APPENDIX 6: AMSTAR MEASUREMENT TOOL TO ASSESS SYSTEMATIC REVIEWS⁵

1.	Was a priori design provided? The research question and inclusion criteria should be established before the conduct of the review.	□ Yes □ No
	should be established before the conduct of the review.	☐ Can't answer
		□ Not applicable
2.	Was there duplicate study selection and data extraction? There should be at	□ Yes
	least two independent data extractors and a consensus procedure for	□ No
	disagreements should be in place.	□ Can't answer
	8	□ Not applicable
3.	Was a comprehensive literature search performed? At least two electronic	□ Yes
	sources should be searched. The report must include years and databases	□ No
	used (e.g. Central, Embase, and MEDLINE). Key words and/or MESH terms	□ Can't answer
	must be stated and where feasible the search strategy should be provided. All	□ Not applicable
	searches should be supplemented by consulting current contents, reviews,	11
	textbooks, specialized registers, or experts in the particular field of study,	
	and by reviewing the references in the studies found.	
4.	Was the status of publication (i.e., grey literature) used as an inclusion	□ Yes
	criterion? The authors should state that they searched for reports regardless	□ No
	of their publication type. The authors should state whether or not they	□ Can't answer
	excluded any reports (from the systematic review), based on their publication	□ Not applicable
	status, language, etc.	
5.	Was a list of studies (included and excluded) provided? A list of included	□ Yes
	and excluded studies should be provided.	□No
		□ Can't answer
	YYY A A A A A A A A A A A A A A A A A A	□ Not applicable
6.	Were the characteristics of the included studies provided? In an aggregated	□ Yes
	form such as a table, data from the original studies should be provided on the	□ No
	participants, interventions, and outcomes. The ranges of characteristics in all	☐ Can't answer
	the studies analyzed e.g. age, race, sex, relevant socioeconomic data, disease status, duration, severity, or other diseases should be reported.	□ Not applicable
7	Was the scientific quality of the included studies assessed and documented?	□ Yes
١.	A priori methods of assessment should be provided (e.g., for effectiveness	□ No
	studies if the author[s] chose to include only randomised, double-blind,	☐ Can't answer
	placebo controlled studies, or allocation concealment as inclusion criteria);	□ Not applicable
	for other types of studies alternative items will be relevant.	
8.	Was the scientific quality of the included studies used appropriately in	□ Yes
	formulating conclusions? The results of the methodological rigor and	□ No
	scientific quality should be considered in the analysis and the conclusions of	□ Can't answer
	the review, and explicitly stated in formulating recommendations.	□ Not applicable
9.	Were the methods used to combine the findings of studies appropriate? For	□ Yes
	the pooled results, a test should be done to ensure the studies were	□ No
	combinable, to assess their homogeneity (i.e., Chi-squared test for	□ Can't answer
	homogeneity, I2). If heterogeneity exists, a random effects model should be	□ Not applicable
	used and/or the clinical appropriateness of combining should be taken into	
	consideration (i.e., is it sensible to combine?).	

10. Was the likelihood of publication bias assessed? An assessment of	□ Yes
publication bias should include a combination of graphical aids (e.g., funnel	□ No
plot, other available tests) and/or statistical tests (e.g., Egger regression test).	□ Can't answer
	□ Not applicable
11. Was the conflict of interest included? Potential sources of support should be	□ Yes □ No
clearly acknowledged in both the systematic review and the included studies.	□ Can't answer
	□ Not applicable