Meta-analysis of the efficacy and safety of vamorolone in Duchenne muscular dystrophy

Supplementary Material

Supplementary Text 1: Complete search string
Search: **((Vamorolone) OR (VBP15)) AND (Duchenne Muscular Dystrophy)** Sort by: **Most Recent**
("vbp15 compound"[Supplementary Concept] OR "vbp15 compound"[All Fields] OR "vamorolone"[All Fields] OR "VBP15"[All Fields]) AND ("muscular dystrophy, duchenne"[MeSH Terms] OR ("muscular"[All Fields] AND "dystrophy"[All Fields] AND "duchenne"[All Fields]) OR "duchenne muscular dystrophy"[All Fields] OR ("duchenne"[All Fields] AND "muscular"[All Fields] AND "dystrophy"[All Fields]))
Translations
Vamorolone: "VBP15 compound"[Supplementary Concept] OR "VBP15 compound"[All Fields] OR "vamorolone"[All Fields]
Duchenne Muscular Dystrophy: "muscular dystrophy, duchenne"[MeSH Terms] OR ("muscular"[All Fields] AND "dystrophy"[All Fields] AND "duchenne"[All Fields]) OR "duchenne muscular dystrophy"[All Fields] OR ("duchenne"[All Fields] AND "muscular"[All Fields] AND "dystrophy"[All Fields])

Supplementary Table 1: Risk of bias (ROB) and quality assessment for each study

Study ID	**ROB assessment for RCT according to ROB2 tool**												
	D1	D2	D3	D4	D5	Other bias							Overall bias
Guglieri 2022 [22]	LR	LR	LR	LR	LR	LR							Low risk of bias
	ROB assessment for NRSI according to ROBINS-I tool												
	D1	D2	D3	D4	D5	D6	D7						Overall bias
Hoffman 2019 [21]	HR	LR	LR	LR	LR	LR	MR						Moderate risk of bias
Smith 2020 [8]	MR	LR	LR	HR	LR	LR	LR						Moderate risk of bias
Mah 2022 [20]	LR	HR	LR	LR	LR	LR	LR						Moderate risk of bias
	Quality assessment for before-after (Pre-Post) Studies with no control group according to NIH tool												
	Q1	Q2	Q3	Q4	Q5	Q6	Q7	Q8	Q9	Q10	Q11	Q12	Overall quality
Conklin 2018 [9]	Y	Y	Y	Y	Y	Y	Y	Y	Y	Y	Y	Y	Excellent (12)

ROB2 tool: bias due to/in: D1: randomization process; D2: deviations from intended interventions; D3: measurement of the outcome; D4 missing outcome data; D5: selection of the reported result. **ROBINS1 tool**: bias due to/in: D1: confounding; D2: selection of participants into the study; D3: classification of interventions; D4: deviations from intended interventions; D5: missing data; D6: measurement of outcomes; D7: selection of the reported result. **NIH tool for pre and post studies**: Q1: Objective clearly stated; Q2: eligibility criteria described; Q3: representative patient population; Q4: all eligible participants enrolled in study; Q5: sufficient sample size; Q6: intervention described; Q7: outcome measures specified; Q8: outcome assessors blinded; Q9: loss to follow-up; Q10: statistical analysis of outcome measures before and after intervention; Q11: interrupted time-series design; Q12: individual data used for group-level effects. LR: low risk; MR: moderate risk; HR, high risk; SC: some concerns; Y, yes; N, no; NA, not applicable; CD: cannot determined.

Section and Topic	Item #	Checklist item	Location where item is reported
TITLE			
Title	1	Identify the report as a systematic review.	1
ABSTRACT			
Abstract	2	See the PRISMA 2020 for Abstracts checklist.	1
INTRODUCTION			
Rationale	3	Describe the rationale for the review in the context of existing knowledge.	1,2
Objectives	4	Provide an explicit statement of the objective(s) or question(s) the review addresses.	1,2
METHODS			
Eligibility criteria	5	Specify the inclusion and exclusion criteria for the review and how studies were grouped for the syntheses.	2
Information sources	6	Specify all databases, registers, websites, organisations, reference lists and other sources searched or consulted to identify studies. Specify the date when each source was last searched or consulted.	2
Search strategy	7	Present the full search strategies for all databases, registers and websites, including any filters and limits used.	2, Supplementary Text 1
Selection process	8	Specify the methods used to decide whether a study met the inclusion criteria of the review, including how many reviewers screened each record and each report retrieved, whether they worked independently, and if applicable, details of automation tools used in the process.	2,3
Data collection process	9	Specify the methods used to collect data from reports, including how many reviewers collected data from each report, whether they worked independently, any processes for obtaining or confirming data from study investigators, and if applicable, details of automation tools used in the process.	2,3
Data items	10a	List and define all outcomes for which data were sought. Specify whether all results that were compatible with each outcome domain in each study were sought (e.g. for all measures, time points, analyses), and if not, the methods used to decide which results to collect.	3
	10b	List and define all other variables for which data were sought (e.g. participant and intervention characteristics, funding sources). Describe any assumptions made about any missing or unclear information.	3
Study risk of bias assessment	11	Specify the methods used to assess risk of bias in the included studies, including details of the tool(s) used, how many reviewers assessed each study and whether they worked independently, and if applicable, details of automation tools used in the process.	3
Effect measures	12	Specify for each outcome the effect measure(s) (e.g. risk ratio, mean difference) used in the synthesis or presentation of results.	1.3
Synthesis methods	13a	Describe the processes used to decide which studies were eligible for each synthesis (e.g. tabulating the study intervention characteristics and comparing against the planned groups for each synthesis (item #5)).	3
	13b	Describe any methods required to prepare the data for presentation or synthesis, such as handling of missing summary statistics, or data conversions.	3
	13c	Describe any methods used to tabulate or visually display results of individual studies and syntheses.	3
	13d	Describe any methods used to synthesize results and provide a rationale for the choice(s). If meta-analysis was performed, describe the model(s), method(s) to identify the presence and extent of statistical heterogeneity, and software package(s) used.	3
	13e	Describe any methods used to explore possible causes of heterogeneity among study results (e.g. subgroup analysis, meta-regression).	3
	13f	Describe any sensitivity analyses conducted to assess robustness of the synthesized results.	N/A
Reporting bias assessment	14	Describe any methods used to assess risk of bias due to missing results in a synthesis (arising from reporting biases).	3
Certainty assessment	15	Describe any methods used to assess certainty (or confidence) in the body of evidence for an outcome.	3
RESULTS			
Study selection	16a	Describe the results of the search and selection process, from the number of records identified in the search to the number of studies included in the review, ideally using a flow diagram.	4-10
	16b	Cite studies that might appear to meet the inclusion criteria, but which were excluded, and explain why they were excluded.	4-10
Study characteristics	17	Cite each included study and present its characteristics.	4-10
Risk of bias in studies	18	Present assessments of risk of bias for each included study.	4, Supplementary table 1
Results of individual studies	19	For all outcomes, present, for each study: (a) summary statistics for each group (where appropriate) and (b) an effect estimate and its precision (e.g. confidence/credible interval), ideally using structured tables or plots.	4-10
Results of syntheses	20a	For each synthesis, briefly summarise the characteristics and risk of bias among contributing studies.	4-10
	20b	Present results of all statistical syntheses conducted. If meta-analysis was done, present for each the summary estimate and its precision (e.g. confidence/credible interval) and measures of statistical heterogeneity. If comparing groups, describe the direction of the effect.	4-10
	20c	Present results of all investigations of possible causes of heterogeneity among study results.	4-10
	20d	Present results of all sensitivity analyses conducted to assess the robustness of the synthesized results.	N/A
Reporting biases	21	Present assessments of risk of bias due to missing results (arising from reporting biases) for each synthesis assessed.	N/A

Section and Topic	Item #	Checklist item	Location where item is reported
Certainty of evidence	22	Present assessments of certainty (or confidence) in the body of evidence for each outcome assessed.	4-10
DISCUSSION			
Discussion	23a	Provide a general interpretation of the results in the context of other evidence.	10-11
	23b	Discuss any limitations of the evidence included in the review.	11
	23c	Discuss any limitations of the review processes used.	11
	23d	Discuss implications of the results for practice, policy, and future research.	11
OTHER INFORMATION			
Registration and protocol	24a	Provide registration information for the review, including register name and registration number, or state that the review was not registered.	1,2
	24b	Indicate where the review protocol can be accessed, or state that a protocol was not prepared.	1,2
	24c	Describe and explain any amendments to information provided at registration or in the protocol.	N/A
Support	25	Describe sources of financial or non-financial support for the review, and the role of the funders or sponsors in the review.	N/A (Statement in supplement)
Competing interests	26	Declare any competing interests of review authors.	N/A (Statement in supplement)
Availability of data, code and other materials	27	Report which of the following are publicly available and where they can be found: template data collection forms; data extracted from included studies; data used for all analyses; analytic code; any other materials used in the review.	Supplement