Checklist item	Sub-item number	Sub-item	Reported by authors?	Notes
Title and abstract				
	1.1	Identify the review as a systematic review, meta-analysis, or	Y	
	1.0	both	V	
	1.2	Summarise the aims and scope of the review	Y	
	1.3	Describe the data set	Y	
	1.4	State the results of the primary outcome	Y	
	1.5	State conclusions	Y	
	1.6	State limitations	Υ	
Aims and questions				
	2.1	Provide a rationale for the review	Υ	
	2.2	Reference any previous reviews or meta-analyses on the topic	Υ	
	2.3	State the aims and scope of the review (including its generality)	Υ	
	2.4	State the primary questions the review addresses (e.g. which moderators were tested)	Υ	
	2.5	Describe whether effect sizes were derived from experimen-	Υ	
		tal and/or observational comparisons		
Review registration				
	3.1	Register review aims, hypotheses (if applicable), and methods in a time-stamped and publicly accessible archive and provide a link to the registration in the methods section of the	N	No registration done
		vide a link to the registration in the methods section of the manuscript. Ideally registration occurs before the search, but it can be done at any stage before data analysis.		
	3.2	Describe deviations from the registered aims and methods	N	No registration done
	3.3	Justify deviations from the registered aims and methods	N	No registration done
Eligibility criteria		,		
Lingibility Criteria	4.1	Report the specific criteria used for including or excluding	Υ	
	7.1	studies when screening titles and/or abstracts, and full texts,	•	
		according to the aims of the systematic review (e.g. study		
		design, taxa, data availability)		
	4.2	Justify criteria, if necessary (i.e. not obvious from aims and	Υ	
	4.2	scope)	1	
Finding studies		. ,		
	5.1	Define the type of search (e.g. comprehensive search, repre-	Υ	
		sentative sample)		
	5.2	State what sources of information were sought (e.g. published	Υ	
		and unpublished studies, personal communications)		
	5.3	Include, for each database searched, the exact search strings	Υ	
		used, with keyword combinations and Boolean operators		
	5.4	Provide enough information to repeat the equivalent search (if	Υ	
		possible), including the timespan covered (start and end dates)		
Study selection				
,				

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	6.1	Describe how studies were selected for inclusion at each	Υ	
	0.1	stage of the screening process (e.g. use of decision trees,	1	
		screening software)		
	6.2	Report the number of people involved and how they con-	Υ	
		tributed (e.g. independent parallel screening)	•	
Data collection process				
	7.1	Describe where in the reports data were collected from (e.g.	Υ	
		text or figures)		
	7.2	Describe how data were collected (e.g. software used to	Υ	
		digitize figures, external data sources)		
	7.3	Describe moderator variables that were constructed from col-	Υ	
		lected data (e.g. number of generations calculated from years		
		and average generation time)		
	7.4	Report how missing or ambiguous information was dealt with	Υ	
		during data collection (e.g. authors of original studies were		
		contacted for missing descriptive statistics, and/or effect sizes		
	7.5	were calculated from test statistics)		
	7.5	Report who collected data	Y	N 1 12 1
	7.6	State the number of extractions that were checked for accuracy by so puthers	N	No checking by secon author
		racy by co-authors		author
Data items				
	8.1	Describe the key data sought from each study	Υ	
	8.2	Describe items that do not appear in the main results, or which	Υ	
		could not be extracted due to insufficient information		
	8.3	Describe main assumptions or simplifications that were made	Υ	
		(e.g. categorising both 'length' and 'mass' as 'morphology')		
	8.4	Describe the type of replication unit (e.g. individuals, broods, study sites)	Υ	
Assessment of individua	l study quality	stady sices,		
	9.1	Describe whether the quality of studies included in the sys-	Υ	
	7.2	tematic review or meta-analysis was assessed (e.g. blinded	•	
		data collection, reporting quality, experimental versus obser-		
		vational)		
	9.2	Describe how information about study quality was incorpo-	Υ	
		rated into analyses (e.g. meta-regression and/or sensitivity		
		analysis)		
Effect size measures				
	10.1	Describe effect size(s) used	Υ	
	10.2	Provide a reference to the equation of each calculated effect	Υ	
		size (e.g. standardised mean difference, log response ratio) and		
		(if applicable) its sampling variance		
	10.3	If no reference exists, derive the equations for each effect size	N	
		and state the assumed sampling distribution(s)		

Checklist item	Sub-item number	Sub-item	Reported by authors?	Notes
	11.1	Describe any steps taken to deal with missing data during	Υ	
		analysis (e.g. imputation, complete case, subset analysis)		
	11.2	Justify the decisions made to deal with missing data	Υ	
Meta-analytic model desc	cription	·		
·	12.1	Describe the models used for synthesis of effect sizes	Υ	
	12.2	The most common approach in ecology and evolution will be	Y	
	12.2	a random-effects model, often with a hierarchical/multilevel	•	
		structure. If other types of models are chosen (e.g. common/-		
		fixed effects model, unweighted model), provide justification		
		for this choice		
Software				
	13.1	Describe the statistical platform used for inference (e.g. R)	Υ	
	13.1	· · · · · · · · · · · · · · · · · · ·	Υ	
		Describe the packages used to run models Describe the functions used to run models	Υ Υ	
	13.3 13.4		Y Y	
	13.4	Describe any arguments that differed from the default set-	ſ	
	13.5	tings Describe the version numbers of all software used	N	Available on Github
	13.3	Describe the version numbers of all software used	1 N	Available off Gitting
Non-independence				
	14.1	Describe the types of non-independence encountered (e.g.	Υ	
		phylogenetic, spatial, multiple measurements over time)		
	14.2	Describe how non-independence has been handled	Υ	
	14.3	Justify decisions made	Υ	
Meta-regression and mod	lel selection			
	15.1	Provide a rationale for the inclusion of moderators (covariates)	Υ	
		that were evaluated in meta-regression models		
	15.2	Justify the number of parameters estimated in models, in rela-	Υ	
		tion to the number of effect sizes and studies (e.g. interaction		
		terms were not included due to insufficient sample sizes)		
	15.3	Describe any process of model selection	Υ	
Publication bias and sens	itivity analyses			
	16.1	Describe assessments of the risk of bias due to missing results	Υ	
		(e.g. publication, time-lag, and taxonomic biases)		
	16.2	Describe any steps taken to investigate the effects of such	Υ	
		biases (if present)		
	16.3	Describe any other analyses of robustness of the results,	Υ	
		e.g. due to effect size choice, weighting or analytical model		
		assumptions, inclusion or exclusion of subsets of the data,		
		or the inclusion of alternative moderator variables in meta-		
		regressions		
Clarification of post hoc a	analyses			
	17.1	When hypotheses were formulated after data analysis, this	N	Not applicable
		should be acknowledged.		

Checklist item	Sub-item number	Sub-item	Reported by authors?	Notes
Metadata, data, and co	de			
	18.1	Share metadata (i.e. data descriptions)	Υ	Available on Github
	18.2	Share data required to reproduce the results presented in the	Υ	Available on Github
		manuscript		
	18.3	Share additional data, including information that was not pre-	Υ	Available on Github
		sented in the manuscript (e.g. raw data used to calculate effect		
		sizes, descriptions of where data were located in papers)		
	18.4	Share analysis scripts (or, if a software package with graph-	Υ	Available on Github
		ical user interface (GUI) was used, then describe full model		
		specification and fully specify choices)		
Results of study select	ion process			
	19.1	Report the number of studies screened	Υ	
	19.2	Report the number of studies excluded at each stage of	Υ	
		screening		
	19.3	Report brief reasons for exclusion from the full text stage	Υ	Available on Github
	19.4	Present a Preferred Reporting Items for Systematic Reviews	Υ	
		and Meta-Analyses (PRISMA)-like flowchart (www.prisma-		
		statement.org).		
Sample sizes and study	characteristics			
	20.1	Report the number of studies and effect sizes for data in-	Υ	
		cluded in meta-analyses		
	20.2	Report the number of studies and effect sizes for subsets of	Υ	
		data included in meta-regressions		
	20.3	Provide a summary of key characteristics for reported out-	Υ	
		comes (either in text or figures; e.g. one quarter of effect sizes		
		reported for vertebrates and the rest invertebrates)		
	20.4	Provide a summary of limitations of included moderators (e.g.	Υ	One moderator only
		collinearity and overlap between moderators)		
	20.5	Provide a summary of characteristics related to individual	Υ	
		study quality (risk of bias)		
Meta-analysis				
	21.1	Provide a quantitative synthesis of results across studies,	Υ	
		including estimates for the mean effect size, with confidence/-		
		credible intervals		
Heterogeneity				
	22.1	Report indicators of heterogeneity in the estimated effect (e.g.	Υ	
		I2, tau2 and other variance components)		
Meta-regression				
	23.1	Provide estimates of meta-regression slopes (i.e. regression	Υ	
		coefficients) and confidence/credible intervals		
	23.2	Include estimates and confidence/credible intervals for all	Υ	
		moderator variables that were assessed (i.e. complete report-		
		ing)		

Checklist item	Sub-item number	Sub-item	Reported by authors?	Notes
	23.3	Report interactions, if they were included	Υ	
	23.4	Describe outcomes from model selection, if done (e.g. R2 and	Υ	
		AIC)		
Outcomes of publicat	tion bias and sensiti	vity analyses		
	24.1	Provide results for the assessments of the risks of bias (e.g.	Υ	
		Egger's regression, funnel plots)		
	24.2	Provide results for the robustness of the review's results (e.g.	Υ	
		subgroup analyses, meta-regression of study quality, results		
		from alternative methods of analysis, and temporal trends)		
Discussion				
	25.1	Summarise the main findings in terms of the magnitude of	Υ	
		effect		
	25.2	Summarise the main findings in terms of the precision of ef-	Υ	
		fects (e.g. size of confidence intervals, statistical significance)		
	25.3	Summarise the main findings in terms of their heterogeneity	Υ	
	25.4	Summarise the main findings in terms of their biological/prac-	Υ	
		tical relevance		
	25.5	Compare results with previous reviews on the topic, if avail-	Υ	
		able		
	25.6	Consider limitations and their influence on the generality	Υ	
	20.0	of conclusions, such as gaps in the available evidence (e.g.		
		taxonomic and geographical research biases)		
Contributions and fu	nding			
	26.1	Provide names, affiliations, and funding sources of all co-	N	Not applicable
		authors		
	26.2	List the contributions of each co-author	N	Not applicable
	26.3	Provide contact details for the corresponding author	N	Not applicable
	26.4	Disclose any conflicts of interest	N	Not applicable
References				
	27.1	Provide a reference list of all studies included in the systematic	Υ	
		review or meta-analysis		
	27.2	List included studies as referenced sources (e.g. rather than	N	Included in Supplemen
		listing them in a table or supplement)		tary Materials