_type = case_when(str_detect(Parameter, "overall") ~ "mean"
ر
                                    TRUE ~ "study"))
  # ManyEcoEvo viz %>%
 # filter(exclusion_set == "complete",
          estimate_type == "Zr",
 #
          model name == "MA mod",
 #
          dataset == "eucalyptus",
  #
          publishable subset == "All",
  #
           expertise_subset == "All") %>%
  #
  # )
min_outlier_euc <- complete_euc_data %>%
  filter(type == "study") %>%
  slice_min(estimate, n = 3) %>%
  pull(term)
sample_size_euc_Zr <- ManyEcoEvo_results %>%
    filter(exclusion set == "complete", dataset == "eucalyptus") %>%
    pluck("data", 1) %>%
    select(id col, sample size) %>%
    rename(term = id col) %>%
    mutate(sample_size = as.numeric(sample_size))
mean_n_euc_Zr <- sample_size_euc_Zr %>%
  drop_na(sample_size) %>%
  pull(sample size) %>%
  mean() %>%
  round(2)
```

```
N_outliers_Zr_euc <- sample_size_euc_Zr %>%
filter(term %in% min_outlier_euc) %>%
arrange(desc(sample_size))
```

Post-hoc analysis: Exploring the effect of including only analyses conducted by analysis teams with at least one member self-rated as "highly proficient" or "expert" in conducting statitistical analyses in their research area

The anonymous Team Identifiers in the reduced subset of "expert" or "highly proficient" analysts are exported internally in the ManyEcoEvo package as ManyEcoEvo:::expert_subset. Analyses from the following teams are retained in the reduced subset: *Bell, Berr, Brim, Bruc, Burr, Byng, Cape, Clar, Clev, Alban, Alpha, Bargo, Berry, Bowen, Bulli, Aldgat, Alding, Anakie, Aramac, August, Bamaga, Barham, Barmah, Batlow, Beltan, Bethan, Beulah, Bindoo, Boonah, Bowral, Bright, Buchan, Burnie, Cairns, Casino, Cattai, Adelong, Angasto, Antwerp, Arltung, Ashford, Babinda, Bargara, Barooga, Barraba, Belmont, Bemboka, Benalla, Bendigo, Berrima, Berwick, Beverle, Bicheno, Biloela, Birchip, Bombala, Bonalbo, Brookto, Bruthen, Buderim, Candelo, Capella, Carcoar, Carnama, Chewton, Anglesea, Ardrossa, Armidale, Atherton, Balaklav, Ballarat, Barellan, Belgrave, Berrigan, Binalong, Binnaway, Blackall, Boggabri, Bridport, Brooklyn, Buckland, Bundeena, Bungonia, Busselto, Calliope, Cardwell, Cassilis, Cessnock, Charlton.*

```
plot_forest <- function(data, intercept = TRUE, MA_mean = TRUE){</pre>
  if (MA mean == FALSE){
    data <- filter(data, Parameter != "overall")</pre>
  }
  p <- ggplot(data, aes(y = estimate,</pre>
                        x = term,
                        ymin = conf.low,
                        ymax = conf.high,
                        shape = parameter_type,
                        colour = parameter_type)) +
    geom_pointrange(fatten = 2) +
    ggforestplot::theme forest() +
    theme(axis.line = element_line(linewidth = 0.10, colour = "black"),
          axis.line.y = element blank(),
          text = element text(family = "Helvetica")#,
          # axis.text.y = element_blank()
    ) +
    guides(shape = guide_legend(title = NULL),
           colour = guide_legend(title = NULL)) +
    coord flip() +
    ylab(bquote(Standardised~Effect~Size~Z[r])) +
    xlab(element_blank()) +
    # scale y_continuous(breaks = c(-4,-3,-2,-1,0,1),
    # minor breaks = seq(from = -4.5, to = 1.5, by = 0.5)) +
    NatParksPalettes::scale_color_natparks_d("Glacier")
```

```
if(intercept == TRUE){
    p <- p + geom_hline(yintercept = 0)</pre>
  }
  if(MA_mean == TRUE){
    p <- p + geom hline(aes(yintercept = meta analytic mean),</pre>
                         data = data,
                         colour = "#01353D",
                         linetype = "dashed")
  }
  return(p)
}
filter_experts <-</pre>
  rlang::exprs(
    exclusion_set == "complete",
    estimate type == "Zr",
    model_name == "MA_mod",
    publishable_subset == "All",
    expertise_subset == "expert")
bt experts only <-
  ManyEcoEvo viz %>%
  filter(!!!filter_experts,
         dataset == "blue tit") %>%
  select(tidy mod summary) %>%
  mutate(plot_data = map(tidy_mod_summary,
                          .f = ~ dplyr::mutate(.x,
                                              point shape =
                                                ifelse(stringr::str_detect(term
, "overall"),
                                                       "diamond",
                                                       "circle"),
                                              Parameter =
                                                forcats::fct_reorder(term,
                                                                      estimate)
%>%
                                                forcats::fct reorder(.,
                                                                      point shap
e,
                                                                      .desc = TR
UE))
  ),
  meta_analytic_mean = map_dbl(plot_data,
                                ~ filter(.x, Parameter == "overall") %>%
                                  pull(estimate))) %>%
  select(plot_data, meta_analytic_mean) %>%
  unnest(cols = c("plot_data")) %>%
```

```
mutate(parameter type = case when(str detect(Parameter, "overall") ~ "mean"
ر
                                    TRUE ~ "study"))
# bt experts only <-</pre>
    bt experts only %>%
#
   rename(id col = term) %>%
#
  group_by(type) %>%
#
#
   group_split() %>%
    set_names(., bt_experts_only$type %>% unique) %>%
#
#
   # map_if(.x = ., names(.) == "study",
#
          # .f = ~ anonymise teams(.x, TeamIdentifier lookup)) %>%
#
   bind rows() %>%
#
    rename(term = id_col)
bt_forest_experts <- bt_experts_only %>%
  arrange(desc(type)) %>%
  mutate(type = forcats::as_factor(type)) %>%
  group by(type) %>%
  arrange(desc(estimate),.by group = TRUE) %>%
  mutate(term = forcats::as factor(term),
         point shape = case when(str detect(type, "summary") ~ "mean",
                                 TRUE ~ "study")) %>%
  plot_forest(intercept = TRUE, MA_mean = TRUE) +
  theme(axis.text.x = element_text(size = 15),
        axis.title.x = element text(size = 15),
        axis.text.y = element blank()
  ) +
  scale y continuous(limits = c(-1.6, 0.65))
euc_experts_only <-</pre>
  ManyEcoEvo_viz %>%
  filter(!!!filter experts,
         dataset == "eucalyptus") %>%
  select(model) %>%
  mutate(plot data = map(model,
                         .f = \sim broom::tidy(.x,
                                             conf.int = TRUE,
                                             include studies = TRUE) %>%
                           dplyr::mutate(point_shape =
                                            ifelse(stringr::str_detect(term, "
overall"),
                                                   "diamond",
                                                   "circle"),
                                          Parameter =
                                            forcats::fct_reorder(term,
                                                                 estimate) %>%
                                            forcats::fct_reorder(.,
                                                                 point_shape,
```

```
.desc = TRUE)
)
  ),
  meta analytic mean = map dbl(plot data,
                               ~ filter(.x, Parameter == "overall") %>%
                                 pull(estimate))) %>%
  select(plot data, meta analytic mean) %>%
  unnest(cols = c("plot_data")) %>%
  mutate(parameter_type = case_when(str_detect(Parameter, "overall") ~ "mean"
                                    TRUE ~ "study"))
# euc_experts_only <-</pre>
  euc experts only %>%
#
# rename(id col = term) %>%
#
   group by(type) %>%
  group split() %>%
#
   set_names(., euc_experts_only$type %>% unique) %>%
#
#
   # map_if(.x = ., names(.) == "study",
          # .f = ~ anonymise_teams(.x, TeamIdentifier_lookup)) %>%
#
#
   bind rows() %>%
    rename(term = id col)
#
euc_forest_experts <- euc_experts_only %>%
  arrange(desc(type)) %>%
  mutate(type = forcats::as factor(type)) %>%
  group_by(type) %>%
  arrange(desc(estimate),.by_group = TRUE) %>%
  mutate(term = forcats::as_factor(term),
         point_shape = case_when(str_detect(type, "summary") ~ "mean",
                                 TRUE ~ "study")) %>%
  plot_forest(intercept = TRUE, MA_mean = TRUE) +
  theme(axis.text.x = element_text(size = 15),
        axis.title.x = element text(size = 15),
        axis.text.y = element_blank()
  ) +
  scale y continuous(limits = c(-5, 1),
                     breaks = c(-5, -4, -3, -2, -1, 0, 1))
# ---- Extract Viz ----
bt_forest_experts
euc_forest_experts
```



(a) Blue tit dataset analyses



⁽b) Eucalyptus dataset analyses

Figure B.4: Estimated meta-analytic mean effect size (Z_r) , standard error, and 95% confidence intervals, from analyses of the primary data set with at least one member self-rated as "highly proficient" or "expert" in conducting statistical analyses in their research area.

Post-hoc analysis: Exploring the effect of excluding analyses of the blue tit dataset containing highly collinear predictor variables

For the blue tit dataset, we created a subset of analyses that excluded effects based on analyses containing highly correlated predictor variables. Excluded analyses are exported internally in the ManyEcoEvo package as ManyEcoEvo::collinearity_subset. Analyses with the following identifiers are excluded in the reduced subset: *Armadal-1-1-1, Babinda-1-1-1, Babinda-2-2-1, Barham-1-1-1, Barham-2-2-1, Bega-1-1-1, Bega-1-1-2, Bega-2-2-1, Bega-2-2-2, Borde-1-1-1, Caigun-1-1-1, Caigun-2-2-1, Adelong-1-1-1, Adelong-2-2-1.*

```
# summary_output_params <- rlang::exprs(tidy_mod_summary, MA_fit_stats, mod_f</pre>
it stats)
ManyEcoEvo viz %>%
 filter(!!!filter_collinear) %>%
  mutate(plot_data = map(tidy_mod_summary,
                         .f = ~ dplyr::mutate(.x,
                                             point shape =
                                               ifelse(stringr::str_detect(term
, "overall"),
                                                      "diamond",
                                                      "circle"),
                                             Parameter =
                                               forcats::fct_reorder(term,
                                                                    estimate)
%>%
                                               forcats::fct_reorder(.,
                                                                    point shap
e,
                                                                    .desc = TR
UE))
  ),
  meta_analytic_mean = map_dbl(plot_data,
                               ~ filter(.x, Parameter == "overall") %>%
                                 pull(estimate))) %>%
  select(plot data, meta analytic mean) %>%
  unnest(cols = c("plot data")) %>%
  mutate(parameter_type = case_when(str_detect(Parameter, "overall") ~ "mean"
ر
                                    TRUE ~ "study")) %>%
  arrange(desc(type)) %>%
  mutate(type = forcats::as_factor(type)) %>%
  group_by(type) %>%
  arrange(desc(estimate),.by_group = TRUE) %>%
  mutate(term = forcats::as factor(term),
         point shape = case when(str detect(type, "summary") ~ "mean",
                                 TRUE ~ "study")) %>%
  plot_forest(intercept = TRUE, MA_mean = TRUE) +
  theme(axis.text.x = element text(size = 15),
        axis.title.x = element_text(size = 15),
        axis.text.y = element_blank()
  ) +
    scale_y_continuous(limits = c(-1.5, 0.5),
                     breaks = c(-1.5, -1, -0.5, 0, 0.5))
```



Figure B.5: Forest plot of meta-analytic estimated effect-sizes Z_r , standard error and 95% confidence intervals of blue tit analyses with highly collinear analyses removed. The meta-analytic mean for the reduced subset is denoted by the black triangle, and a dashed vertical line, with error bars representing the 95% confidence interval. The solid black vertical line demarcates effect size of 0.

B.1.2 Out of sample predictions y_i

```
Excluded analyses with constructed variables
by <- join_by(response_variable_name) # don't join on id_col: inc. other excl</pre>
# Analyst Constructed Variables
all_constructed_vars <-
  ManyEcoEvo %>%
    pull(data, dataset) %>%
    list rbind(names to = "dataset") %>%
    filter(str_detect(response_variable_type, "constructed")) %>%
    distinct(dataset,response variable name) %>%
    drop_na() %>%
    arrange()
# Constructed Variables Included in the ManyAnalysts meta-analysis
# (i.e. we have included them in the parameter tables)
ManyEcoEvo_yi_constructed_vars <-</pre>
 ManyEcoEvo:::analysis data param tables %>%
  distinct(variable, dataset) %>%
  rename(response_variable_name = variable) %>%
  semi join(all constructed vars, by) %>%
  filter(!str_detect(response_variable_name,
                     "average.proportion.of")) # was excluded
yi_constructed <-</pre>
  ManyEcoEvo_yi_results %>%
    pull(data, dataset) %>%
    list rbind(names to = "dataset") %>%
    filter(str_detect(response_variable_type, "constructed")) %>%
    distinct(dataset, id_col, TeamIdentifier, response_variable_name) %>%
    drop_na()
excluded_yi_constructed <-</pre>
  ManyEcoEvo %>%
  pull(data, dataset) %>%
  list_rbind(names_to = "dataset") %>%
  filter(str detect(response variable type, "constructed"),
         str_detect(exclusions_all, "retain")) %>%
  distinct(dataset, id col, TeamIdentifier, response variable name) %>%
  drop_na() %>%
  anti join(yi constructed, by) #rm response vars in yi constructed
n_dropped_analyses <-</pre>
  excluded yi constructed %>%
  n_distinct("id_col")
n teams w dropped analyses <-
```

```
excluded_yi_constructed %>%
group_by(TeamIdentifier) %>%
count() %>%
n_distinct("TeamIdentifier")
```

We standardized the y_i estimates and their standard errors for the blue tit analyses using the population mean and standard deviations of the corresponding dependent variable for that analysis, as shown in Equation B.1, using the function ManyEcoEvo::Z_VZ_preds(). Note that this is NOT the same process as standardizing the effect sizes Z_r . We used the mean and standard deviation of the relevant raw datasets as our 'population' parameters.

Equation B1

$$Z_j = \frac{\mu_i - \bar{x}_j}{SD_j}$$
$$VAR_j = \frac{SE_{\mu_i}}{SD_j}$$

Where μ is the population parameter taken from our original dataset for variable *i*, and \bar{x}_i and SD_i are the out of sample point estimate values supplied for analysis j. SE_{μ_i} is the standard error of the population mean for variable i, while VAR_{Z_i} and Z_j are the standardized variance and mean estimate for analysis *j*. Note that for the response variables that were scaled-andcentered, or else mean-centred before model fitting, we do not need to standardise because these are already on the Z-scale. In doing so we make the assumption that analysts' data subsetting will have little effect on the outcomes. For some analyses of the blue tit dataset, analysts constructed their own unique response variables, which meant we needed to reconstruct these variables in order to calculate the population parameters. Unfortunately we were not able to re-construct all variables used by the analysts, as we were unable to reproduce the data required for their re-construction, e.g. we were unable to reproduce principal component analyses or fitted models for extracting residuals Table B.2. A total of 15 were excluded from out-of-sample meta-analysis, from 10 teams, including the following analysis identifiers: Bruc-1-1-1, Clar-2-2-1, Clar-1-1-1, Batlow-1-1-1, Batlow-1-1-2, Bindoo-1-1-1, Bourke-1-1-1, Buchan-1-1-1, Arltung-4-4-1, Bargara-1-1-1, Bendigo-5-5-1, Booligal-3-3-1, Booligal-2-2-1, Booligal-4-4-1 and Booligal-5-5-1.

```
all_constructed_vars %>%
semi_join(ManyEcoEvo_yi_constructed_vars, by) %>%
mutate(included_in_yi = TRUE) %>%
bind_rows(
    {
        all_constructed_vars %>%
        anti_join(ManyEcoEvo_yi_constructed_vars, by) %>%
        mutate(included_in_yi = FALSE)
    }
) %>%
dplyr::mutate(included_in_yi =
        case match(included in yi,
```

```
TRUE ~ "check",
                                            FALSE ~ "xmark" ),
                response_variable_name =
                  gluedown::md_code(response_variable_name)) %>%
  group_by(dataset) %>%
  gt::gt() %>%
  gt::cols_label(response_variable_name = "Constructed Variable",
                 included in yi = gt::md("Variable reconstructed for meta-ana
lysis?")) %>%
  gt::fmt icon(included in yi) %>%
  gt::tab_style(style = cell_text(style = "italic", transform = "capitalize")
ر
                locations = cells_row_groups(groups = "eucalyptus")) %>%
  gt::tab_style(style = cell_text(align = "center"),
                locations = cells_body(columns = included_in_yi)) %>%
  gt::tab style(style = cell text(align = "left"),
                locations = cells_body(columns = response_variable_name)) %>%
  gt::tab style(style = cell text(align = "left"),
                locations = cells column labels(response variable name)) %>%
  gt::tab_style(locations = cells_body(columns = response_variable_name),
                     style = cell_text(size = "small")) %>%
  gt::fmt_markdown(columns = response_variable_name) %>%
  gt::opt_stylize(style = 6, color = "gray", add_row_striping = TRUE) %>%
  gt::opt row striping(row striping = TRUE)
```

Table B.2: Analyst-constructed variables and their inclusion in meta-analyses of out-of-sample predictions, y_i .

Constructed Variable	Variable reconstructed for meta-analysis?			
blue tit				
day_14_weight/day_14_tarsus_length	~			
<pre>day_14_weight/(day_14_tarsus_length^2)</pre>	\checkmark			
SMI	×			
<pre>day_14_tarsus_length_group_deviation</pre>	×			
<pre>day_14_weight_group_deviation</pre>	×			
PC1.day_14_weight.day_14_tarsus_length	×			
<pre>day_14_tarsus_length_deviation</pre>	×			
<pre>residual_day14_weight</pre>	×			
<pre>residual_day_14_weight_males</pre>	×			
Eucalyptus				
euc_sdlgs0_2m	~			
euc_sdlgs_all	\checkmark			
euc_sdlgs>50cm	\checkmark			
<pre>small*0.25+medium*1.25+large*2.5</pre>	\checkmark			
average.proportion.of.plots.containing.at.least.one.euc.seedling.of.any.size	×			

Non-truncated y_i meta-analysis forest plot

Below is the non-truncated version of Figure 4.3 showing a forest plot of the out-of-sample predictions, y_i , on the response-scale (stem counts), for *Eucalyptus* analyses, showing the full error bars of all model estimates.

```
plot_forest_2 <- function(data, intercept = TRUE, MA_mean = TRUE, y_zoom = nu
meric(2L)){
    if(MA_mean == FALSE){
        data <- filter(data, study_id != "overall")
    }
    plot_data <- data %>%
        group_by(study_id) %>%
        group_nest() %>%
```

```
hoist(data, "estimate",.remove = FALSE) %>%
  hoist(estimate, y50 = 2) %>%
  select(-estimate) %>%
  unnest(data) %>%
  arrange(desc(type)) %>%
  mutate(type = forcats::as_factor(type)) %>%
  group by(type) %>%
  arrange(desc(y50),.by_group = TRUE) %>%
  mutate(study_id = forcats::as_factor(study_id),
         point_shape = case_when(str_detect(type, "summary") ~ "diamond",
                                 TRUE ~ "circle"))
p <- ggplot(plot_data, aes(y = estimate,</pre>
                      x = study_id,
                      ymin = conf.low,
                      ymax = conf.high,
                      # shape = type,
                      shape = point shape,
                      colour = estimate_type
                      )) +
  geom_pointrange(position = position_dodge(width = 0.5)) +
  ggforestplot::theme forest() +
  theme(axis.line = element_line(linewidth = 0.10, colour = "black"),
        axis.line.y = element blank(),
        text = element_text(family = "Helvetica")) +
  guides(shape = "none", colour = "none") +
  coord flip(ylim = y zoom) +
  labs(y = "Model estimated out of sample predictions, stem counts",
       x = element blank()) +
  scale_y_continuous(breaks = scales::breaks_extended(10)) +
  NatParksPalettes::scale_color_natparks_d("Glacier")
if(intercept == TRUE){
  p <- p + geom_hline(yintercept = 0)</pre>
}
if(MA_mean == TRUE){
  p <- p +
    geom_hline(aes(yintercept = plot_data %>%
                     filter(type == "summary", estimate_type == "y25") %>%
                     pluck("estimate")),
               data = data,
               colour = "#01353D",
               linetype = "dashed") +
    geom hline(aes(yintercept = plot data %>%
                     filter(type == "summary", estimate type == "y50") %>%
                     pluck("estimate")),
               data = data,
               colour = "#088096",
```

```
linetype = "dashed") +
      geom hline(aes(yintercept = plot data %>%
                       filter(type == "summary", estimate_type == "y75") %>%
                       pluck("estimate")),
                 data = data,
                 colour = "#58B3C7"
                 linetype = "dashed")
  }
  print(p)
}
# ---- new code ----
eucalyptus_yi_plot_data <-</pre>
  ManyEcoEvo yi viz %>%
  filter(dataset == "eucalyptus", model_name == "MA_mod") %>%
  unnest(cols = tidy mod summary) %>%
  mutate(response scale = list(log back(estimate, std.error, 1000)),
         .by = c(dataset, estimate_type, term, type),
         .keep = "used") %>%
  select(-estimate, -std.error) %>%
  unnest wider(response scale) %>%
  rename(estimate = mean origin, conf.low = lower, conf.high = upper) %>%
  nest(tidy mod summary = c(-dataset, -estimate type)) %>% #extract euc data
for plotting (on count scale, not log scale)
  select(dataset, estimate type, tidy mod summary) %>%
  unnest(cols = tidy_mod_summary) %>%
  rename(study_id = term) %>%
  ungroup()
max x axis <-</pre>
  eucalyptus yi plot data %>%
  pluck("conf.high", max) %>%
  round() + 10
eucalyptus yi plot data %>%
  plot_forest_2(MA_mean = T, y_zoom = c(0, max_x_axis)) +
theme(axis.text.y = element_blank())
```



Figure B.6: Forest plot of meta-analytic estimated out of sample predictions, y_i , on the response-scale (stem counts) for Eucalyptus analyses. Circles represent individual analysis estimates. Triangles represent the meta-analytic mean for each prediction scenario. Navy blue coloured points correspond to y_{25} scenario, blue coloured points correspond to the y_{50} scenario, while light blue points correspond to the y_{75} scenario. Error bars are 95% confidence intervals. Outliers (i.e. observations with mean estimates more than 3SD above the population parameter mean, see Section B.1.2.1) have been removed prior to model fitting.

SM C: Explaining Variation in Deviation Scores

```
library(withr)
library(here)
library(tidyverse)
library(performance)
library(broom.mixed)
library(gt)
library(lme4)
library(parameters) #must be loaded directly else parameters fail
library(MuMIn)
library(ManyEcoEvo)
library(tidymodels)
library(multilevelmod)
library(rlang)
set.seed(1234)
source(here::here("utils.R"))
plot_forest <- function(data, intercept = TRUE, MA_mean = TRUE){</pre>
  if (MA mean == FALSE) {
    data <- filter(data, study_id != "overall")</pre>
  }
  data <- data %>%
    group_by(study_id) %>%
    group_nest() %>%
    hoist(data, "estimate",.remove = FALSE) %>%
    hoist(estimate, y50 = 2) %>%
    select(-estimate) %>%
    unnest(data) %>%
    arrange(y50) %>%
    mutate(point_shape = case_when(str_detect(type, "summary") ~ "diamond",
                                    TRUE ~ "circle"))
  p <- ggplot(data, aes(y = estimate,</pre>
                        x = reorder(study id, y50),
                        ymin = conf.low,
                        ymax = conf.high,
                        shape = point_shape,
                        colour = estimate_type
  )) +
    geom pointrange(position = position jitter(width = 0.1)) +
    ggforestplot::theme_forest() +
    theme(axis.line = element line(linewidth = 0.10, colour = "black"),
          axis.line.y = element_blank(),
          text = element_text(family = "Helvetica")) +
    guides(shape = "none", colour = "none") +
    coord flip() +
```
```
labs(y = "Standardised Out of Sample Predictions, Z",
         x = element blank()) +
    scale_y_continuous(breaks = seq(from = round(min(data$conf.low)), to = ro
und(max(data$conf.high)), by = 1),
                       minor_breaks = seq(from = -4.5, to = 1.5, by = 0.5)) +
    NatParksPalettes::scale_color_natparks_d("Glacier")
  if (intercept == TRUE) {
    p <-p + geom hline(yintercept = 0)
  }
  if (MA mean == TRUE) {
   # p <- p + geom_hline(aes(yintercept = meta_analytic_mean),</pre>
   #
                          data = data,
   #
                          colour = "#01353D",
                          linetype = "dashed")
   #
  }
  print(p)
}
plot forest 2 <- function(data, intercept = TRUE, MA mean = TRUE){</pre>
  if (MA mean == FALSE) {
    data <- filter(data, study id != "overall")</pre>
  }
  plot_data <- data %>%
    group by(study id) %>%
    group_nest() %>%
    hoist(data, "estimate",.remove = FALSE) %>%
    hoist(estimate, y50 = 2) %>%
    select(-estimate) %>%
    unnest(data) %>%
    arrange(y50)
  p <- ggplot(plot data, aes(y = estimate,</pre>
                             x = reorder(study_id, y50),
                             ymin = conf.low,
                             ymax = conf.high,
                              # shape = type,
                              colour = estimate_type
  )) +
    geom_pointrange(position = position_dodge(width = 0.5)) +
    ggforestplot::theme_forest() +
    theme(axis.line = element line(linewidth = 0.10, colour = "black"),
          axis.line.y = element blank(),
          text = element_text(family = "Helvetica")) +
    guides(shape = "none", colour = "none") +
    coord flip() +
```

```
labs(y = "Model estimated out of sample predictions",
         x = element blank()) +
    scale_y_continuous(breaks = scales::breaks_extended(10)) +
    NatParksPalettes::scale color natparks d("Glacier")
  if (intercept == TRUE) {
    p <-p + geom hline(yintercept = 0)
  }
  if (MA mean == TRUE) {
    p <- p +
      geom_hline(aes(yintercept = plot data %>%
                       filter(type == "summary", estimate_type == "y25") %>%
                       pluck("estimate")),
                 data = data,
                 colour = "#01353D",
                 linetype = "dashed") +
      geom_hline(aes(yintercept = plot_data %>%
                       filter(type == "summary", estimate type == "y50") %>%
                       pluck("estimate")),
                 data = data,
                 colour = "#088096",
                 linetype = "dashed") +
      geom hline(aes(yintercept = plot data %>%
                       filter(type == "summary", estimate type == "y75") %>%
                       pluck("estimate")),
                 data = data,
                 colour = "#58B3C7"
                 linetype = "dashed")
  }
  print(p)
}
create model workflow <- function(outcome, fixed effects, random intercepts){</pre>
  # https://community.rstudio.com/t/programmatically-generate-formulas-for-lm
er/8575
  # ---- Define random effects constructor function ----
  randomify <- function(feats) {</pre>
    paste0("(1|", feats, ")", collapse = " + ")
  }
 # ---- Construct formula ----
  randomify <- function(feats) paste0("(1|", feats, ")", collapse = " + ")</pre>
  fixed <- paste0(fixed_effects, collapse = " + ")</pre>
  random <- randomify(random intercepts)</pre>
  model formula <- as.formula(paste(outcome, "~", fixed, "+", random))</pre>
```

```
# ---- Construct Workflow ----
  model <- linear_reg() %>%
    set_engine("lmer")
  workflow_formula <- workflow() %>%
    add variables(outcomes = all of(outcome),
                  predictors = all_of(c(fixed_effects, random_intercepts)))
%>%
    add_model(model, formula = model_formula) #%>%
  # add case weights(weight)
  return(workflow_formula)
}
# Define Plotting Function
plot_model_means_box_cox_cat <- function(dat,</pre>
                                          variable,
                                          predictor_means,
                                          new_order,
                                          title,
                                          back_transform = FALSE) {
  dat <- mutate(dat,</pre>
                "{{variable}}" := #
                  fct_relevel(.f = {{variable}},
                               new_order)
  )
  if (back_transform == TRUE) {
    dat <- dat %>%
      mutate(box cox abs deviation score estimate =
               sae::bxcx(unique(dat$lambda),
                         x = box_cox_abs_deviation_score_estimate, InverseQ =
TRUE))
    predictor means <- predictor means %>%
      as tibble() %>%
      mutate(lambda = dat$lambda %>% unique()) %>%
      mutate(across(.cols = -PublishableAsIs,
                    ~ sae::bxcx(unique(dat$lambda),x = .x, InverseQ = TRUE)))
  }
  p <- ggplot(dat, aes(x = {{variable}},</pre>
                       y = box cox abs deviation score estimate)) +
    # Add base dat
    geom_violin(aes(fill = {{variable}}),
                trim = TRUE,
                # scale = "count",
                colour = "white") +
```

```
see::geom jitter2(width = 0.05, alpha = 0.5) +
    # Add pointrange and line from means
    geom_line(dat = predictor_means, aes(y = Mean, group = 1), linewidth = 1)
+
    geom_pointrange(
      dat = predictor_means,
      aes(y = Mean, ymin = CI_low, ymax = CI_high),
      linewidth = 1,
      color = "white",
      alpha = 0.5
    ) +
    # Improve colors
    see::scale fill material d(discrete = TRUE,
                                name = "",
                                palette = "ice",
                                labels = pull(dat, {{variable}}) %>%
                                  levels() %>%
                                  capwords(),
                                reverse = TRUE) +
    EnvStats::stat_n_text() +
    see::theme modern() +
    theme(axis.text.x = element_text(angle = 90))
  if (back_transform == TRUE) {
    p <- p +
      labs(x = "Categorical Peer Review Rating",
           y = "Absolute Deviation from\n Meta-Anaytic Mean Zr")
  } else {
    p <- p + labs(x = "Categorical Peer Review Rating",</pre>
                  y = "Deviation from\nMeta-Analytic Mean Effect Size")
  }
  return(p)
}
possibly check convergence <- possibly(performance::check convergence,
                                        otherwise = NA)
possibly_check_singularity <- possibly(performance::check_singularity,</pre>
                                        otherwise = NA)
# define plotting fun for walk plotting
plot continuous rating <- function(plot data){</pre>
  plot_data %>%
    plot cont rating effects(response = "box cox abs_deviation_score_estimate")
",
                              predictor = "RateAnalysis",
                              back_transform = FALSE,
                              plot = FALSE) %>%
```

```
pluck(2) +
    ggpubr::theme pubr() +
    ggplot2::xlab("Rating") +
    ggplot2::ylab("Deviation In Effect Size from Analytic Mean")
}
walk plot effects diversity <- function(model, plot data, back transform = FA
LSE){
  out_plot <- plot_effects_diversity(model, plot_data, back_transform) +</pre>
    ggpubr::theme pubr()
 return(out plot)
}
plot model means RE <- function(data, variable, predictor means) {</pre>
  p <- ggplot(data, aes(x = as.factor({{variable}}),</pre>
                        y = box_cox_abs_deviation_score_estimate)) +
    # Add base data
    geom_violin(aes(fill = as.factor({{variable}})), color = "white") +
    see::geom jitter2(width = 0.05, alpha = 0.5) +
    # Add pointrange and line from means
    geom line(data = predictor means, aes(y = Mean, group = 1), linewidth = 1
) +
    geom_pointrange(
      data = predictor_means,
      aes(y = Mean, ymin = CI low, ymax = CI high),
      linewidth = 1,
     color = "white"
    ) +
    # Improve colors
    scale x discrete(labels = c("0" = "No Random Effects", "1" = "Random Effe
cts")) +
    see::scale_fill_material(palette = "ice",
                              discrete = TRUE,
                              labels = c("No Random Effects", "Random effects"
),
                              name = "") +
    see::theme modern() +
    EnvStats::stat_n_text() +
    labs(x = "Random Effects Included",
         y = "Deviation from meta-analytic mean")+
    guides(fill = guide_legend(nrow = 2)) +
    theme(axis.text.x = element text(angle = 90))
  return(p)
}
poss_fit <- possibly(fit, otherwise = NA, quiet = FALSE)</pre>
```

```
create_model_formulas <- function(outcome, fixed_effects, random_intercepts){
    # https://community.rstudio.com/t/programmatically-generate-formulas-for-lm
er/8575
    # ---- Define random effects constructor function ----
randomify <- function(feats) {
    paste0("(1|", feats, ")", collapse = " + ")
    }
    # ---- Construct formula -----
randomify <- function(feats) paste0("(1|", feats, ")", collapse = " + ")
fixed <- paste0(fixed_effects, collapse = " + ")
random <- randomify(random_intercepts)
model_formula <-- as.formula(paste(outcome, "~", fixed, "+", random))
return(model_formula)
}</pre>
```

C.1 Box-Cox transformation of response variable for model fitting

To aid in interpreting explanatory models where the response variable has been Box-Cox transformed, we plotted the transformation relationship for each of our analysis datasets (Figure C.1). Note that timetk::step_box_cox() directly optimises the estimation of the transformation parameter lambda, λ , using the "Guerrero" method such that λ minimises the coefficient of variation for sub-series of a numeric vector (Dancho and Vaughan 2023, see ?timetk::step_box_cox() for further details). Consequently, each dataset has its own unique value of λ , and therefore a unique transformation relationship.

```
prep math label estimate type <- function(estimate string){</pre>
  paste0(substring(estimate_string, 1, 1),
         "[", substring(estimate string, 2, 3), "]")
}
filter_vars_main_analysis <- rlang::exprs(estimate_type == "Zr",
                                          exclusion set == "complete",
                                          publishable subset == "All",
                                          expertise subset == "All",
                                           collinearity_subset == "All")
transformation plot data <-
  ManyEcoEvo::ManyEcoEvo yi results %>%
  bind rows(ManyEcoEvo results %>%
              filter(!!!filter_vars_main_analysis)) %>%
  select(dataset, estimate type, effects analysis) %>%
  hoist(effects_analysis, "abs_deviation_score estimate",
        "box_cox_abs_deviation_score_estimate") %>%
  hoist(effects_analysis, "lambda", .simplify = TRUE, .transform = ~unique(.x
)) %>%
  select(-effects_analysis) %>%
  unnest(cols = c(abs deviation score estimate,
                  box cox abs deviation score estimate))
```

```
transformation plot data %>%
  mutate(estimate type = forcats::as factor(estimate type),
         estimate_type = forcats::fct_relabel(estimate_type, prep_math_label_
estimate type),
         dataset = case_match(dataset,
                              "eucalyptus" ~ "Eucalyptus",
                              .default = dataset),
         dataset = dplyr::if_else(str_detect(dataset, "blue"),
                                   latex2exp::TeX(dataset, output = "characte")
r"),
                                   latex2exp::TeX(dataset, italic = TRUE, out
put = "character") )
  ) %>%
  ggplot(aes(y = abs_deviation_score_estimate,
             x = box_cox_abs_deviation_score_estimate)) +
  geom point() +
  ggh4x::facet_grid2(c("dataset", "estimate_type"),
                     scales = "free",
                     independent = "all",
                     labeller = labeller(estimate_type = label_parsed, datase
t = label parsed)) +
  geom_label(aes(x = -Inf, y = Inf,
                 label = latex2exp::TeX(paste("$\\lambda =$", round(lambda, d
igits = 4)), output = "character"),
                 hjust = -0.2, vjust = 2),
             size = 4, parse = TRUE) +
  theme bw() +
  xlab("Box-Cox transformed absolute deviation score") +
ylab("Absolute deviation score")
```



Figure C.1: Box-Cox transformed absolute deviation scores plotted against (untransformed) absolute deviation scores.

C.2 Model Convergence and Singularity problems

During model fitting, especially during fitting of models with random effects using lme4:: (Bates et al. 2015), some models failed to converge while others were accompanied with console warnings of singular fit. However, the convergence checks from lme4:: are known to be overly strict (see ?performance::check_convergence() documentation for a discussion of this issue), consequently we checked for model warnings of convergence failure using the performance::check_convergence() function from the performance:: package (Lüdecke, Ben-Shachar, et al. 2021). For all models we double-checked that they did not have singular fit by using performance::check_singularity(). Despite passing singularity checks with the performance:: package, parameters::parameters() was unable to properly estimate SE and confidence intervals for the random effects of some models, which suggests singular fit. For all models we also checked whether the SE of random effects estimates could be calculated, and if not, marked these models as being singular. Analyses of singularity and convergence are presented throughout this document under the relevant section-heading for the analysis type and outcome, i.e. effect size (Z_r) or out-of-sample predictions (y_i).

C.3 Deviation Scores as explained by Reviewer Ratings

C.3.1 Effect Sizes Z_r

Models pf deviation explained by categorical peer ratings all had singular fit or failed to converge for both blue tit and *Eucalyptus* datasets when random effects were included for both the effect ID and the reviewer ID (Table C.1). For the *Eucalyptus* dataset, when a random effect was included for effect ID only, the model failed to converge. The same was true for the blue tit dataset. As for the effect-size analysis, we included a random-effect for Reviewer ID only when fitting models of deviation explained by categorical peer ratings (See Table 4.6).

For models of deviation explained by continuous peer-review ratings, when including both random effects for effect ID and Reviewer ID model fits were singular for both datasets (Table C.1). The models passed the performance::check_singularity() check, however, however, the SD and CI could not be estimated by parameters::model_parameters() with a warning stating this was likely due to singular fit. For models with a random effect for effect ID, the same occurred for the blue tit dataset, whereas for the *Eucalyptus* dataset, the model did not converge at all. Consequently, for both blue tit and *Euclayptus* datasets, we fitted and analysed models of deviation explained by continuous peer review ratings with a random effect for Reviewer ID only (See Table 4.6).

```
library(multilevelmod)
```

```
poss_extract_fit_engine <- purrr::possibly(extract_fit_engine, otherwise = NA</pre>
poss_parameters <- purrr::possibly(parameters::parameters, otherwise = NA)</pre>
model <- linear reg() %>%
  set_engine("lmer", control = lmerControl(optimizer = "nloptwrap"))
base wf <- workflow() %>%
  add_model(model)
formula study id <- workflow() %>%
  add_variables(outcomes = box_cox_abs_deviation_score_estimate,
                predictors = c(publishable_as_is, study_id)) %>%
  add model(model, formula = box cox abs deviation score estimate ~ publishab
le_as_is + (1 | study_id ))
formula_ReviewerId <- workflow() %>%
  add_variables(outcomes = box_cox_abs_deviation_score_estimate,
                predictors = c(publishable_as_is, reviewer_id)) %>%
  add model(model,
            formula = box cox abs deviation score estimate ~ publishable as i
s + (1 | reviewer_id ))
formula_both <- workflow() %>%
```

```
add variables(outcomes = box cox abs deviation score estimate,
                predictors = c(publishable_as_is, reviewer_id, study_id)) %>
%
  add model(model,
            formula = box_cox_abs_deviation_score_estimate ~ publishable_as_i
s + (1 | study id) + (1 | reviewer id))
# ---- Create DF for combinatorial model specification ----
model_vars <-</pre>
  bind rows(
    tidyr::expand grid(outcome = "box cox abs deviation score estimate",
                       fixed_effects = c("publishable_as_is",
                                          "rate analysis"),
                       random_intercepts = c("study_id",
                                              "reviewer_id")) %>%
      rowwise() %>%
      mutate(random_intercepts = as.list(random_intercepts)),
    tidyr::expand grid(outcome = "box cox abs deviation score estimate",
                       fixed_effects = c("publishable_as_is",
                                          "rate analysis"),
                       random_intercepts = c("study_id",
                                              "reviewer id")) %>%
      group_by(outcome, fixed_effects) %>%
      reframe(random intercepts = list(random intercepts))
  )
# ----- Run all models for all combinations of dataset, exclusion_set, and pu
blishable subset ----
# And Extract
set.seed(1234)
all_model_fits <-
  model vars %>%
  cross_join(.,
             {ManyEcoEvo::ManyEcoEvo results %>%
                 select(estimate type, ends with("set"), effects analysis) %>
%
                 dplyr::filter(expertise_subset == "All",
                               collinearity_subset == "All") %>%
                 select(-c(expertise subset, collinearity subset))}) %>%
  ungroup() %>%
  filter(publishable subset == "All",
         exclusion set == "complete") %>%
  select(-c(exclusion_set, publishable_subset)) %>%
  mutate(effects_analysis =
           map(effects_analysis,
               ~ .x %>%
                 unnest(review data) %>%
                 select(any_of(c("id_col", "study_id")),
```

```
starts with("box cox abs dev"),
                        RateAnalysis,
                        PublishableAsIs,
                        ReviewerId,
                        box_cox_var) %>%
                 janitor::clean_names() %>%
                 mutate if(is.character, factor)
           ),
         model_workflows = pmap(.1 = list(outcome,
                                          fixed effects,
                                          random_intercepts),
                                .f = create model workflow),
         fitted mod workflow = map2(model workflows, effects analysis, poss f
it), #NOT MEANT TO BE TEST DAT
         fitted_model = map(fitted_mod_workflow, extract_fit_engine),
         convergence = map lgl(fitted model, performance::check convergence),
         singularity = map_lgl(fitted_model, performance::check_singularity),
         params = map(fitted model, parameters::parameters)
  ) %>%
  unnest_wider(random_intercepts, names_sep = "_") %>%
  select(-outcome,
         -model_workflows,
         -fitted_mod_workflow,
         -effects analysis,
         estimate type) %>%
  replace_na(list(convergence = FALSE))
# If singularity == FALSE and convergence == TRUE, but the model appears here
, then that's because
# the SD and CI's couldn't be estimated by parameters::
Zr_singularity_convergence <-</pre>
  all_model_fits %>%
  left_join({all_model_fits %>%
      unnest(params) %>%
      filter(Effects == "random") %>%
      filter(if any(contains("SE"), list(is.infinite, is.na))) %>%
      distinct(fixed effects,
               random_intercepts_1,
               random_intercepts_2,
               dataset,
               estimate_type,
               convergence,
               singularity) %>%
      mutate(SE_calc = FALSE)}) %>%
  left_join({all_model_fits %>%
      unnest(params) %>%
      filter(Effects == "random") %>%
      filter(if_any(contains("CI"), list(is.infinite, is.na))) %>%
      distinct(fixed_effects,
```

```
random intercepts 1,
              random_intercepts_2,
              dataset,
              estimate_type,
              convergence,
              singularity) %>%
     mutate(CI calc = FALSE)}) %>%
 rowwise() %>%
 mutate(across(ends_with("_calc"),
               ~ replace_na(.x, TRUE))) %>%
 mutate(across(c(SE_calc, CI_calc, singularity), ~ ifelse(is_false(convergen
ce), NA, .x)))
# ----- new code showing ALL model fits not just bad fits
Zr_singularity_convergence %>%
 select(-fitted_model, -params, -estimate_type) %>%
 arrange(dataset,
         fixed effects,
         random intercepts 1,
         random intercepts 2
 ) %>%
 Hmisc::capitalize() %>%
                 str_replace("id", "ID")),
        dataset =
          case when(dataset == "eucalyptus" ~ Hmisc::capitalize(dataset),
                    TRUE ~ dataset)) %>%
 group_by(dataset) %>%
 gt::gt() %>%
 gt::text_transform(
   locations = cells_body(
     columns = fixed effects,
     rows = random_intercepts_1 != "Reviewer ID"
   ),
   fn = function(x)
     paste0("")
   }
 ) %>%
 tab_style(
   style = list(
     cell_fill(color = scales::alpha("red", 0.6)),
     cell_text(color = "white", weight = "bold")
   ),
   locations = list(
     cells_body(columns = "singularity", rows = singularity == TRUE),
     cells_body(columns = "convergence", rows = convergence == FALSE),
     cells_body(columns = "SE_calc", rows = SE_calc == FALSE),
     cells_body(columns = "CI_calc", rows = CI_calc == FALSE)
```

```
)
  ) %>%
  gt::text_transform(fn = function(x) ifelse(x == TRUE, "yes",
                                             ifelse(x == FALSE, "no", x)),
                     locations = cells_body(columns = c("singularity", "conve
rgence", "SE_calc", "CI_calc"))) %>%
  gt::opt_stylize(style = 6, color = "gray") %>%
  gt::cols_label(dataset = "Dataset",
                 fixed_effects = "Fixed Effect",
                 singularity = "Singular Fit?",
                 convergence = "Model converged?",
                 SE calc = gt::md("Can random effects $\\text{SE}$ be calcula
ted?"),
                 CI_calc = "Can random effect 95% CI be calculated?") %>%
  gt::tab_spanner(label = "Random Effects",
                  columns = gt::starts with("random")) %>%
  gt::sub_missing() %>%
  gt::cols_label_with(columns = gt::starts_with("random"),
                      fn = function(x) paste0("")) %>%
  gt::tab_style(locations =
                  cells_body(rows = str_detect(dataset, "Eucalyptus"),
                             columns = dataset),
                style = cell_text(style = "italic")) %>%
  gt::text_transform(fn = function(x) str_replace(x, "publishable_as_is", "Ca
tegorical Peer Rating") %>%
                       str_replace(., "rate_analysis", "Continuous Peer Ratin
g"),
                     locations = cells body(columns = c("fixed effects"))) %>
%
  gt::tab_style(style = cell_text(style = "italic", transform = "capitalize"
),
                locations = cells_row_groups(groups = "Eucalyptus"))
```

Table C.1: Singularity and convergence checking outcomes for models of deviation in effectsizes Z_r explained by peer-review ratings for different random effect structures. Problematic checking outcomes are highlighted in red.

	Randon	n Effects	Model		Can random effects SE be	Can random effect 95% CI be
Fixed Effect			converged?	Singular Fit?	calculated?	calculated?
blue tit						
Categorical Peer Rating	Reviewer ID	_	yes	no	yes	yes
	Study ID	Reviewer ID	yes	no	yes	no
	Study ID	_	no	_	_	_
Continuous Peer Rating	Reviewer ID	_	yes	no	yes	yes
	Study ID	Reviewer ID	yes	no	no	no
	Study ID	_	yes	no	no	no
Eucalyptus						
Categorical Peer Rating	Reviewer ID	_	yes	no	yes	yes
	Study ID	Reviewer ID	yes	yes	no	no
	Study ID	-	no	—	-	—
Continuous Peer Rating	Reviewer ID	_	yes	no	yes	yes
	Study ID	Reviewer ID	yes	no	no	no
	Study ID	_	no	-	_	_

C.3.2 Out of sample predictions y_i

As for effect-size estimates Z_r , we encountered convergence and singularity problems when fitting models of deviation in out-of-sample predictions y_i explained by categorical peer ratings for both datasets (Table C.2). For all continuous models across both datasets, we encountered convergence and singularity problems when including random effects for both effect ID and Reviewer ID, as well as when including random effects for the effect ID only. In the latter case, for many prediction scenarios, across both blue tit and *Eucalyptus* datasets, estimated random effect coefficient CI's and SE could not be estimated. For models of deviation in out-of-sample predictions explained by continuous peer review ratings, when a random effect was included

for effect ID only, CI's returned values of 0 for both bounds and model means estimated with modelbased::estimate_means() could not be reliably estimated and were equal for every peer-rating category (Table C.2). Consequently, we fitted models of deviation in out-of-sample predictions explained by continuous peer ratings with a random effect for Reviewer ID only (Table C.3). These model structures matched converging and non-singular model structures for effect-size estimates Z_r (Table C.1).

```
all_model_fits_yi <-</pre>
  model_vars %>%
  cross_join(.,
             {ManyEcoEvo::ManyEcoEvo_yi_results %>%
                 select(estimate_type, ends_with("set"), effects_analysis)})
%>%
  ungroup() %>%
  mutate(effects analysis =
           map(effects_analysis,
               ~ .x %>%
                 select(any of(c("id col", "study id")),
                        starts_with("box_cox_abs_dev"),
                        RateAnalysis,
                        PublishableAsIs,
                        ReviewerId,
                        box_cox_var) %>%
                 janitor::clean names() %>%
                 mutate_if(is.character, factor)
           ),
         model workflows = pmap(.1 = list(outcome,
                                           fixed_effects,
                                           random intercepts),
                                 .f = create model workflow),
         fitted mod workflow = map2(model workflows, effects analysis, poss f
it), #NOT MEANT TO BE TEST DAT
         fitted_model = map(fitted_mod_workflow, poss_extract_fit_engine),
         convergence = map(fitted_model, possibly_check_convergence),
         singularity = map(fitted model, possibly check singularity),
         params = map(fitted_model, poss_parameters)) %>%
  mutate(
    across(where(is.list),
           .fns = ~ coalesce(.x, list(NA)))
  ) %>%
  mutate(convergence = list c(convergence),
         singularity = list c(singularity)) %>%
  unnest_wider(random_intercepts, names_sep = "_") %>%
  select(-outcome,
         -model_workflows,
         -fitted mod workflow,
         -effects analysis,
         estimate_type)
```

```
yi singularity convergence all <-
 all model fits yi %>%
 left_join({all_model_fits_yi %>%
     unnest(params) %>%
     filter(Effects == "random") %>%
     filter(if_any(contains("SE"), list(is.infinite, is.na))) %>%
     distinct(fixed effects,
              random intercepts 1,
              random_intercepts_2,
              dataset,
              estimate_type,
              convergence,
              singularity) %>%
     mutate(SE_calc = FALSE)}) %>%
 left_join({all_model_fits %>%
     unnest(params) %>%
     filter(Effects == "random") %>%
     filter(if any(contains("CI"), list(is.infinite, is.na))) %>%
     distinct(fixed effects,
              random_intercepts_1,
              random_intercepts_2,
              dataset,
              estimate_type,
              convergence,
              singularity) %>%
     mutate(CI_calc = FALSE)}) %>%
 rowwise() %>%
 mutate(across(ends_with("_calc"),
               ~ replace_na(.x, TRUE))) %>%
 mutate(across(c(SE calc, CI calc, singularity), ~ ifelse(is false(convergen
ce), NA, .x)))
yi singularity convergence all %>%
    select(-fitted model, -params) %>%
 arrange(dataset,
         estimate_type,
         fixed_effects,
         random intercepts 1,
         random intercepts 2
 ) %>%
 Hmisc::capitalize() %>%
                 str_replace("id", "ID")),
        dataset =
          case_when(dataset == "eucalyptus" ~ Hmisc::capitalize(dataset),
                    TRUE ~ dataset)) %>%
 mutate(fixed_effects = forcats::fct_recode(fixed_effects,
                                            "Categorical Peer Rating" = "pub
lishable as is",
```

```
"Continuous Peer Rating" = "rate
analysis")) %>%
  group_by(fixed_effects) %>%
  arrange(fixed_effects, dataset, pick(starts_with("random"))) %>%
  relocate(estimate_type,.after = dataset) %>%
  gt::gt(rowname_col = "dataset") %>%
  gt::text transform(
    locations = cells body(
      columns = fixed effects,
      rows = random intercepts 1 != "Reviewer ID"
    ),
   fn = function(x)
     paste0("")
    }
  ) %>%
  tab style(
    style = list(
      cell fill(color = scales::alpha("red", 0.6)),
      cell text(color = "white", weight = "bold")
    ),
    locations = list(
      cells_body(columns = "singularity", rows = singularity == TRUE),
      cells_body(columns = "convergence", rows = convergence == FALSE),
      cells_body(columns = "SE_calc", rows = SE_calc == FALSE),
      cells_body(columns = "CI_calc", rows = CI_calc == FALSE)
    )
  ) %>%
  gt::text_transform(fn = function(x) ifelse(x == TRUE, "yes",
                                             ifelse(x == FALSE, "no", x)),
                     locations = cells body(columns = c("singularity", "conve
rgence", "SE_calc", "CI_calc"))) %>%
  gt::opt_stylize(style = 6, color = "gray") %>%
  gt::cols_label(dataset = "Dataset",
                 estimate_type = "Prediction Scenario",
                 fixed_effects = "Fixed Effect",
                 singularity = "Singular Fit?",
                 convergence = "Model converged?",
                 SE_calc = gt::md("Can random effects $\\text{SE}$ be calcula
ted?"),
                 CI_calc = "Can random effect 95% CI be calculated?") %>%
  gt::tab_spanner(label = "Random Effects",
                  columns = gt::starts_with("random")) %>%
  gt::sub_missing() %>%
  gt::cols_label_with(columns = gt::starts_with("random"),
                      fn = function(x) paste0("")) %>%
  gt::tab_style(locations =
                  cells_body(rows = str_detect(dataset, "Eucalyptus"),
                             columns = dataset),
                style = cell_text(style = "italic")) %>%
  gt::text_transform(fn = function(x) str_replace(x, "publishable_as_is", "Ca
```

```
tegorical Peer Rating") %>%
                      str_replace(., "rate_analysis", "Continuous Peer Ratin
g"),
                     locations = cells_body(columns = c("fixed_effects"))) %>
%
   gt::text_transform(
   locations = cells_stub(
     rows = estimate_type != "y25"
    ),
    fn = function(x)
     paste0("")
    }
  ) %>%
  gt::tab_style(locations = cells_stub(rows = str_detect(dataset, "Eucalyptu
s")),
                style = cell_text(style = "italic")) %>%
gt_fmt_yi(columns = "estimate_type")
```

Table C.2: Singularity and convergence checking outcomes for models of deviation in out-ofsample predictions y_r explained by peer-review ratings for different random effect structures. Problematic checking outcomes are highlighted in red.

	Randon	n Effects				Can random	Can random
			Prediction Scenario	Model converged?	Singular Fit?	effects SE be calculated?	effect 95% CI be calculated?
Categorical	Peer Rating	9					
Eucalyptus	Reviewer ID	_	y_{25}	yes	no	yes	yes
	Reviewer ID	_	${y}_{50}$	yes	no	yes	yes
	Reviewer ID	_	y_{75}	yes	yes	no	yes
Eucalyptus	Study ID	Reviewer ID	y_{25}	yes	no	yes	yes
	Study ID	Reviewer ID	y_{50}	yes	no	yes	yes
	Study ID	Reviewer ID	y_{75}	yes	yes	no	yes
Eucalyptus	Study ID	_	y_{25}	yes	no	yes	yes
	Study ID	_	y_{50}	yes	no	yes	yes
	Study ID	_	y_{75}	yes	no	yes	yes

blue tit	Reviewer ID	_	y_{25}	yes	yes	no	yes
	Reviewer ID	_	${y}_{50}$	yes	no	yes	yes
	Reviewer ID	_	y_{75}	yes	no	yes	yes
blue tit	Study ID	Reviewer ID	y_{25}	yes	no	yes	yes
	Study ID	Reviewer ID	${y}_{50}$	yes	yes	no	yes
	Study ID	Reviewer ID	y_{75}	yes	no	yes	yes
blue tit	Study ID	_	y_{25}	yes	no	yes	yes
	Study ID	_	${y}_{50}$	yes	no	yes	yes
	Study ID	_	y_{75}	yes	no	yes	yes

Continuous	Peer Rating	g					
Eucalyptus	Reviewer ID	_	y_{25}	yes	no	yes	yes
	Reviewer ID	_	${y}_{50}$	yes	yes	no	yes
	Reviewer ID	_	y_{75}	yes	yes	no	yes
Eucalyptus	Study ID	Reviewer ID	y_{25}	yes	no	no	yes
	Study ID	Reviewer ID	y_{50}	no	-	_	_
	Study ID	Reviewer ID	y_{75}	yes	no	no	yes
Eucalyptus	Study ID	_	y_{25}	yes	no	no	yes
	Study ID	_	${y}_{50}$	yes	no	no	yes
	Study ID	_	y_{75}	yes	no	yes	yes

blue tit	Reviewer ID	_	y_{25}	yes	yes	no	yes
	Reviewer ID	_	y_{50}	yes	no	yes	yes
	Reviewer ID	_	y_{75}	yes	no	yes	yes
blue tit	Study ID	Reviewer ID	y_{25}	yes	no	no	yes
	Study ID	Reviewer ID	y_{50}	yes	no	no	yes
	Study ID	Reviewer ID	y_{75}	yes	no	no	yes
blue tit	Study ID	-	y_{25}	yes	no	no	yes
	Study ID	_	${y}_{50}$	yes	no	no	yes
	Study ID	_	y_{75}	yes	no	yes	yes

```
yi fitted mods <-
  ManyEcoEvo::ManyEcoEvo yi viz %>%
  filter(model_name %in% c("box_cox_rating_cat",
                           "box_cox_rating_cont",
                           "sorensen glm",
                           "uni mixed effects")) %>%
  select(-ends_with("_plot"), -MA_fit_stats, -contains("mod_")) %>%
  rowwise() %>%
  mutate(singularity = possibly_check_singularity(model),
         convergence = list(possibly_check_convergence(model))) %>%
  ungroup() %>% mutate(
    across(where(is.list),
           .fns = ~ coalesce(.x, list(NA)))
  ) %>%
  mutate(convergence = list_c(convergence),
         singularity = case_when(is.na(convergence) ~ NA,
                                 TRUE ~ singularity))
```

yi_convergence_singularity <-</pre>

```
vi fitted mods %>%
  left join({ # Check if SE and CI can be calculated
    yi_fitted_mods %>%
      unnest(model params) %>%
      filter(Effects == "random") %>%
      filter(if_any(contains("SE"), list(is.infinite, is.na))) %>%
      distinct(dataset, estimate type, model name) %>%
      mutate(SE_calc = FALSE)
  }, by = join_by(dataset, estimate_type, model_name)) %>%
  left_join({
    yi_fitted_mods %>%
      unnest(model params) %>%
      filter(Effects == "random") %>%
      filter(if_any(contains("CI_"), list(is.infinite, is.na))) %>%
      distinct(dataset, estimate_type, model_name) %>%
      mutate(CI calc = FALSE)
  }, by = join_by(dataset, estimate_type, model_name)) %>%
  rowwise() %>%
  mutate(across(ends with(" calc"),
                ~ replace_na(.x, TRUE)),
         across(c(SE_calc, CI_calc, singularity), ~ ifelse(is_false(convergen))
ce) | is_na(convergence), NA, .x)),
         model_name = forcats::as_factor(model_name),
         model name = forcats::fct relevel(model name,
                                           c("box cox rating cat",
                                             "box cox_rating_cont",
                                             "sorensen glm",
                                             "uni_mixed_effects")),
         model name =
           forcats::fct recode(
             model name,
             `Deviation explained by categorical ratings` = "box_cox_rating_c
at",
             `Deviation explained by continuous ratings` = "box_cox_rating_co
nt",
             `Deviation explained by Sorensen's index` = "sorensen glm",
             `Deviation explained by inclusion of random effects` =
               "uni_mixed_effects"),
         dataset = case_when(str_detect(dataset, "eucalyptus") ~ "Eucalyptus"
                             TRUE ~ dataset)) %>%
  ungroup() %>%
  select(-model)
yi_singularity_convergence_sorensen_mixed_mod <-</pre>
  yi_convergence_singularity %>%
 filter(str detect(model name, "Sorensen") | str detect(model name, "random"
))
```

We fitted the same deviation models on the out-of-sample-predictions dataset that we fitted for the effect-size dataset. However, while all models of deviation explained by categorical peer-ratings converged, the following datasets and prediction scenarios suffered from singular fit: blue tit - y_{25} , *Eucalyptus* - y_{75} (Table C.3). Models of deviation explained by *continuous* ratings all converged, however models for the out-of-sample predictions model fit was singular. Similarly to the effect-size (Z_r) dataset, SD and CI could not be estimated for random effects in some models (Table C.3), consequently we interpreted this to mean the models had singular fit (See Section C.3.1). Results of all deviation models are therefore presented only for models with non-singular fit, and that converged (Table C.3).

```
yi_convergence_singularity %>%
  filter(stringr::str detect(model name, "ratings")) %>%
  select(-model params) %>%
  group_by(model_name) %>%
  gt::gt(rowname col = "dataset") %>%
  gt::tab_style(locations =
                  cells_body(rows = str_detect(dataset, "Eucalyptus"),
                             columns = dataset),
                style = cell_text(style = "italic")) %>%
  gt::cols_label(dataset = "Dataset",
                 estimate_type = "Prediction Scenario",
                 singularity = "Singular Fit?",
                 convergence = "Model converged?",
                 SE_calc = gt::md("Can random effects $\\text{SE}$ be calcula
ted?"),
                 CI_calc = "Can random effect CI be calculated?") %>%
  gt::opt_stylize(style = 6, color = "gray") %>%
  gt::text_transform(fn = function(x) ifelse(x == TRUE, "yes",
                                             ifelse(x == FALSE, "no", x)),
                     locations = cells_body(columns = c("singularity",
                                                        "convergence",
                                                        "SE calc",
                                                        "CI calc")
                     )) %>%
  gt::text transform(
    locations = cells stub(
      rows = estimate_type != "y25"
    ),
    fn = function(x){
      paste0("")
    }
  ) %>%
  gt_fmt_yi("estimate_type") %>%
  gt::tab_style(locations = cells_stub(rows = str_detect(dataset, "Eucalyptus
")),
                style = cell_text(style = "italic")) %>%
 tab style(
    style = list(
     cell_fill(color = scales::alpha("red", 0.6)),
```

```
cell_text(color = "white", weight = "bold")
),
locations = list(
   cells_body(columns = "singularity", rows = singularity == TRUE),
   cells_body(columns = "convergence", rows = convergence == FALSE),
   cells_body(columns = "SE_calc", rows = SE_calc == FALSE),
   cells_body(columns = "CI_calc", rows = CI_calc == FALSE)
)
) %>%
gt::sub_missing()
```

Table C.3: Singularity and convergence checking for models of deviation in out-of-samplepredictions y_i explained by peer-ratings.

	Prediction Scenario	Singular Fit?	Model converged?	Can random effects SE be calculated?	Can random effect CI be calculated?
Deviation explai	ned by continuou	s ratings			
blue tit	y_{25}	yes	yes	no	no
	${y}_{50}$	no	yes	yes	yes
	y_{75}	no	yes	yes	yes
Eucalyptus	y_{25}	no	yes	yes	yes
	${y}_{50}$	yes	yes	no	no
	y_{75}	yes	yes	no	no

Deviation explained by categorical ratings

blue tit	y_{25}	yes	yes	no	no
	${y}_{50}$	no	yes	yes	yes
	y_{75}	no	yes	yes	yes
Eucalyptus	y_{25}	no	yes	yes	yes
	${y}_{50}$	no	yes	yes	yes
	y_{75}	yes	yes	no	no

Group means and 95% confidence intervals for different categories of peer-review rating are all overlapping (Figure C.2). The fixed effect of peer review rating also explains virtually no variability in deviation scores for out-of-sample predictions y_i (Table C.3).

```
yi_violin_cat_plot_data <-</pre>
  ManyEcoEvo::ManyEcoEvo yi viz %>%
  filter(model_name %in% "box_cox_rating_cat") %>%
  left_join(.,
            {ManyEcoEvo::ManyEcoEvo yi results %>%
                select(dataset, estimate_type, effects_analysis) %>%
                hoist(effects_analysis, "lambda", .transform = unique) %>%
                select(-effects analysis)},
            by = join by(dataset, estimate type)) %>%
 mutate( dataset = case_when(str_detect(dataset, "eucalyptus") ~ "Eucalyptus")
",
                              TRUE ~ dataset)) %>%
  semi_join({
    yi convergence singularity %>%
      filter( str_detect(model_name, "categorical"),
              !singularity, convergence, SE_calc, CI_calc)
    }, by = join_by("dataset", "estimate_type")) %>%
  select(dataset, estimate_type, model_name, model) %>%
  mutate(predictor_means =
           map(model, modelbased::estimate means, backend = "marginaleffects"
),
         model data = map(model, ~pluck(.x, "frame") %>%
                            drop_na() %>%
                            as_tibble()),
         plot_name = paste(dataset,
                           estimate type,
                           "violin cat",
                           sep = " ")) %>%
 mutate(model data = map(model data,
                          .f = ~ mutate(.x, PublishableAsIs =
                                           str replace(PublishableAsIs,
                                                       "publishable with ", ""
) %>%
                                           str replace("deeply flawed and ", "
") %>%
                                           capwords())),
         predictor means = map(predictor means,
                                .f = ~ mutate(.x, PublishableAsIs =
                                                str_replace(PublishableAsIs,
                                                            "publishable with
", "") %>%
                                                str_replace("deeply flawed and
", "") %>%
                                                capwords())) %>%
  select(-model)
```

```
yi violin cat plots <- yi violin cat plot data %>%
  pmap(.1 = list(.$model data, .$predictor means, .$plot name),
       .f = ~ plot_model_means_box_cox_cat(..1,
                                           PublishableAsIs,
                                           ...2,
                                           new order =
                                             c("Unpublishable",
                                               "Major Revision",
                                               "Minor Revision",
                                               "Publishable As Is"),
                                           purrr::set names({yi violin cat plot data %>%
      pull(plot name) %>%
      stringr::str_split("_violin_cat", 2) %>%
      map_chr(pluck, 1) })
subfigcaps_yi_cat <- yi_violin_cat_plot_data %>%
  mutate(dataset =
           case when(dataset == "Eucalyptus" ~ paste0("*", dataset, "*"),
                     TRUE ~ Hmisc::capitalize(dataset))) %>%
  unite(plot name, dataset, estimate type, sep = ", ") %>%
  pull(plot_name)
fig cap yi deviation cat rating <-
  paste0("Violin plot of Box-Cox transformed deviation from the meta-analytic
mean for $y i$ estimates as a function of categorical peer-review ratings rat
ings. Note that higher (less negative) values of the deviation score result f
rom greater deviation from the meta-analytic mean. Grey points for each ratin
g group denote model-estimated marginal mean deviation, and error bars denote
95% CI of the estimate. ", subfigcaps_yi_cat %>%
           paste0(paste0(paste0("**", LETTERS[1:length(subfigcaps_yi_cat)], "
**", sep = ""), sep = ": "), ., collapse = ", "), ".")
library(patchwork)
patchwork::wrap_plots(yi_violin_cat_plots, ncol = 2, nrow = 2, guides = 'coll
ect') +
patchwork::plot annotation(tag levels = 'A')
```



Figure C.2: Violin plot of Box-Cox transformed deviation from the meta-analytic mean for y_i estimates as a function of categorical peer-review ratings ratings. Note that higher (less negative) values of the deviation score result from greater deviation from the meta-analytic mean. Grey points for each rating group denote model-estimated marginal mean deviation, and error bars denote 95% CI of the estimate. **A**: Blue tit, y50, **B**: Blue tit, y75, **C**: Eucalyptus, y25, **D**: Eucalyptus, y50.

There was a lack of any clear relationships between quantitative review scores and y_i deviation scores (Table C.11). Plots of these relationships indicated either no relationship or extremely weak positive relationships (Figure C.3). Recall that positive relationships mean that as review scores became more favorable, the deviation from the meta-analytic mean increased, which is surprising. Because almost no variability in y_i deviation score was explained by reviewer ratings (Table C.11), this pattern does not appear to merit further consideration.

```
vi cont plot data <-
  ManyEcoEvo::ManyEcoEvo yi viz %>%
  filter(model_name %in% c("box_cox_rating_cont")) %>%
  mutate(dataset = case_match(dataset, "eucalyptus" ~ "Eucalyptus",.default =
dataset)) %>%
  semi_join({yi_convergence_singularity %>%
      filter( str detect(model name, "cont"), # Omit all in-estimable models
              !singularity,
              convergence,
              #SE calc,
              #CI calc
      )},
      by = join_by("dataset", "estimate_type")) %>%
  select(dataset, estimate_type, model_name, model) %>%
  mutate(plot_data = map(model, pluck, "frame"))
subfigcaps <- yi cont plot data %>%
  mutate(dataset =
           case_when(dataset == "Eucalyptus" ~ paste0("*", dataset, "*"),
                     TRUE ~ Hmisc::capitalize(dataset))) %>%
  unite(plot name, dataset, estimate type, sep = ", ") %>%
  pull(plot name)
fig_cap_yi_deviation_cont_rating <-</pre>
  paste0("Scatterplots explaining Box-Cox transformed deviation from the meta
-analytic mean for $y i$ estimates as a function of continuous ratings. Note
that higher (less negative) values of the deviation score result from greater
deviation from the meta-analytic mean. ", subfigcaps %>%
           paste0(paste0(paste0("**", LETTERS[1:length(subfigcaps)], "**", se
p = ""), sep = ": "), ., collapse = ", "), ".")
yi_cont_plots <-</pre>
  yi_cont_plot_data$plot_data %>%
  map(.f = ~ plot_continuous_rating(.x)) %>%
  purrr::set_names({yi_cont_plot_data %>%
      unite(plot name, dataset, estimate type, sep = " ") %>%
      pull(plot name)})
patchwork::wrap_plots(yi_cont_plots, heights = 4, byrow = TRUE) +
  patchwork::plot annotation(tag levels = 'A')
```



Figure C.3: Scatterplots explaining Box-Cox transformed deviation from the meta-analytic mean for y_i estimates as a function of continuous ratings. Note that higher (less negative) values of the deviation score result from greater deviation from the meta-analytic mean. **A**: Blue tit, y50, **B**: Blue tit, y75, **C**: Eucalyptus, y25.

```
ManyEcoEvo yi viz %>%
  filter(
    model name %nin% c("MA mod",
                        "box cox rating cat no int",
                       "MA_mod_mv")) %>%
  mutate( dataset = case_when(str_detect(dataset, "eucalyptus") ~ "Eucalyptus")
۳,
                              TRUE ~ dataset),
          model_name = forcats::as_factor(model_name) %>%
            forcats::fct_relevel(c("box_cox_rating_cat",
                                    "box_cox_rating_cont",
                                    "sorensen glm",
                                    "uni mixed effects")) %>%
            forcats::fct_recode(
              `Deviation explained by categorical ratings` = "box_cox_rating_
cat",
              `Deviation explained by continuous ratings` = "box_cox_rating_c
ont",
              `Deviation explained by Sorensen's index` = "sorensen_glm",
```

```
`Deviation explained by inclusion of random effects` = "uni_mi
xed effects")
  ) %>%
  semi_join(
    {yi_convergence_singularity %>%
        filter(!singularity,
               convergence,
               SE_calc,
               CI_calc) },
    by = join_by("dataset", "estimate_type", "model_name")
  ) %>%
  select(dataset,
         estimate type,
         model_name,
         model_params) %>%
  unnest(model params) %>%
  mutate(
    Group = case_match(Group,
                       "study_id" ~ "Effect ID",
                       "ReviewerId" ~ "Reviewer ID",
                       "" ~ NA,
                       .default = Group),
    df_error = as.integer(df_error),
    Parameter = str remove(Parameter, "PublishableAsIs") %>%
      str_replace("diversity", "Sorensen's") %>%
      str_replace_all(., "_", " ") %>%
      str remove(., "1") %>%
      Hmisc::capitalize() ) %>%
  group by(model name) %>%
  arrange(model name,
          dataset, estimate_type) %>%
  select(-CI) %>%
  gt::gt(rowname_col = "dataset") %>%
  gt::fmt(columns = "p",
          fns = function(x) gtsummary::style_pvalue(x)) %>%
  gt::cols label(CI low = gt::md("95\\%CI"),
                 estimate_type = "Prediction Scenario",
                 SE = gt::md("$\\text{SE}$"),
                 df_error = gt::md("$\\mathit{df}$"),
                 t = gt::md("$t$"),
                 p = gt::md("*p*")) %>%
  gt::cols_merge(columns = starts_with("CI_"),
                 pattern = "[{1},{2}]") %>%
  gt::cols_move(columns = CI_low, after = SE) %>%
  gt::opt_stylize(style = 6, color = "gray") %>%
  gt::fmt(columns = c(Coefficient, SE, starts_with("CI_"), t) ,
          rows = Parameter %nin% c("RateAnalysis", "SD (Observations)", "mixe
d model1"),
          fns = function(x) format(round(x, 2),nsmall = 2)) %>%
gt::fmt(columns = c(Coefficient, SE, t, starts_with("CI_")) ,
```

```
rows = Parameter %in% c("RateAnalysis", "SD (Observations)", "mixed
model1"),
         fns = function(x) ifelse(x < 0.0009,
                                   format(x, nsmall = 2, digits = 1),
                                   round(x, digits = 2))) %>%
 gt::cols_move(columns = c(Effects, Group), after = Parameter) %>%
 gt::sub missing(columns = c(Effects, Group, t, df error, p),
                  missing_text = "") %>%
 gt::cols_hide(Effects) %>%
 gt::text_transform(fn = function(x) map(x, gt::md),
                     locations = gt::cells_row_groups()) %>%
 gt::text transform(
    locations = cells stub(
     rows = Parameter != "(Intercept)"
    ),
   fn = function(x)
     paste0("")
    }
 ) %>%
 gt::fmt_number(columns = c(Coefficient, SE, t, starts_with("CI_")), decimal
s = 2,drop_trailing_zeros = TRUE, drop_trailing_dec_mark = TRUE) %>%
 gt::fmt_scientific(columns = c( starts_with("CI_")),
                     rows = abs(CI low) < 0.01 | abs(CI high) < 0.01 | abs(CI
low) > 1000 | abs(CI high) > 1000,
                     decimals = 2) %>%
 gt::fmt_scientific(columns = c( starts_with("Coefficient")),
                     rows = abs(Coefficient) < 0.01 | abs(Coefficient) > 1000
ر
                     decimals = 2) %>%
 gt::fmt scientific(columns = c( starts with("SE")),
                     rows = abs(SE) < 0.01 | abs(SE) > 1000,
                     decimals = 2) %>%
 gt::tab style(locations = gt::cells stub(rows = str detect(dataset, "Eucaly
ptus")),
                style = cell text(style = "italic")) %>%
 gt::cols label(Group = "Random Effect") %>%
gt_fmt_yi("estimate_type")
```

Table C.4: Parameter estimates for univariate models of Box-Cox transformed deviation from the mean y_i estimate as a function of categorical peer-review rating, continuous peer-review rating, and Sorensen's index for blue tit and Eucalyptus analyses, and also for the inclusion of random effects for Eucalyptus analyses.

	Prediction Scenario	Parameter	Random Effect	Coefficient	${ m SE}$	95%Cl	t	df	p
Deviation expla	ined by catego	rical ratings							
Eucalyptus	y_{25}	(Intercept)		-0.35	0.32	[-0.98,0.28]	-1.1	168	0.3
	y_{25}	Publishable with major revision		0.29	0.35	[-0.39,0.98]	0.84	168	0.4
	${y}_{25}$	Publishable with minor revision		0.41	0.34	[-0.26,1.07]	1.21	168	0.2
	y_{25}	Publishable as is		0.35	0.36	[-0.35,1.06]	0.99	168	0.3
	y_{25}	SD (Intercept)	Reviewer ID	0.23	0.15	[0.07,0.79]			
	y_{25}	SD (Observations)	Residual	0.92	0.06	[0.82,1.04]			
Eucalyptus	y_{50}	(Intercept)		-0.61	0.4	[-1.4,0.18]	-1.52	165	0.13
	y_{50}	Publishable with major revision		0.18	0.44	[-0.69,1.05]	0.4	165	0.7
	y_{50}	Publishable with minor revision		0.25	0.42	[-0.59,1.09]	0.58	165	0.6
	${y}_{50}$	Publishable as is		0.09	0.47	[-0.83,1.01]	0.19	165	0.8
	${y}_{50}$	SD (Intercept)	Reviewer ID	0.12	0.42	$[1.50 \times 10^{-4}, 1.02 \times 10^{2}]$			
	y_{50}	SD (Observations)	Residual	1.29	0.08	[1.14,1.46]			

blue tit	y_{50}	(Intercept)		-1.23	0.29	[-1.81,-0.65]	-4.18	218	<0.001
	y_{50}	Publishable with major revision		-0.21	0.3	[-0.81,0.39]	-0.69	218	0.5
	y_{50}	Publishable with minor revision		-0.25	0.3	[-0.85,0.34]	-0.83	218	0.4
	y_{50}	Publishable as is		-0.42	0.32	[-1.05,0.21]	-1.33	218	0.2
	y_{50}	SD (Intercept)	Reviewer ID	0.22	0.09	[0.11,0.47]			
	y_{50}	SD (Observations)	Residual	0.71	0.04	[0.64,0.8]			
blue tit	y_{75}	(Intercept)		-1.51	0.27	[-2.05,-0.97]	-5.52	231	<0.001
	y_{75}	Publishable with major revision		0.09	0.28	[-0.46,0.64]	0.33	231	0.7
		Publishable							
	y_{75}	with minor revision		0.35	0.28	[-0.2,0.9]	1.25	231	0.2
	y_{75} y_{75}			0.35	0.28	[-0.2,0.9] [-0.19,0.97]	1.25 1.33	231 231	0.2
		revision Publishable as	Reviewer ID						
Deviation explained by continuous ratings

Eucalyptus	y_{25}	(Intercept)		-0.46	0.26	[-0.97,0.06]	-1.76	170	0.08
	y_{25}	RateAnalysis		6.14 × 10 ⁻³	3.54 × 10 ⁻³	[-8.61 × 10 ⁻⁴ ,1.31 × 10 ⁻²]	1.73	170	0.08
	y_{25}	SD (Intercept)	Reviewer ID	0.12	0.21	[4.27 × 10 ⁻³ ,3.64]			
	y_{25}	SD (Observations)	Residual	0.93	0.06	[0.82,1.05]			
olue tit	y_{50}	(Intercept)		-1.35	0.24	[-1.82,-0.88]	-5.65	220	<0.00
	y_{50}	RateAnalysis		-1.82 × 10 ⁻³	3.05 × 10 ⁻³	[-7.83 × 10 ⁻³ ,4.18 × 10 ⁻³]	-0.6	220	0.0
	y_{50}	SD (Intercept)	Reviewer ID	0.24	0.08	[0.12,0.47]			
	y_{50}	SD (Observations)	Residual	0.71	0.04	[0.64,0.79]			
olue tit	y_{75}	(Intercept)		-1.66	0.22	[-2.1,-1.23]	-7.52	233	<0.00
	y_{75}	RateAnalysis		5.62 × 10 ⁻³	2.79 × 10 ⁻³	[1.22 × 10 ⁻⁴ , 1.11 × 10 ⁻²]	2.01	233	0.045
	y_{75}	SD (Intercept)	Reviewer ID	0.3	0.06	[0.2,0.45]			
	y_{75}	SD (Observations)	Residual	0.63	0.03	[0.57,0.7]			

Deviation explain	ned by Sorens	sen's index						
Eucalyptus	y_{25}	(Intercept)	-0.3	1.48	[-3.2,2.59]	-0.21	36	0.8
	y_{25}	Mean Sorensen's index	0.44	2.19	[-3.86,4.74]	0.2	36	0.8
Eucalyptus	y_{50}	(Intercept)	-1.32	2.1	[-5.43,2.79]	-0.63	36	0.5
	${y}_{50}$	Mean Sorensen's index	1.32	3.16	[-4.87,7.51]	0.42	36	0.7
Eucalyptus	y_{75}	(Intercept)	-0.71	1.78	[-4.19,2.78]	-0.4	36	0.7
	y_{75}	Mean Sorensen's index	0.34	2.71	[-4.96,5.64]	0.12	36	>0.9
blue tit	y_{25}	(Intercept)	-0.77	0.6	[-1.94,0.4]	-1.29	61	0.2
	y_{25}	Mean Sorensen's index	-0.23	1.04	[-2.27,1.82]	-0.22	61	0.8
blue tit	y_{50}	(Intercept)	-0.43	0.73	[-1.86,0.99]	-0.6	58	0.6
	${y}_{50}$	Mean Sorensen's index	-1.77	1.27	[-4.26,0.71]	-1.4	58	0.2
blue tit	y_{75}	(Intercept)	-1.72	0.74	[-3.16,-0.28]	-2.34	61	0.019
	y_{75}	Mean Sorensen's index	0.78	1.28	[-1.73,3.29]	0.61	61	0.5

Deviation explained by inclusion of random effects

Eucalyptus	y_{25}	(Intercept)	-0.53	0.26	[-1.05,-0.02]	-2.03	36	0.042
	y_{25}	Mixed model	0.74	0.31	[0.13,1.35]	2.37	36	0.018
Eucalyptus	y_{50}	(Intercept)	-0.57	0.4	[-1.36,0.21]	-1.43	36	0.2
	y_{50}	Mixed model	0.18	0.48	[-0.75,1.11]	0.38	36	0.7
Eucalyptus	y_{75}	(Intercept)	-0.35	0.39	[-1.11,0.41]	-0.9	36	0.4
	y_{75}	Mixed model	-0.19	0.46	[-1.1,0.71]	-0.41	36	0.7

C.4 Deviation scores as explained by the distinctiveness of variables in each analysis

C.4.1 Out of sample predictions y_i

Given the convergence and singularity issues encountered with most other analyses, we also checked for convergence and singularity issues in models of deviation explained by Sorensen's similarity index for y_i estimates (Table C.5). All models fitted without problem.

```
yi_sorensen_plot_data <-</pre>
  ManyEcoEvo_yi_viz %>%
  filter(str_detect(model_name, "sorensen_glm")) %>%
  mutate( dataset =
            case when(str detect(dataset, "eucalyptus") ~ "Eucalyptus",
                      TRUE ~ dataset),
          model_name = forcats::as_factor(model_name) %>%
            forcats::fct_relevel(c("box_cox_rating_cat",
                                    "box cox rating cont",
                                   "sorensen glm",
                                    "uni mixed_effects")) %>%
            forcats::fct recode(
              `Deviation explained by categorical ratings` = "box_cox_rating_
cat",
              `Deviation explained by continuous ratings` = "box cox rating c
ont",
              `Deviation explained by Sorensen's index` = "sorensen glm",
              `Deviation explained by inclusion of random effects` = "uni mi
xed effects")) %>%
  select(dataset, estimate type, model name, model) %>%
  semi join(
    {yi convergence singularity %>%
        filter(!singularity,
               convergence,
               SE_calc, CI_calc) },
    by = join_by("dataset", "estimate_type", "model_name")
  ) %>%
  mutate(
    plot_data = map(model, ~ pluck(.x, "fit", "data") %>%
                      rename(box_cox_abs_deviation_score_estimate = ..y))) %>
%
  unite(plot_names, dataset, estimate_type, sep = ", ")
yi sorensen subfigcaps <-</pre>
  yi_sorensen_plot_data$plot_names %>%
  paste0(paste0(paste0("**", LETTERS[1:length(yi_sorensen_plot_data$plot_name
s)], "**", sep = ""), sep = ": "), ., collapse = ", ")
```

yi_sorensen_fig_cap <- paste0("Scatter plots examining Box-Cox transformed de



Figure C.4: Scatter plots examining Box-Cox transformed deviation from the meta-analytic mean for y_i estimates as a function of Sorensen's similarity index. Note that higher (less negative) values of the deviation score result from greater deviation from the meta-analytic mean (Figure C.1). A: blue tit, y25, B: blue tit, y50, C: blue tit, y75, D: Eucalyptus, y25, E: Eucalyptus, y50, F: Eucalyptus, y75.

We checked the fitted models for the inclusion of random effects for the *Eucalyptus* dataset, and for models of deviation explained by Sorensen's similarity index for y_i estimates (Table C.5). All models converged, and no singular fits were encountered.

```
yi singularity convergence sorensen mixed mod %>%
  drop_na(convergence) %>%
  mutate(across(c(SE_calc, CI_calc, singularity), ~ ifelse(is_false(convergen
ce), NA, .x))) %>%
  select(-model params) %>%
  group by(model name) %>%
  gt::gt(rowname_col = "dataset") %>%
 gt::tab_style(locations = cells body(rows = str_detect(dataset, "Eucalyptus
"),
                                       columns = dataset),
                style = cell_text(style = "italic")) %>%
  gt::cols_label(dataset = "Dataset",
                 estimate_type = "Prediction Scenario",
                 singularity = "Singular Fit?",
                 convergence = "Model converged?",
                 SE_calc = gt::md("Can random effects $\\text{SE}$ be calcula
ted?"),
                 CI calc = "Can random effects CI be calculated?") %>%
  gt::opt_stylize(style = 6, color = "gray") %>%
  tab_style(
    style = list(
      cell fill(color = scales::alpha("red", 0.6)),
      cell_text(color = "white", weight = "bold")
    ),
    locations = list(
      cells_body(columns = "singularity", rows = singularity == TRUE),
      cells_body(columns = "convergence", rows = convergence == FALSE),
      cells_body(columns = "SE_calc", rows = SE_calc == FALSE),
      cells_body(columns = "CI_calc", rows = CI_calc == FALSE)
    )) %>%
  gt::text transform(fn = function(x) ifelse(x == TRUE, "yes",
                                              ifelse(x == FALSE, "no", x)),
                     locations = cells_body(columns = c("singularity",
                                                         "convergence",
                                                        "SE calc",
                                                        "CI calc")
                     )) %>%
  gt::text_transform(
```

```
locations = cells_stub(
    rows = estimate_type != "y25"
    ),
    fn = function(x){
        paste0("")
     }
        ) %>%
    gt_fmt_yi("estimate_type") %>%
    gt::tab_style(locations = cells_stub(rows = str_detect(dataset, "Eucalyptus
")),
        style = cell_text(style = "italic"))
```

Table C.5: Singularity and convergence checks for models of deviation explained by Sorensen's similarity index and inclusion of random effects for out-of-sample predictions, y_i . Models of Deviation explained by inclusion of random effects are not presented for blue tit analyses because the number of models not using random effects was less than our preregistered threshold.

	Prediction Scenario	Singular Fit?	Model converged?	Can random effects SE be calculated?	Can random effects CI be calculated?
Deviation expla	ined by Sorense	n's index			
blue tit	y_{25}	no	yes	yes	yes
	${y}_{50}$	no	yes	yes	yes
	y_{75}	no	yes	yes	yes
Eucalyptus	y_{25}	no	yes	yes	yes
	${y}_{50}$	no	yes	yes	yes
	y_{75}	no	yes	yes	yes

Deviation explained by inclusion of random effects

Eucalyptus	y_{25}	no	yes	yes	yes
	${y}_{50}$	no	yes	yes	yes
	y_{75}	no	yes	yes	yes

C.5 Deviation scores as explained by the inclusion of random effects

C.5.1 Out of sample predictions y_i

Only 1 of the Blue tit out-of-sample analyses y_i included random effects, which was below our preregistered threshold of 5 for running the models of Box-Cox transformed deviation from the meta-analytic mean explained by the inclusion of random-effects. However, 14 *Eucalyptus* analyses included in the out-of-sample y_i results included only fixed effects, which crossed our pre-registered threshold.

Consequently, we performed this analysis for the *Eucalyptus* dataset only, here we present results for the out of sample prediction y_i results. There is inconsistent evidence of somewhat higher Box-Cox-transformed deviation values for models including a random effect, meaning the analyses of the *Eucalyptus* dataset that included random effects averaged slightly higher deviation from the meta-analytic mean out-of-sample estimate in the relevant prediction scenario. This is most evident for the y_{25} predictions which both shows the greatest difference in Box-Cox transformed deviation values (Figure C.5) and explains the most variation in y_i deviation score (Table C.4).

```
yi_deviation_RE_plot_data <-</pre>
  ManyEcoEvo yi results %>%
  mutate(dataset = Hmisc::capitalize(dataset)) %>%
  semi_join({yi_singularity_convergence_sorensen_mixed_mod %>% filter(!singul
arity, convergence, SE calc, CI calc, str detect(model name, "random"))}, by
= join_by(dataset, estimate_type)) %>%
  select(dataset, estimate_type, model = uni_mixed_effects) %>%
  rowwise() %>%
  filter(!is_logical(model)) %>% ungroup %>%
  mutate(predictor_means = map(model, .f = ~ pluck(.x, "fit") %>%
                                 modelbased::estimate means(.)),
         plot_data = map(model, pluck, "fit", "data"),
         plot_data = map(plot_data,
                         rename,
                         box_cox_abs_deviation_score_estimate = ...y)) %>%
  mutate(dataset = case when(str detect(dataset, "Eucalyptus") ~ paste0("*",
dataset, "*"), TRUE ~ dataset)) %>%
  unite(plot_names, dataset, estimate_type, sep = ", ")
yi_deviation_RE_plot_subfigcaps <- yi_deviation_RE_plot_data %>%
  pull(plot names)
yi deviation RE plot figcap <-
  paste0("Violin plot of Box-Cox transformed deviation from meta-analytic mea
n as a function of presence or absence of random effects in the analyst's mod
el. White points for each rating group denote model-estimated marginal mean d
eviation, and error bars denote 95% CI of the estimate. Note that higher (les
```

s negative) values of Box-Cox transformed deviation result from greater devia

tion from the meta-analytic mean. ",



```
Figure C.5: Violin plot of Box-Cox transformed deviation from meta-analytic mean as a function of presence or absence of random effects in the analyst's model. White points for each rating group denote model-estimated marginal mean deviation, and error bars denote 95% CI of the estimate. Note that higher (less negative) values of Box-Cox transformed deviation result from greater deviation from the meta-analytic mean. A: Eucalyptus, y25, B: Eucalyptus, y50, C: Eucalyptus, y75.
```

No Random Effects Random effects

C.6 Multivariate Analysis

C.6.1 Effect Sizes Z_r

```
collinearity_subset == "All")
multivar_mods <-</pre>
  ManyEcoEvo viz %>%
  dplyr::filter(!!!filter vars, model name == "MA mod mv") %>%
  hoist(mod fit stats, "R2 conditional", "R2 marginal", "Sigma")
bt multivar mod R <-
  multivar mods %>%
  ungroup %>%
  filter(dataset == "blue tit") %>%
  select(R2_marginal, R2_conditional) %>%
  transpose() %>%
  flatten_dbl()
euc multivar mod R <-
  multivar mods %>%
  ungroup %>%
  filter(dataset == "eucalyptus") %>%
  select(R2_marginal, R2_conditional) %>%
  transpose() %>%
  flatten_dbl()
bt multivar mod sigma <- multivar mods %>%
  filter(dataset == "blue tit") %>%
  round_pluck("Sigma")
euc multivar mod sigma <- multivar mods %>%
  filter(dataset == "eucalyptus") %>%
  round pluck("Sigma")
multivar_mods %>%
  select(dataset, model params) %>%
  unnest(model_params) %>%
  select(-CI) %>%
  mutate(
    dataset =
      str_replace(dataset, "eucalyptus", "*Eucalyptus*"),
    Parameter =
      str_replace(Parameter, "mixed_model", "random_included")) %>%
  group by(dataset) %>%
  gt::gt() %>%
  gt::fmt_number(columns = c(Coefficient, SE, starts_with("CI_"), t),
                 decimals = 2,
                 drop trailing zeros = TRUE,
                 drop_trailing_dec_mark = TRUE) %>%
  gt::fmt scientific(
    columns = c( starts with("CI ")),
    rows = abs(CI_low) < 0.01 | abs(CI_high) < 0.01 | abs(CI_low) > 1000 | ab
```

```
s(CI high) > 1000,
    decimals = 2) %>%
  gt::fmt_scientific(
    columns = c( starts_with("Coefficient")),
    rows = abs(Coefficient) < 0.01 | abs(Coefficient) > 1000,
    decimals = 2) %>%
  gt::fmt(columns = "p",
          fns = function(x) gtsummary::style pvalue(x,
                                                      prepend_p = TRUE)
  ) %>%
  gt::cols_label(CI_low = gt::md("95\\%CI"),
                 df error = "df",
                 p = gt::md("*p*"),
                 SE = gt::md("$\\text{SE}$")) %>%
  gt::cols_merge(columns = starts_with("CI_"),
                 pattern = "[\{1\}, \{2\}]") %>%
  gt::cols_move(columns = CI_low, after = SE) %>%
  gt::opt stylize(style = 6, color = "gray") %>%
  gt::cols move(columns = c(Effects, Group), after = Parameter) %>%
  gt::text_transform(fn = function(x){
    str_remove(x, "PublishableAsIs") %>%
    str_replace_all("_", " ") %>%
      str_replace("diversity", "Sorensen's") %>%
      Hmisc::capitalize()
  },
  locations = cells_body(columns = Parameter)) %>%
  gt::text transform(fn = function(x) str replace(x, "ReviewerId", "Reviewer
ID")) %>%
  gt::text_transform(fn = function(x) map(x, gt::md),
                     locations = gt::cells row groups()) %>%
  gt::sub_missing(missing_text = "") %>%
  gt::cols_hide(Effects) %>%
  gt::cols_label(Group = "Random Effect")
multivar_mod_tidy <- multivar_mods %>%
  pull(model, name = "dataset") %>%
  map dfr(broom.mixed::tidy, conf.int = TRUE, .id = "dataset")
multivar performance tidy <- multivar mods %>%
  pull(model, name = "dataset") %>%
  map_dfr(performance::performance, .id = "dataset")
```

Table C.6: Parameter estimates from models explaining Box-Cox transformed deviation scores from the mean Z_r as a function of continuous and categorical peer-review ratings in multivariate analyses. Standard Errors (SE), 95% confidence intervals (95% CI) are reported for all estimates, while t values, degrees of freedom (df) and p-values are presented for fixed-effects.

Parameter	Random Effect	Coefficient	SE	95%CI	t	df	p
blue tit							
(Intercept)		-1.24	0.24	[-1.71,-0.77]	-5.16	465	p<0.001
RateAnalysis		-2.84×10^{-3}	0	$[-7.81 \times 10^{-3}, 2.13 \times 10^{-3}]$	-1.12	465	p=0.3
Publishable as is		0.06	0.2	[-0.33,0.45]	0.3	465	p=0.8
Publishable with major revision		-0.09	0.14	[-0.36,0.18]	-0.67	465	p=0.5
Publishable with minor revision		-0.02	0.17	[-0.36,0.31]	-0.14	465	p=0.9
Mean Sorensen's index		0.33	0.27	[-0.2,0.87]	1.22	465	p=0.2
SD (Intercept)	Reviewer ID	0.16	0.03	[0.11,0.25]			
SD (Observations)	Residual	0.5	0.02	[0.46,0.53]			

Eucalyptus							
(Intercept)		-2.8	0.77	[-4.32,-1.27]	-3.61	337	p<0.001
RateAnalysis		-0.01	0.01 [-2	2.27 × 10 ⁻² ,1.74 × 10 ⁻³]	-1.69	337	p=0.093
Publishable as is		0.76	0.54	[-0.3,1.82]	1.42	337	p=0.2
Publishable with major revision		0.65	0.36	[-0.06,1.35]	1.79	337	p=0.074
Publishable with minor revision		0.55	0.44	[-0.32,1.41]	1.24	337	p=0.2
Mean Sorensen's index		0.36	0.91	[-1.43,2.15]	0.39	337	p=0.7
Random included		0.19	0.2	[-0.2,0.59]	0.97	337	p=0.3
SD (Intercept)	Reviewer ID	0.38	0.09	[0.24,0.61]			
SD (Observations)	Residual	1.06	0.04	[0.98,1.15]			

The multivariate models did a poor job of explaining how different from the meta-analytic mean each analysis would be. For the blue tit analyses the R^2 value for the whole model was 0.11 and for the fixed effects component was 0.01, and the residual standard deviation for the model was 0.5. Further, all of the fixed effects had 95% confidence intervals that overlaped 0. This evidence is all consistent with none of the predictor variables in this model (continuous review rating, categorical review rating, distinctiveness of variables included) having any meaningful effect on how far Z_r estimates fell from the meta-analytic mean for the blue tit analyses. The pattern is largely similar for the *Eucalyptus* multivariate analysis, in which R^2 for the whole model was 0.13 and for the fixed effects component was 0.02, and the residual standard deviation for the model was 1.06. There is somewhat more of a hint of a pattern when examining the parameter estimates from the *Eucalyptus* analysis. In the case of the fixed effect of categorical reviewer ratings, analyses that were reviewed as 'publishable as is' and 'publishable with major revisions' appeared to produce results more different from the metaanalytic mean than those that were in the reference class of 'deeply flawed and unpublishable'. However, the estimates are very uncertain (*Eucalyptus* fixed effect for 'publishable as is' 0.76 (95% CI -0.29,1.81), and for 'publishable with major revision' 0.06 (95% CI -0.33,0.45)). Further, the collinearity between the categorical and continuous ratings make interpretation of effects involving either of these two variables unclear, and so we recommend against interpreting the pattern observed here. We report this analysis only for the sake of transparency.

```
multivar performance tidy %>%
  select(dataset, starts_with("R2_"), ICC, RMSE, Sigma) %>%
  mutate(dataset =
           case_when(str_detect(dataset, "eucalyptus") ~ "Eucalyptus",
                     TRUE ~ dataset)) %>%
  gt::gt() %>%
  gt::fmt(columns = function(x) rlang::is_bare_numeric(x),
          fns = function(x) round(x, 2)) %>%
  gt::opt_stylize(style = 6, color = "gray") %>%
  gt::cols label(R2 conditional = gt::md("$$R^{2} \\text{Conditional}$$"),
                 R2 marginal = gt::md("$$R^{2}_\\text{Marginal}$$"),
                 Sigma = gt::md("$$\\sigma$$"),
                 dataset = "Dataset") %>%
  gt::tab_style(locations =
                  cells_body(rows = str_detect(dataset, "Eucalyptus"),
                            columns = dataset),
                style = cell_text(style = "italic")) %>%
  gt::fmt_number(columns = c(R2_conditional, R2_marginal, ICC, Sigma),
                 decimals = 2,
                 drop_trailing_zeros = TRUE,
                 drop_trailing_dec_mark = TRUE)
```

Table C.7: Model summary metrics for multivariate models. σ is the residual standard deviation, ICC is the intra-class correlation coefficient, and R_M^2 and R_C^2 are the marginal and conditional R^2 , respectively.

Dataset	$R^2_{ m Conditional}$	$R^2_{ m Marginal}$	ICC	RMSE	σ
blue tit	0.11	0.01	0.1	0.48	0.5
Eucalyptus	0.13	0.02	0.11	1.02	1.06

C.6.2 Out of sample predictions y_i

For the blue tit analyses, the only models that did converge, which were not singular and that had estimable random effect variances were the y_{50} and y_{75} prediction scenarios with Reviewer ID as the model random effect (Table C.8). Of the different random effects structures we trialled for the *Eucalyptus* analyses, only the model that included Reviewer ID as the random effect successfully fitted to the y_{50} and y_{75} prediction scenarios, with other models either failing to converge due to complete separation (1me4:: error: Downdated VtV is not positive definite, see https://github.com/Ime4/Ime4/Issues/483).

```
possibly_parameters <- possibly(parameters::parameters, otherwise = NA)</pre>
poss_extract_fit_engine <- possibly(extract_fit_engine, otherwise = NA)</pre>
# ---- Create DF for combinatorial model specification ----
model formulas multivar <-
  tidyr::expand_grid(outcome = "box_cox_abs_deviation_score_estimate",
                     random_intercepts = list("study_id",
                                               "reviewer_id",
                                               c("study_id",
                                                 "reviewer_id")),
                     fixed_effects = list(c("publishable_as_is",
                                             "rate_analysis",
                                             "mean diversity index",
                                             "mixed_model"),
                                           c("publishable_as_is",
                                             "rate analysis",
                                             "mean_diversity_index"))) %>%
  rowwise() %>%
  mutate(dataset = case when(length(fixed_effects) == 4 ~ "eucalyptus",
                              TRUE ~ "blue tit"),
         wflow_id = paste0("RE:",
                           paste0(random intercepts, collapse = " "))) %>%
  unite(wflow_id, dataset, wflow_id, remove = FALSE) %>%
  rowwise() %>%
  mutate(model formulas =
           list(create_model_formulas(outcome,
```

```
fixed effects,
                                      random intercepts)) %>%
           set_names(wflow_id),
         model_workflows = list(create_model_workflow(outcome,
                                                      fixed_effects,
                                                       random_intercepts)) %>%
           set names(wflow id))
all model fits multivar <-
  ManyEcoEvo yi results %>%
  select(dataset, estimate type, effects analysis) %>%
  group by(dataset, estimate type) %>%
  nest_join(model_formulas_multivar %>%
              select(dataset,
                     model workflows,
                     fixed effects,
                     random intercepts),
            by = join by(dataset),
            name = "model_workflow_sets") %>%
  unnest(model_workflow_sets) %>%
  rowwise() %>%
  mutate(effects analysis =
           list(effects analysis %>%
                  select(study id,
                         starts_with("box_cox_abs_dev"),
                         RateAnalysis,
                         PublishableAsIs,
                         ReviewerId,
                         box cox var,
                         mean diversity_index,
                         mixed model) %>%
                  janitor::clean names() %>%
                  mutate if(is.character, factor)),
         fitted_mod_workflow = list(poss_fit(model_workflows,
                                             effects analysis)),
         fitted_model = list(poss_extract_fit_engine(fitted_mod_workflow)),
         convergence = list(if (!is.na(fitted_model))
           possibly check convergence(fitted_model)),
         singularity = list(if (!is.na(fitted model))
           possibly check singularity(fitted model)),
         params = list(if (!is.na(fitted model))
           possibly_parameters(fitted_model)),
        fixed_effects = paste0(fixed_effects, collapse = ", ")
  ) %>%
  unnest wider(random intercepts, names sep = " ") %>%
  unnest(c(convergence, singularity)) %>%
  rowwise() %>%
  replace na(list(convergence = FALSE)) %>%
  select(-model_workflows, -fitted_mod_workflow, -effects_analysis)
```

```
yi multivar singularity convergence <-
  all model fits multivar %>%
  left_join({all_model_fits_multivar %>%
      unnest(params) %>%
      filter(Effects == "random") %>%
      filter(is.na(SE) | is.infinite(SE)) %>%
      distinct(fixed effects,
               random intercepts 1,
               random_intercepts_2,
               dataset,
               estimate_type) %>%
      mutate(SE calc = FALSE)},
      by = join_by(dataset,
                   estimate_type,
                   random_intercepts_1,
                   random_intercepts_2,
                   fixed_effects)) %>%
  left_join({all_model_fits multivar %>%
      unnest(params) %>%
      filter(Effects == "random") %>%
      filter(if any(contains("CI"),
                    list(is.infinite, is.na))) %>%
      distinct(fixed_effects,
               random intercepts 1,
               random_intercepts_2,
               dataset,
               estimate type) %>%
      mutate(CI_calc = FALSE)},
      by = join_by(dataset,
                   estimate type,
                   random_intercepts_1,
                   random_intercepts_2,
                   fixed_effects)) %>%
  rowwise() %>%
  mutate(across(c(SE_calc, CI_calc), ~ ifelse(is.na(.x), TRUE, .x)),
         across(c(SE calc, CI calc, singularity),
                ~ ifelse(is_false(convergence), NA, .x)))
# If singularity == FALSE and convergence == TRUE,
# but the model appears here, then that's because
# the SD and CI's couldn't be estimated by parameters::
yi_multivar_singularity_convergence %>%
  select(-fixed effects, -fitted model, -params) %>%
  arrange(random_intercepts_1,
          random_intercepts_2,
          dataset,
          estimate_type) %>%
  mutate(across(starts_with("random"),
                ~ str_replace_all(.x, "_", " ") %>%
```

```
Hmisc::capitalize() %>%
                  str_replace("id", "ID")),
         dataset = str_replace(dataset, "eucalyptus", "*Eucalyptus*")) %>%
  group by(dataset) %>%
  gt::gt(rowname_col = "estimate_type") %>%
  tab style(
    style = list(
      cell fill(color = scales::alpha("red", 0.6)),
      cell_text(color = "white", weight = "bold")
    ),
    locations = list(
      cells_body(columns = "singularity", rows = singularity == TRUE),
      cells_body(columns = "convergence", rows = convergence == FALSE
                                                                            ce
lls_body(columns = "SE_calc", rows = SE_calc == FALSE),
      cells_body(columns = "CI_calc", rows = CI_calc == FALSE)
    )
  ) %>%
  gt::text transform(fn = function(x) ifelse(x == TRUE, "yes",
                                             ifelse(x == FALSE, "no", x)),
                     locations = cells body(columns = c("singularity",
                                                        "convergence",
                                                        "SE_calc",
                                                        "CI calc"))) %>%
  gt::opt stylize(style = 6, color = "gray") %>%
  gt::cols label(dataset = "Dataset",
                 singularity = "Singular Fit?",
                 convergence = "Model converged?",
                 SE_calc = gt::md("Can random effects $\\text{SE}$ be calcula
ted?"),
                 CI calc = "Can random effect CI be calculated?"
                 ) %>%
  gt::tab_spanner(label = "Random Effects",
                  columns = gt::starts with("random")) %>%
  gt::sub missing() %>%
  gt::cols label with(columns = gt::starts with("random"),
                      fn = function(x) paste0("")) %>%
  gt::text_transform(fn = function(x) map(x, gt::md),
                     locations = cells_row_groups()) %>%
 gt_fmt_yi(columns = "estimate_type")
```

Table C.8: Singularity and convergence checks for all combinations of random effects specifications trialled for across subsets of out of sample predictions y_i from multivariate models.

	Randor	n Effects	Model converged?	Singular Fit?	Can random effects SE be calculated?	Can random effect CI be calculated?
blue tit						
${y}_{25}$	Reviewer ID	_	yes	yes	no	no
${y}_{50}$	Reviewer ID	_	yes	no	yes	yes
y_{75}	Reviewer ID	_	yes	no	yes	yes
${y}_{25}$	Study ID	Reviewer ID	yes	yes	no	no
${y}_{50}$	Study ID	Reviewer ID	yes	no	no	no
y_{75}	Study ID	Reviewer ID	yes	yes	no	no
${y}_{25}$	Study ID	_	yes	no	no	no
${y}_{50}$	Study ID	_	yes	no	no	no

Eucalyptus						
${y}_{25}$	Reviewer ID	_	yes	yes	no	no
${y}_{50}$	Reviewer ID	_	yes	no	yes	yes
y_{75}	Reviewer ID	-	yes	no	yes	yes
${y}_{25}$	Study ID	Reviewer ID	yes	no	no	no
${y}_{50}$	Study ID	Reviewer ID	yes	yes	no	no
${y}_{25}$	Study ID	-	yes	no	no	no
${y}_{50}$	Study ID	_	yes	no	no	no

Consequently, we deviated from our intended plan of using random effects for both Effect ID and Reviewer ID, instead using a single random effect for Reviewer ID for the y_{50} and y_{75} prediction scenarios for both blue tit and *Eucalyptus* datasets (Table C.10, Table C.9).

```
yi_multivar_singularity_convergence %>%
  filter(SE calc == TRUE) %>%
  filter(random_intercepts_1 != "study_id" | dataset != "blue tit") %>% #rm e
liminated modl
  select(dataset, estimate_type, params) %>%
  unnest(params) %>%
  relocate(c(Effects, Group), .after = Parameter) %>%
  gt::gt(rowname_col = "estimate_type", groupname_col = "dataset") %>%
  gt::fmt_number(columns = c(-dataset, -estimate_type),
                 decimals = 2,
                 drop_trailing_zeros = TRUE,
                 drop trailing dec mark = TRUE
  ) %>%
  gt::text_transform(fn = function(x) str_replace(x, "publishable_as_is", "Ca
tegorical Peer Rating") %>%
                       str_replace(., "rate_analysis", "Continuous Peer Ratin
g") %>%
                       str_replace(., "mean_diversity_index", "Sorensen's Ind
```

```
ex") %>%
                       str replace(., "mixed model", "Random Included"),
                     locations = cells_body(columns = c("Parameter"))) %>%
  gt::opt_stylize(style = 6, color = "gray") %>%
  gt::sub_missing(missing_text = "") %>%
  gt::fmt(columns = "p",
          fns = function(x) gtsummary::style pvalue(x)) %>%
  gt::text_transform(
    locations = cells stub(
      rows = Parameter != "(Intercept)"
    ),
   fn = function(x)
     paste0("")
    }
  ) %>%
  gt::text_transform(locations =
                       cells body(columns = Group,
                                  rows = Group %in% c("reviewer id", "study i
d")),
                     fn = function(x){
   str_replace(x, "_", " ") %>%
                         Hmisc::capitalize() %>%
                         str replace("id", "ID")
                     }) %>%
 gt::cols label(CI low = gt::md("95\\%CI"), df error = "df", p = gt::md("*p*
"), SE = gt::md("$\\text{SE}$")) %>%
  gt::tab style(style = cell text(style = "italic", transform = "capitalize")
ر
                locations = cells_row_groups(groups = "eucalyptus")) %>%
  gt fmt yi(columns = "estimate type") %>%
  fmt_number(columns = c(gt::contains("CI"), "SE", "t"),
             drop_trailing_zeros = TRUE,
             drop trailing dec mark = TRUE,
             decimals = 2) %>%
  gt::fmt scientific(columns = c("Coefficient"),
                     rows = abs(Coefficient) < 0.01 | abs(Coefficient) > 1000
                     decimals = 2) %>%
  gt::fmt_scientific(columns = c("SE"),
                     rows = abs(SE) < 0.01 | abs(SE) > 1000,
                     decimals = 2) %>%
  gt::fmt scientific(columns = t,
                     rows = abs(t) < 0.01,
                     decimals = 2) %>%
  gt::fmt_scientific(columns = CI_low,
                     rows = abs(CI low) < 0.01 | abs(CI low) > 1000) %>%
  gt::fmt_scientific(columns = CI_high,
                     rows = abs(CI high) < 0.01 | abs(CI high) > 1000,
                     decimals = 2) %>%
  gt::cols_hide(Effects) %>%
```

Table C.9: Parameter estimates for converging, non-singular multivariate models fitted to blue tit out-of-sample-prediction
estimates y _i .

,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,	Parameter	Random Effect	Coefficient	${ m SE}$	95%Cl	t	df	p
olue tit								
y_{50}	(Intercept)		-0.42	0.49	[-1.39,0.56]	-0.84	216	0.4
	Categorical Peer Ratingpublishable as is		-0.59	0.43	[-1.44,0.26]	-1.37	216	0.2
	Categorical Peer Ratingpublishable with major revision		-0.27	0.33	[-0.92,0.38]	-0.81	216	0.4
	Categorical Peer Ratingpublishable with minor revision		-0.4	0.38	[-1.16,0.35]	-1.05	216	0.3
	Continuous Peer Rating		3.62×10^{-3}	4.94×10^{-3}	[-6.13 × 10 ⁻³ ,0.01]	0.73	216	0.5
	Sorensen's Index		-1.69	0.66	[-2.98,-0.39]	-2.57	216	0.011
	SD (Intercept)	Reviewer ID	0.2	0.09	[0.08,0.49]			
	SD (Observations)	Residual	0.71	0.04	[0.64,0.79]			

y_{75}	(Intercept)		-1.57	0.48	[-2.51,-0.63]	-3.3	229	0.001
	Categorical Peer Ratingpublishable as is		0.41	0.41	[-0.4,1.22]	1.01	229	0.3
	Categorical Peer Ratingpublishable with major revision		0.1	0.31	[-0.51,0.71]	0.32	229	0.7
	Categorical Peer Ratingpublishable with minor revision		0.37	0.37	[-0.36,1.1]	1	229	0.3
	Continuous Peer Rating		-4.10×10^{-4}	4.70×10^{-3} [-	-9.66 × 10 ⁻³ ,8.84 × 10 ⁻³]	-0.09	229	>0.9
	Sorensen's Index		0.13	0.63	[-1.12,1.38]	0.21	229	0.8
	SD (Intercept)	Reviewer ID	0.29	0.07	[0.18,0.45]			
	SD (Observations)	Residual	0.64	0.03	[0.57,0.71]			

calyptus								
y_{50}	(Intercept)		-2.08	1.21	[-4.47,0.31]	-1.72	162	0.087
	Categorical Peer Ratingpublishable as is		-0.05	0.86	[-1.75,1.66]	-0.06	162	>0.9
	Categorical Peer Ratingpublishable with major revision		0.12	0.58	[-1.03,1.26]	0.2	162	0.8
	Categorical Peer Ratingpublishable with minor revision		0.16	0.67	[-1.16,1.48]	0.24	162	0.8
	Continuous Peer Rating		4.04×10^{-4}	0.01	[-0.02,0.02]	0.04	162	>0.9
	Sorensen's Index		1.99	1.58	[-1.13,5.1]	1.26	162	0.2
	Random Included	I	0.31	0.26	[-0.21,0.83]	1.19	162	0.2
	SD (Intercept)	Reviewer ID	0.09	0.61	[1.68 × 10 ⁻⁷ ,4.89 × 10 ⁴]			
	SD (Observations)	Residual	1.3	0.08	[1.15,1.46]			

y_{75}	(Intercept)		-0.54	1.01	[-2.53,1.44]	-0.54	161	0.6
	Categorical Peer Ratingpublishable as is		0.19	0.81	[-1.42,1.79]	0.23	161	0.8
	Categorical Peer Ratingpublishable with major revision		-0.01	0.55	[-1.11,1.08]	-0.02	161	>0.9
	Categorical Peer Ratingpublishable with minor revision		0.17	0.63	[-1.08,1.41]	0.26	161	0.8
	Continuous Peer Rating		-1.91 × 10 ⁻³	9.63 × 10 ⁻³	[-0.02,0.02]	-0.2	161	0.8
	Sorensen's Index		0.31	1.31	[-2.27,2.9]	0.24	161	0.8
	Random Included	1	-0.11	0.25	[-0.6,0.38]	-0.46	161	0.6
	SD (Intercept)	Reviewer ID	0.04	1.3 [1.	$69 \times 10^{-30}, 9.00 \times 10^{26}]$			
	SD (Observations)	Residual	1.26	0.08	[1.11,1.42]			

```
ManyEcoEvo vi viz %>%
  filter(model name == "MA mod mv") %>%
  rowwise() %>%
  mutate(converged =
           possibly_check_convergence(model),
         singularity = possibly_check_singularity(model)) %>%
  select(dataset, estimate_type, mod_fit_stats, mod_glance) %>%
  hoist(mod_fit_stats, "RMSE", "Sigma", "R2_conditional", "R2_marginal", "ICC
") %>%
  hoist(mod_glance, "nobs") %>%
  select(-mod_glance, -mod_fit_stats) %>%
  semi join({ManyEcoEvo yi viz %>%
      filter(model_name == "MA_mod_mv") %>%
      rowwise() %>%
      transmute(dataset,
                estimate type,
                converged = possibly_check_convergence(model),
                singularity = possibly check singularity(model)) %>%
      filter(converged, !singularity)},
      by = join_by(dataset, estimate_type)) %>%
  relocate(nobs, .after = "ICC") %>%
  gt::gt(groupname_col = "dataset", rowname_col = "estimate_type") %>%
  gt::opt stylize(style = 6, color = "gray") %>%
  gt::cols_label(estimate_type = "Prediction Scenario",
                 R2 conditional = gt::md("$$R^{2} \\text{Conditional}$$"),
                 R2_marginal = gt::md("$$R^{2}_\\text{Marginal}$$"),
                 Sigma = gt::md("$$\\sigma$$"),
                 dataset = "Dataset",
                 nobs = gt::md("$N_{Obs}$")) %>%
 gt::tab_style(locations = cells_body(rows = str_detect(dataset, "Eucalyptus
"),
                                       columns = dataset),
                style = cell text(style = "italic")) %>%
  gt::cols hide(dataset) %>%
  gt_fmt_yi(columns = "estimate_type") %>%
  gt::fmt number(columns = c(gt::starts with("R2"), "ICC", "Sigma", "RMSE"),
                 drop_trailing_zeros = TRUE,
                 drop_trailing_dec_mark = TRUE,
                 decimals = 2) %>%
  gt::fmt_scientific(columns = c("RMSE"),
                     rows = abs(RMSE) < 0.01 | abs(RMSE) > 1000,
                     decimals = 2) %>%
  gt::fmt_scientific(columns = c("Sigma"),
                     rows = abs(Sigma) < 0.01 | abs(Sigma) > 1000,
                     decimals = 2) %>%
  gt::tab_style(style = cell_text(style = "italic", transform = "capitalize")
                locations = cells row groups(groups = "eucalyptus"))
```

			$R^2_{ m Conditional}$	$R^2_{ m Marginal}$		
	RMSE	σ	Conditional	wiarginai	ICC	N_{Obs}
blue tit						
y_{50}	0.68	0.71	0.11	0.05	0.07	224
y_{75}	0.59	0.64	0.2	0.03	0.17	237
Eucalyptus						
y_{50}	1.27	1.3	0.02	0.02	0	171
y_{75}	1.23	1.26	0.01	0.01	0	170

Table C.10: Model summary statistics for non-singular, converging multivariate models fit to
out-of-sample estimates y _i .

C.7 Model Summary Metrics for out-of-sample predictions y_i

```
tbl_data_yi_deviation_model_params %>%
  gt::gt(rowname col = "dataset") %>%
  gt::opt_stylize(style = 6, color = "gray") %>%
  gt::sub_missing(missing_text = "") %>%
  gt::cols_label(dataset = "Dataset",
                 R2 = gt::md("\$\$R^2\$\$"),
                 R2_conditional = gt::md("$$R^{2}_\\text{Conditional}$$"),
                 R2_marginal = gt::md("$$R^{2}_\\text{Marginal}$$"),
                 Sigma = gt::md("$$\\sigma$$"),
                 nobs = gt::md("$$N_{\\text{Obs.}}$$"),
                 estimate_type = "Prediction Scenario") %>%
 gt::tab_style(locations = cells_body(rows = str_detect(dataset, "Eucalyptus
"),
                                       columns = dataset),
                style = cell_text(style = "italic")) %>%
  gt::text_transform(
    locations = cells_stub(
     rows = estimate_type != "y25"
    ),
    fn = function(x)
     paste0("")
```

```
) %>%
  gt::tab style(locations = gt::cells stub(rows = str detect(dataset, "Eucaly
ptus")),
                style = cell text(style = "italic")) %>%
  gt::fmt_number(columns = gt::contains(c("ICC", "RMSE")),
                 drop_trailing_dec_mark = TRUE,
                 drop trailing zeros = T,
                 decimals = 2) %>%
  gt::fmt_scientific(columns = c("RMSE"),
                     rows = abs(RMSE) < 0.01 | abs(RMSE) > 1000,
                     drop_trailing_dec_mark = TRUE,
                     drop trailing zeros = T,
                     decimals = 2) %>%
  gt::fmt_scientific(columns = c(gt::contains(("R2_marginal"))),
                     rows = str_detect(model_name, "continuous|categorical"),
                     drop_trailing_zeros = T,
                     decimals = 2) %>%
  gt::fmt_number(columns = gt::contains(c("R2_conditional")),
                 drop trailing dec mark = TRUE,
                 drop_trailing_zeros = T,
                 decimals = 2) %>%
  gt::fmt_number(columns = gt::contains(c("Sigma")),
                 drop_trailing_dec_mark = TRUE,
                 drop_trailing_zeros = T,
                 decimals = 2) %>%
  gt::fmt_scientific(columns = c("Sigma"),
                     rows = abs(Sigma) < 0.01 | abs(Sigma) > 1000,
                     drop_trailing_zeros = T,
                     decimals = 2) %>%
  gt::fmt number(columns = "R2",
                 decimals = 2) %>%
  gt::fmt_scientific(columns = c("R2"),
                     rows = abs(R2) < 0.01 | abs(R2) > 1000,
                     drop_trailing_zeros = T,
                     decimals = 2) %>%
  gt::fmt scientific(columns = c("R2 conditional"),
                     rows = abs(R2_conditional) < 0.01 | abs(R2_conditional)</pre>
> 1000,
                     drop_trailing_zeros = T,
                     decimals = 2) %>%
  gt::fmt_scientific(columns = c("R2_marginal"),
                     rows = abs(R2 marginal) < 0.01 | abs(R2 marginal) > 1000
                     drop_trailing_zeros = T,
                     decimals = 2) %>%
  gt::fmt_scientific(columns = c("ICC"),
                     rows = abs(ICC) < 0.01 | abs(ICC) > 1000,
                     drop trailing zeros = T,
                     decimals = 2) %>%
gt_fmt_yi("estimate_type")
```

Table C.11: Model summary metrics for models of Box-Cox transformed deviation from the mean y_i estimate as a function of categorical peer-review rating, continuous peer-review rating, and Sorensen's index for blue tit and Eucalyptus analyses, and also for the inclusion of random effects for Eucalyptus analyses. Coefficient of determination, R^2 , is reported for models of deviation as a function of Sorensen diversity scores and presence of random effects, while $R^2_{Conditional}$, $R^2_{Marginal}$ and the intra-class correlation (ICC) are reported for models of deviation as explained by peer-review ratings. For all models the residual standard deviation σ , root mean squared error (RMSE) were calculated. The number of observations ($N_{Obs.}$) is displayed for reference.

	Prediction Scenario	$N_{ m Obs.}$	RMSE	σ	R^2	$R^2_{ m Conditional}$	$R^2_{ m Marginal}$	ICC			
Deviation explained	Deviation explained by Sorensen's index										
blue tit	y_{25}	63	0.58	0.59	7.76 × 10 ⁻⁴						
	y_{50}	60	0.72	0.73	0.03						
	y_{75}	63	0.71	0.73	5.99 × 10 ⁻³						
Eucalyptus	y_{25}	38	0.91	0.94	1.13 × 10 ⁻³						
	y_{50}	38	1.29	1.33	4.82 × 10 ⁻³						
	y_{75}	38	1.26	1.29	4.32×10^{-4}						
Deviation explained	d by continuous ratin	gs									
blue tit	y_{25}	237	0.58	0.58			2.48×10^{-4}				
	y_{50}	224	0.68	0.71		0.1	1.74 × 10 ⁻³	0.1			
	y_{75}	237	0.59	0.63		0.2	1.86 × 10 ⁻²	0.18			
Eucalyptus	y_{25}	174	0.92	0.93		0.04	1.79 × 10 ⁻²	0.02			
	y_{50}	171	1.28	1.29			5.75×10^{-4}				
	y_{75}	170	1.23	1.24			1.44×10^{-4}				

Deviation explained	l by categorical ratings	;						
blue tit	y_{25}	237	0.58	0.58			4.35×10^{-3}	
	y_{50}	224	0.68	0.71		0.1	1.3 × 10 ^{−2}	0.09
	y_{75}	237	0.59	0.63		0.2	3.36 × 10 ^{−2}	0.17
Eucalyptus	y_{25}	174	0.89	0.92		0.07	1.1 × 10 ^{−2}	0.06
	y_{50}	171	1.27	1.29		0.01	3.39 × 10 ^{−3}	9.14 × 10 ⁻³
	y_{75}	170	1.23	1.25			1.95 × 10 ^{−3}	
Deviation explained	by inclusion of rando	n effects						
Eucalyptus	y_{25}	38	0.85	0.87	0.14			
	y_{50}	38	1.29	1.33	3.91 × 10 ^{−3}			
	y_{75}	38	1.26	1.29	4.76 × 10 ⁻³			
C.8 Post-hoc analysis: checking the use of model weights in all models explaining deviation from the meta-analytic mean

As we describe in Note 3.1, for models of deviation from the meta-analytic mean effect-size, we had intended to use the inverse variance of the Box-Cox transformed deviation from the meta-analytic mean as model weights. Unfortunately using our intended weights specification resulted in invalid transformed response variables for some models whereby extreme outliers were weighted more heavily (two orders of magnitude) than other effect sizes, which caused both issues in the estimated model parameters as well as convergence issues, in particular for models analysing the effect of categorical peer-review rating on deviation from the meta-analytic mean.

(i) Model Weight Calculation Details

We intended to use the invariance of the Box-Cox transformed deviation scores as model weights in our models of deviation from the meta-analytic mean. The variance of the Box-Cox transformation scores is calculated using the delta method (Equation C.1).

$$\mu_{\text{folded}} = \sigma \sqrt{\frac{2}{\pi}} \exp\left(-\frac{\mu^2}{2\sigma^2}\right) + \mu \left(1 - 2 \times \boldsymbol{\Phi}\left(-\frac{\mu}{\sigma}; 0, 1\right)\right)$$

$$SE_{\text{folded}} = \sqrt{\mu^2 + \sigma^2 - \mu_{\text{folded}}^2}$$

$$VAR_{\text{folded}} = SE_{\text{folded}}^2$$

$$VAR_{\text{Box-Cox}} = VAR_{\text{folded}} \times \left(\lambda \mu_{\text{folded}}^{\lambda - 1}\right)^2$$

$$(C.1)$$

Where:

- μ_{folded} is the folded mean of the deviation scores from the mean for effect *i*,
- VAR_{folded} is the folded variance of the deviation scores from the mean for effect *i*,
- μ is the deviation score from the mean for effect *i*, calculated as the difference between effect size *i* and the mean for all effects: $\overline{Z}_r Z_{r_i}$,
- σ is the square root of the variance of effect *i*, VZ_r,
- λ is the Box-Cox transformation parameter (Figure C.1), and
- ϕ is the standard normal cumulative distribution function.

Which is executed in the function from the the package, illustrated in the following code snippet:

```
variance_box_cox <- function(folded_mu, folded_v, lambda){
  variance_bc <- folded_v * (lambda * folded_mu^(lambda - 1))^2 # delta
method
  return(variance_bc)
}
folded_params <- function(abs_dev_score, VZr){
  mu <- abs_dev_score
  sigma <- sqrt(VZr)
  fold_mu <- sigma * sqrt(2/pi) * exp((-mu^2)/(2 * sigma^2)) +
    mu * (1 - 2 * pnorm(-mu/sigma)) # folded abs_dev_score
  fold_se <- sqrt(mu^2 + sigma^2 - fold_mu^2)
  fold_v <- fold_se^2 # folded VZr
  return(list(fold_mu = fold_mu, fold_v = fold_v))
}</pre>
```

We systematically investigated the impact of using different weighting schemes (no weights, inverse-Box-Cox transformed variance-, and inverse folded variance-, of the absolute deviation

scores) on model convergence, singularity and other model checking metrics to aid decisionmaking about the appropriate weighting scheme and random-effects structure for these models. Given the convergence issues we encountered when using the intended weights, and that the desired random effects structure could not be fitted, we also investigated the impact of using different random-effects structures (effect ID, reviewer ID, or both effect ID and reviewer ID) on model convergence and singularity. For each weighting scheme, we fitted models with each different random effects structure to both the blue tit and *Eucalyptus* analyst data, and evaluated model convergence, singularity and model performance using the performance::compare_performance function.

```
# Create filter argument expressions
filter_args = rlang::exprs(exclusion_set == "complete",
                            publishable subset == "All",
                            expertise_subset == "All",
                            collinearity_subset == "All")
# Create function to prepare ratings data
prepare_ratings_data <- function(effects_analysis){</pre>
  data tbl <-
    effects analysis %>%
    unnest(cols = c(review data)) %>%
    select(study_id,
           TeamIdentifier,
           starts_with("box_cox_"),
           ReviewerId,
           PublishableAsIs,
           # lambda,
           folded_v_val) %>%
    ungroup() %>%
    mutate(PublishableAsIs =
             forcats::fct_relevel(PublishableAsIs,
                                   c("deeply flawed and unpublishable",
                                      "publishable with major revision",
                                      "publishable with minor revision",
                                     "publishable as is")),
           obs_id = 1:n())
  return(data_tbl)
}
# Create base model formulat
base_formula <- rlang::new_formula(</pre>
  rlang::expr(box cox abs deviation score estimate),
  rlang::expr(PublishableAsIs))
# Create weight functions
calc inv bc var <- rlang::as function(~ 1/pull(.x, box cox var))</pre>
calc inv folded v <- rlang::as function(~ 1/pull(.x, folded v val))</pre>
no_weights <- NA</pre>
```

```
weight formulas <- list(no weights,</pre>
                         calc inv bc var,
                         calc_inv_folded_v
) %>%
  purrr::set_names("no_weights",
                    "inv bc_var",
                    "inv folded v")
# Create random effect expressions
RE_rev <- expr((1 | ReviewerId))</pre>
RE_study <- expr((1 | study_id))</pre>
RE both <- expr(!!RE rev + !!RE study)</pre>
random expressions <- list(</pre>
  RE_rev,
  RE_study,
  RE both
) %>% purrr::set_names("RE_rev",
                        "RE_study",
                        "RE_both")
# Create model fitting wrapper function
lmer wrap <- function(data tbl,</pre>
                       random effect,
                       weight_form,
                       . . . ,
                       env = caller env()){
 f <- rlang::new_formula(expr(box_cox_abs_deviation_score_estimate),</pre>
                           expr(PublishableAsIs + !!random effect),
                           env = env)
  weights <- if ( rlang::is_na(weight_form) ) NULL else weight_form(data_tbl)</pre>
  rlang::inject(lme4::lmer(!!f,
                            data = data_tbl,
                            weights = weights,
                             ...))
}
# Fit all models
all models <-
  ManyEcoEvo_results %>%
  ungroup %>%
 filter(!!!filter_args) %>%
  select(dataset, effects_analysis) %>%
  hoist(.col = effects_analysis,
        "lambda",
        .simplify = TRUE,
        .transform = unique) %>%
  mutate(model data = map(effects analysis,
```

```
prepare ratings data),
         .keep = "unused") %>%
  expand grid(
    expand grid(weight formulas, random expressions) %>%
      mutate(weight_forms = names(weight_formulas),
             random_effect = names(random_expressions)) %>%
      unite("model spec", weight forms, random effect, sep = ".")
  ) %>%
  # hoist("model_data", weights = list("study_id", "box_cox_var", "folded_v_v
al"),.remove = F) %>%
  mutate(model = pmap(list(model_data,
                           random expressions,
                           weight_formulas),
                      lmer wrap),
         .keep = "unused") %>%
  mutate(singularity = map_lgl(model,
                               performance::check singularity),
         convergence = map lgl(model,
                               performance::check convergence))
possibly estimate means <- possibly(modelbased::estimate means, otherwise = N</pre>
ULL)
# Extract Parameter Estimates
estimate means <-
  all models %>%
  filter(singularity == F, convergence == T) %>%
  reframe(model = set names(model, model spec),
          dataset = dataset,
          model_spec = model_spec,
          weights = case when(!str detect(model spec, "no weights") ~ "(weigh
ts)",
                                   .default = NA)) %>%
  rowwise() %>%
  mutate(weights = modify_if(list(weights), ~ is.na(.x), ~ NULL),
         results = list(possibly estimate means(model,
                        by = "PublishableAsIs", weights = weights))) %>%
  ungroup() %>%
  mutate(results = set_names(results, dataset)) %>%
  drop_na(results) # model means couldn't be estimated due to convergence iss
ues, drop those models
# evaluate and compare performance for remaining models
model_comparison_results <-</pre>
  all models %>%
 filter(model_spec != "inv_bc_var.RE_study" | dataset != "eucalyptus") %>% #
rm nearly unidentifiable model
  semi_join(estimate_means,
            by = join_by(dataset, model_spec)) %>%
```

```
group by(dataset) %>%
  summarise(model = set names(model, model spec) %>% list,
            results = map(model,
                          performance::compare performance,
                          rank = T),
            results = set_names(results,
                                unique(dataset)),
            .groups = "keep")
model means results <-
  estimate means %>%
  left join(model comparison results %>%
              select(-model) %>% unnest(results),
            by = join_by("dataset", "model_spec" == "Name")) %>%
  mutate(label = paste(dataset, model spec, sep = ".")) %>%
  arrange(dataset, desc(Performance Score)) %>%
  select(Performance_Score, dataset, model_spec, results) %>%
  mutate(label = paste(dataset, model_spec, sep = "."))
```

C.8.1 Model Weight Investigation Findings

For the blue tit models of deviation influenced by categorical peer-review rating, no models including study ID as a random-effect were able to be properly fitted, across all model weight specifications Table C.12. For *Eucalyptus* models with either no weights, or inverse folded variance as weights, only models with Reviewer ID as the random-effect fitted properly. While models with either Reviewer ID or Study ID as the random-effect passed singularity and convergence fit-checks, and had estimable parameter means when the Inverse Box-Cox variance was used as a model weight Table C.12.

```
all models %>%
  select(dataset, model_spec, singularity, convergence) %>%
  left_join(estimate_means %>% select(-model, -results, -weights) %>%
              mutate(estimate means = T)) %>%
  separate(model_spec, into = c("model_spec_weight",
                                "model spec random effect"), sep = "\\.") %>%
  replace_na(replace = list(estimate_means = FALSE)) %>%
  group_by(dataset) %>%
  gt::gt(rowname_col = "model_spec_weight") %>% #
  gt::opt_stylize(style = 6, color = "gray") %>%
  gt::tab_stubhead(label = "Model Weight") %>%
  gt::tab style(
    style = list(
      cell_fill(color = scales::alpha("red", 0.6)),
      cell_text(color = "white", weight = "bold")
    ),
    locations = list(
      cells_body(columns = "singularity", rows = singularity == TRUE),
      cells_body(columns = "convergence", rows = convergence == FALSE),
      cells_body(columns = "estimate_means", rows = estimate_means == FALSE)
```

```
)
  ) %>%
  gt::tab_style(style = cell_text(style = "italic", transform = "capitalize")
ر
                locations = cells_row_groups(groups = "eucalyptus")) %>%
  gt::cols_label(dataset = "Dataset",
                 singularity = "Singular Fit?",
                 convergence = "Model converged?",
                 model_spec_random_effect = "Random Effects",
                 estimate means = "Means Estimable?") %>%
  gt::text_transform(fn = function(x) ifelse(x == TRUE, "yes",
                                               ifelse(x == FALSE, "no", x)),
                     locations = cells_body(columns = c("singularity",
                                                          "convergence",
                                                          "estimate_means"))) %
>%
  gt::text_transform(
    locations = cells stub(
      rows = model spec random effect != "RE rev"
    ),
    fn = function(x)
    paste0("")
    }
  ) %>%
  gt::text transform(
    locations = cells_body(columns = "model_spec_random_effect"),
    fn = function(x)
      str_replace(x, "RE_rev", "Reviewer ID") %>%
str_replace("RE_study", "Study ID") %>%
        str replace("RE both", "Reviewer ID and Study ID")
    }
  ) %>%
  gt::text_transform(
    locations = cells stub(),
    fn = function(x){
      str_replace(x, "no_weights", "None") %>%
        str_replace("inv_bc_var", "Inverse Box-Cox variance") %>%
        str_replace("inv_folded_v", "Inverse folded variance")
    }
  ) %>%
  gt::tab_style(style = cell_text(style = "italic", transform = "capitalize"
),
                locations = cells row groups(groups = "eucalyptus"))
```

Table C.12: Singularity and convergence checks for all combinations of model weights and random-effects structure in models of the effect of categorical peer rating on deviation from the analytic mean. For some models, mean estimates of parameter levels for peer-review rating were not able to be estimated.

Model Weight	Random Effects	Singular Fit?	Model converged?	Means Estimable?
blue tit				
None	Reviewer ID	no	yes	yes
	Study ID	no	no	no
	Reviewer ID and Study ID	yes	yes	no
Inverse Box-Cox variance	Reviewer ID	no	yes	yes
	Study ID	no	yes	no
	Reviewer ID and Study ID	yes	yes	no
Inverse folded variance	Reviewer ID	no	yes	yes
	Study ID	no	yes	no
	Reviewer ID and Study ID	yes	yes	no
Eucalyptus				
None	Reviewer ID	no	yes	yes
	Study ID	no	no	no
	Reviewer ID and Study ID	no	yes	no
Inverse Box-Cox variance	Reviewer ID	no	yes	yes
	Study ID	no	yes	yes
	Reviewer ID and Study ID	yes	yes	no
Inverse folded variance	Reviewer ID	no	yes	yes
	Study ID	no	yes	no
	Reviewer ID and Study ID	yes	yes	no

To check that the alternative weighting methods generated sensible parameter estimates, we generated marginal effects plots for all models that passed the convergence and singularity checks where marginal effects were estimable for both blue tit Figure C.7 and *Eucalyptus* datasets Figure C.6.

Using the inverse Box-Cox transformed variance for model weights resulted in the marginal mean being pulled towards zero for both datasets, however, this was quite extreme for the *Eucalyptus* dataset Figure C.6A. When the random-effect for study ID is substituted in place of the reviewer ID, the model means seem more fitting to the data, but uncertainty of estimates seemed artificially small Figure C.6B. For this model, marginal means were all equal in both mean estimate and their standard error when the inverse Box-Cox variance was used as a weight with Study ID as the random effect because the model was nearly unidentifiable Table C.13, so we eliminated this specification from further consideration.

```
modify_plot <- function(p, .y){</pre>
  p +
   labs(subtitle = as label(.y),
         title = NULL,
         x = "",
         y = "Box-Cox transformed\nabsolute deviation from\nmeta-analytic mea
n") +
    see::theme_lucid() +
    theme(axis.text.x = element text(angle = 50, hjust = 1))
}
model_means_results %>%
  filter(dataset == "eucalyptus") %>%
  arrange(model spec) %>%
  pull(results, name = model_spec) %>%
  map(., plot, at = "PublishableAsIs") %>%
  map2(., names(.), modify_plot) %>%
  patchwork::wrap_plots() +
  patchwork::plot annotation(tag levels = 'A')
```



Figure C.6: Effect plots for each non-singular model that converged with estimable fixed effect group means for the Eucalyptus dataset.

```
modify_plot <- function(p, .y){
    p +
    labs(subtitle = as_label(.y),
        title = NULL,
        x = "",
        y = "Box-Cox transformed\nabsolute deviation from\nmeta-analytic mea
n") +
    see::theme_lucid() +
    theme(axis.text.x = element_text(angle = 50, hjust = 1))
}
model_means_results %>%
    filter(dataset == "blue tit") %>%
```

```
pull(results, name = model_spec) %>%
map(., plot, at = "PublishableAsIs") %>%
map2(., names(.), modify_plot) %>%
patchwork::wrap_plots() +
patchwork::plot_annotation(tag_levels = 'A')
```



Figure C.7: Effect plots for each non-singular model that converged with estimable fixed effect group means for the blue tit dataset.

```
paste0("")
  }
) %>%
gt::text_transform(
  locations = cells_body(columns = "model_spec_random effect"),
  fn = function(x)
    str_replace(x, "RE_rev", "Reviewer ID") %>%
      str_replace("RE_study", "Study ID") %>%
str_replace("RE_both", "Reviewer ID and Study ID")
  }
) %>%
gt::text transform(
  locations = cells stub(),
  fn = function(x){
    str_replace(x, "no_weights", "None") %>%
    str_replace("inv_bc_var", "Inverse Box-Cox variance") %>%
      str_replace("inv_folded_v", "Inverse folded variance")
  }
) %>%
gt::cols_label(model_spec_random_effect = "Random Effects",
                PublishableAsIs = "Peer Rating",
                CI_low = gt::md("95\\%CI"),
                SE = gt::md("$\\text{SE}$")) %>%
gt::cols_merge(columns = starts_with("CI_"),
                pattern = "[{1},{2}]") %>%
gt::fmt_number(columns = c("Mean", "SE", "CI_low", "CI_high"),
                decimals = 2) %>%
gt::text_transform(
  locations = cells body(
    columns = "model spec random effect",
    rows = PublishableAsIs != "deeply flawed and unpublishable"
  ),
  fn = function(x)
   paste0("")
  }
) %>%
gt::tab_style(
  style = list(gt::cell_text(transform = "capitalize"),
                gt::cell_text(style = "italic")),
  locations = gt::cells_row_groups(groups = "eucalyptus")
) %>%
gt::tab style(style = cell text(transform = "capitalize"),
              locations = cells body(columns = "PublishableAsIs"))
```

Table C.13: Marginal means estimate across weight and random effects specifications for all estimable models for both Eucalyptus and blue tit datasets.

Model Weight	Random Effects	Peer Rating	Mean	${ m SE}$	95%CI
blue tit					
None	Reviewer ID	Deeply Flawed And Unpublishable	-1.11	0.11	[-1.33,-0.89]
		Publishable With Major Revision	-1.30	0.05	[-1.39,-1.21]
		Publishable With Minor Revision	-1.30	0.04	[-1.38,-1.22]
		Publishable As Is	-1.24	0.07	[-1.38,-1.10]
Inverse folded variance	Reviewer ID	Deeply Flawed And Unpublishable	-1.40	0.14	[-1.67,-1.12]
		Publishable With Major Revision	-1.48	0.06	[-1.61,-1.35]
		Publishable With Minor Revision	-1.43	0.05	[-1.53,-1.32]
		Publishable As Is	-1.18	0.08	[-1.35,-1.01]
Inverse Box-Cox variance	Reviewer ID	Deeply Flawed And Unpublishable	-0.95	0.12	[-1.18,-0.72]
		Publishable With Major Revision	-1.15	0.06	[-1.26,-1.03]
		Publishable With Minor Revision	-1.05	0.05	[-1.15,-0.95]
		Publishable As Is	-0.72	0.07	[-0.85,-0.58]

Eucalyptus					
None	Reviewer ID	Deeply Flawed And Unpublishable	-2.66	0.27	[-3.19,-2.12]
		Publishable With Major Revision	-2.37	0.12	[-2.61,-2.12]
		Publishable With Minor Revision	-2.65	0.11	[-2.86,-2.43]
		Publishable As Is	-2.60	0.17	[-2.94,-2.27]
Inverse folded variance	Reviewer ID	Deeply Flawed And Unpublishable	-3.31	0.26	[-3.82,-2.81]
		Publishable With Major Revision	-2.97	0.13	[-3.21,-2.72]
		Publishable With Minor Revision	-3.02	0.11	[-3.23,-2.80]
		Publishable As Is	-3.08	0.16	[-3.40,-2.77]
Inverse Box-Cox variance	Reviewer ID	Deeply Flawed And Unpublishable	-0.51	0.39	[-1.27,0.24]
		Publishable With Major Revision	-1.00	0.30	[-1.59,-0.42]
		Publishable With Minor Revision	-2.22	0.30	[-2.81,-1.64]
		Publishable As Is	-2.76	0.34	[-3.42,-2.09]
Inverse Box-Cox variance	Study ID	Deeply Flawed And Unpublishable	-2.58	0.06	[-2.70,-2.45]
		Publishable With Major Revision	-2.58	0.06	[-2.70,-2.45]
		Publishable With Minor Revision	-2.58	0.06	[-2.70,-2.45]
		Publishable As Is	-2.58	0.06	[-2.70,-2.45]

After discarding models based on the above criteria, we were left with a subset of models that passed convergence and singularity checks, and had estimable parameter means, all of which included reviewer ID as the only random effect. Although it seemed that the inverse Box-Cox variance resulted skewed estimated marginal means, it was unclear whether the inverse folded variance was a better alternative over using no weights, as the inverse folded variance seemed to exhibit a similar pattern in pulling the estimated marginal effects towards zero, but to a lesser extent than for the inverse Box-Cox variance.

To further aid in decision-making about which model weights to use, we compared the performance of all models that passed the convergence and singularity checks, and had estimable parameter means. We used the performance::compare_performance function to calculate performance metrics for this subset of models, and plotted the results using the performance package's in-built plotting method, which creates spider plots of normalised performance metrics for each model Figure C.8. For completeness, all model performance results of this final subset are reported in Table C.14.

The performance comparison plots confirmed our suspicions that the inverse Box-Cox variance was not a suitable weight for these models, and it performed relatively poorly across all metrics for both the *Eucalyptus* and blue tit datasets Figure C.8. The inverse folded variance weighted models performed similarly poorly for both datasets across all metrics except for RMSE in the case of the blue tit dataset, and both Sigma and RMSE for the *Eucalyptus* dataset. For both datasets using no weights when there is only a random-effect for Reviewer ID resulted in the best model fit and performance across all metrics. In keeping with our informed preference of using no weights, model comparison analysis highlighted that using no weights in the models was preferable.

```
# plot performance for remaining models
model_comparison_plots <-</pre>
  model comparison results %>%
  mutate(dataset = str_replace(dataset,
                                "eucalyptus",
                                "*Eucalyptus*")) %>%
  pull(results, "dataset") %>%
  map(plot)
# for printing plot name on figure
# model comparison plots %>%
   map2(.x = ., .y = names(.), ~ .x + ggtitle(.y) #+
#
           theme(title = ggtext::element_markdown())
#
#
         )
model_comparison_plots
```



Comparison of Model Indices

(a) Blue tit models.



Comparison of Model Indices

(b) Eucalyptus models.

Figure C.8: Model performance comparison plots for a subset of models that passed convergence and singularity checks and had estimable marginal effects. Models are compared based on their performance metrics, including $R_{Conditional'}^2$, $R_{Marginal'}^2$, Intra-Class Correlation, Root Mean Squared Error (RMSE), and the weighted AIC, corrected AIC and BIC. Values of performance are normalised across models for each metric, against the top-most performing model across all metrics. Greater distance from the centre on each metric axis indicates dominance in performance. For both blue tit and Eucalyptus models, all included a random effect for Reviewer ID, and weights consisted of the inverse Box-Cox variance for weights (blue line), inverse folded variance (yellow line), or none (red line).

```
results = set names(results,
                                unique(dataset)),
            .groups = "keep") %>%
  unnest(results) %>%
  select(-Model, -model) %>%
  gt::gt() %>%
  gt::cols label(Name = "Model Weight",
                 R2_conditional = gt::md("$$R^{2}_\\text{Conditional}$$"),
                 R2_marginal = gt::md("$$R^{2}_\\text{Marginal}$$"),
                 Sigma = gt::md("$$\\sigma$$"),
                 AICc = gt::md("$$AIC_c$$"),
                 AICc wt = gt::md("$$AIC c$$ (wt)"),
                 BIC_wt = gt::md("$$BIC$$ (wt)"),
                 AIC_wt = gt::md("$$AIC$$ (wt)"),
                 AIC = gt::md("$$AIC$$"),
                 BIC = gt::md("$$BIC$$")) %>%
  gt::fmt_number(columns = contains(c("AIC", "AICc", "BIC", "R2_", "ICC", "Sig
ma")),
                 drop trailing zeros = TRUE,
                 drop_trailing_dec_mark = TRUE,
                 decimals = 2) %>%
  gt::fmt_scientific(columns = "R2_conditional",
                     rows = R2_conditional < 0.01 | R2_conditional > 1000,
                     drop trailing zeros = T,
                     decimals = 2) %>%
  gt::fmt_scientific(columns = "R2_marginal",
                     rows = R2 marginal < 0.01 | R2 marginal > 1000,
                     drop_trailing_zeros = T,
                     decimals = 2) %>%
  gt::fmt scientific(columns = "RMSE",
                     rows = RMSE < 0.01 | RMSE > 1000,
                     drop_trailing_zeros = T,
                     decimals = 2) %>%
  gt::fmt_scientific(columns = c("ICC"),
                     rows = ICC < 0.01 | ICC > 1000,
                     drop trailing zeros = T,
                     decimals = 2) %>%
  gt::fmt_scientific(columns = c("Sigma"),
                     rows = Sigma < 0.01 | Sigma > 1000,
                     drop_trailing_zeros = T,
                     decimals = 2) %>%
  gt::fmt_scientific(columns = gt::contains(c("_wt")),
                     rows = Name != "no_weights.RE_rev",
                     drop_trailing_zeros = F,
                     decimals = 2) %>%
  gt::opt stylize(style = 6, color = "gray") %>%
  gt::tab_style(
    style = list(gt::cell text(transform = "capitalize"),
                 gt::cell_text(style = "italic")),
   locations = gt::cells_row_groups(groups = "eucalyptus")
```

```
) %>%
gt::fmt(
    columns = gt::contains("Name"),
    fns = function(x) str_remove(x, ".RE_rev") %>%
        str_replace("no_weights", "None") %>%
        str_replace("inv_bc_var", "Inverse Box-Cox variance") %>%
        str_replace("inv_folded_v", "Inverse folded variance")
)
```

Table C.14: Model performance metric values (non-normalised) for final subset of models considered in weights analysis. All models in final subset included random-effect of Reviewer ID. Metrics included $R_{Conditional}^2$, $R_{Marginal}^2$, Intra-Class Correlation, Root Mean Squared Error (RMSE), and the weighted AIC, corrected AIC and BIC.

Model Weight	AIC	AIC (wt)	AIC_c	$\int_{c} AIC_{c}$ (wt)		BIC (wt)	$R^2_{ m Conditional}$	$R^2_{ m Marginal}$	ICC	RMSE	σ
blue tit											
None	723.15	1	723.33	1	748.1	1	0.09	7.47 × 10 ⁻³	0.08	0.4827981	0.5
Inverse Box- Cox variance	840.52	3.26×10^{-26}	840.7	3.26 × 10 ⁻²⁶	865.48	3.26 × 10 ⁻²⁶	7.67 × 10 ⁻⁴	1.31 × 10 ⁻⁴	6.36 × 10 ⁻⁴	0.5822505	11.83
Inverse folded variance	1,026.75	1.19 × 10 ⁻⁶⁶	1,026.93	1.19 × 10 ⁻⁶⁶	1,051.7	1.19 × 10 ⁻⁶⁶	4.71 × 10 ⁻⁴	5.14 × 10 ⁻⁵	4.2×10^{-4}	0.5026704	13.19
Eucalyptus											
None	1,063.74	1	1,063.99	1	1,086.82	1	0.13	0.01	0.12	1.0173409	1.06
Inverse Box- Cox variance	1,386.88	6.80 × 10 ⁻⁷¹	1,387.12	6.80 × 10 ⁻⁷¹	1,409.96	6.80 × 10 ⁻⁷¹	2.28 × 10 ⁻⁹	2.47 × 10 ⁻¹⁰	2.03 × 10 ⁻⁹	1.4993872	4.68 × 10 ⁴
Inverse folded variance	1,109.35	1.25 × 10 ^{−10}	1,109.6	1.25 × 10 ^{−10}	1,132.43	1.25 × 10 ⁻¹⁰	5.66 × 10 ⁻⁴	1.82 × 10 ⁻⁵	5.48 × 10 ⁻⁴	1.1266586	19.68

SM D: Correlation Matrices of Case Study Data

```
library(tidyverse)
library(ManyEcoEvo)
library(GGally)
set.seed(1234)
source(here::here("utils.R"))
```

Pairwise-correlation plots for the *Eucalyptus* and blue tit case-study data provided to analysts are shown in Figure D.1 and Figure D.2, respectively. Plots were created with R package GGally (Schloerke et al. 2024).

```
euc_data %>%
select(where(is_double),
        -Date,
        -`Quadrat no`,
        -Easting,
        -Northing,
        -`small*0.25+medium*1.25+large*2.5`,
        -`average.proportion.of.plots.containing.at.least.one.euc.seedling.o
f.any.size`) %>%
GGally::ggpairs()
```

0.06 0.04 0.02 -		Corr:	Corr:	Corr:	Corr:	Corr:	Corr:	Corr:	Corr:	Corr:	Corr:	Corr:	Corr:	Corr:	Corr:	Corr:	Corr:	Corr:	Corr:	Corr:	Corr:	Corr:	Corr:	Corr:	Corr:	Corr:	Corr:	SRad_Jul Corr: -0.104.	Corr:		
0.00 - 80 - 60 - 40 - 20 -		Corr: 0.183***	Corr: 0.121*		Corr: 0.103.		Corr: 0.056		Corr: -0.048		Corr: 0.365**		Corr: 0.088		Corr: 0.196***	Corr: 0.058	Corr: -0.134*	Corr: -0.058		Corr: 0.336***	Corr: 0.333***	Corr: 0.382**	Corr: 0.347**	Corr: 0.426***	Corr: -0.125*	Corr: 0.280***	Corr: -0.054	Corr: 0.080	Corr: -0.031	Corr: -0.022	Corr: -0.145** He
60 40 20 0	÷.		Corr: 0.060	Corr: 0.039	Corr: -0.001	Corr: 0.082	Corr: 0.177***		Corr: -0.000	Corr: -0.165**	Corr: 0.294**		Corr: -0.011	Corr: -0.030		Corr: -0.022		Corr: -0.046	Corr:).329***	Corr: 0.324***	Corr: 0.319***	Corr: 0.331**	Corr: 0.233**	Corr: 0.071	Corr: 0.015	Corr: 0.208***	Corr: -0.110*	Corr: -0.023	Corr: -0.070	Corr: -0.063	Corr: -0.105*
80 60 40 20	.			Corr: -0.024	Corr: -0.065		Corr: -0.001		Corr: -0.024	Corr: -0.045	Corr: 0.200**			Corr: -0.063	Corr: 0.173**			Corr: -0.004	Corr: 0.199***	Corr: 0.206***	Corr: 0.184***	Corr: 0.218**	Corr: -0.053	Corr: 0.019		Corr: 0.112*	Corr: 0.018	Corr: 0.029		Corr: -0.071	Corr: -0.096. G
3 2 1 0		. <u>.</u>		Ļ	Corr: -0.018	Corr: 0.043	Corr: 0.043		Corr: -0.010	Corr: -0.061	Corr: -0.014			-0.039	0.056		-0.033			Corr: 0.032	0.016		Corr: -0.030		0.026	Corr: -0.041				Corr: -0.026	3
10 5 80 6	<u>.</u>	<u>.</u>	<u>ŀ</u>	i		Corr: 0.066	Corr: 0.098.		Corr: -0.023		Corr: -0.073			Corr: 0.078	Corr: -0.050			Corr: -0.006		Corr: 0.021			Corr: -0.139**		Corr: 0.012					Corr: 0.097.	8
40 20 0 15	.		K	. :	a		Corr: -0.032				Corr: 0.311** Corr:		Corr: -0.046 Corr:			Corr: -0.047 Corr:		Corr: -0.074 Corr:	Corr: 0.027 Corr:			Corr: 0.008 Corr:	Corr: 0.188** Corr:	Corr: -0.017 Corr:		Corr: -0.017 Corr:	Corr: 0.071 Corr:	Corr: 0.031 Corr:	Corr: -0.011 Corr:	Corr: 0.000	Corr: nii) 0.046 Corr: en
10 - 5 - 5 - 5 - 5 - 5 - 5 - 5 - 5 - 5 -	.	. 1 1	.	! . :	.	.	<u> </u>				-0.080	0.110*		0.054	0.054		0.067				0.317***			0.092.	0.171**		-0.110*	-0.031 Corr:	0.030 Corr:	0.025	
10 - 5 0	.	· •	<u>.</u>	!	. حب	<u>ier.</u>	. .		-0.022	Corr:	Corr:	Corr:	Corr:	Corr:	Corr:	Corr:	Corr:	Corr:	Corr:	Corr:	Corr:	Corr:	Corr:	Corr:	-0.029 Corr:	0.061 Corr:	Corr:	0.091. Corr:	Corr:	Corr:	Corr:
50 100 75 50	Ē.	. <u>6-</u> •	Ľ	1 · ·		<u>».</u> k.	84 - k	ь L		1	0.028 Corr:	Corr:	Corr:	Corr:	Corr:	Corr:		Corr:	Corr:	0.011 Corr:	Corr:	-0.007 Corr:	-0.063 Corr:		Corr:	Corr:	Corr:	-0.010 Corr:	Corr:	Corr:	0.021 Gran
25 0 100 75 50			k.			the last		i	6.		\wedge	Corr:	Corr:	Corr:	Corr:	Corr:	Corr:	Corr:	Corr:	Corr:	Corr:	Corr:	Corr:	Corr:	Corr:	Corr:	Corr:	0.084 Corr: -0.048	Corr:	Corr:	Corr:
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Figure D.1: Pairwise correllation plot for all Eucalyptus dataset variables except for Date, Quadrat no, Easting, Northing.

```
blue tit data %>%
  naniar::replace_with_na_all(condition = ~ .x == ".") %>%
 mutate(across(c(contains("_ring"),
                  rear_nest_trt,
                  hatch_year,
                  hatch_nest_breed_ID,
                  hatch Area,
                  hatch_Box,
                  day14_measurer,
                  contains("hatch_Box"),
                  starts_with("rear_"),
                  starts with("hatch nest"),
                  home or away,
                  -rear_d0_rear_nest_brood_size,
                  contains("manipulation"),
                  chick_sex_molec,
                  Date_of_day14,
                  `Extra-pair paternity`,
                  -rear_Cs_in,
                  -rear_Cs_out,
                  chick_survival_to_first_breed_season,
                  -rear_Cs_at_start_of_rearing),
                as.factor),
         across(where(is.character), as.numeric),
         across(c(rear_Cs_out,
                  rear_Cs_in,
                  rear_Cs_at_start_of_rearing),
                as.integer)) %>%
  select(where(is.numeric), -`day 14 weight`) %>%
  GGally::ggpairs()
```



Figure D.2: Pairwise correlation plot of all numeric variables in blue tit case study dataset