SUPPLEMENTARY MATERIAL

Table S1: PRISMA 2020 Main Checklist

Topic	No.	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	
INTRODUCTION			
Rationale	3	Describe the rationale for the review in the context of existing knowledge.	5,6
Objectives	4	Provide an explicit statement of the objective(s) or question(s) the review addresses.	6
METHODS			
Eligibility criteria	Specify the inclusion and exclusion criteria for the review and how studies were grouped for the syntheses.		6,7
Information sources	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.		8
Search strategy	7 Present the full search strategies for all databases, registers and websites, including any filters and limits used.		8
Selection process	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.		8,9
Data collection process	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.		8,9

Topic	No.	Item	Location where item is reported
Data items 10		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.	
	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.	7
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.	9
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.	9
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)).	8,9
	13b	Describe any methods required to prepare the data for presentation or synthesis, such as handling of missing summary statistics, or data conversions.	9
	13c	Describe any methods used to tabulate or visually display results of individual studies and syntheses.	9
	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.	9
	13e	Describe any methods used to explore possible causes of heterogeneity among study results (e.g. subgroup analysis, meta-regression).	Not applicable
	13f	Describe any sensitivity analyses conducted to assess robustness of the synthesized results.	Not applicable
Reporting bias assessment	14	Describe any methods used to assess risk of bias due to missing results in a synthesis (arising from reporting biases).	Not applicable
Certainty assessment	15	Describe any methods used to assess certainty (or confidence) in the body of evidence for an outcome.	9
RESULTS			

Topic	No.	Item	Location where item is reported
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.	10
	16b	Cite studies that might appear to meet the inclusion criteria, but which were excluded, and explain why they were excluded.	10
Study characteristics	17	Cite each included study and present its characteristics.	10,11
Risk of bias in studies	18	Present assessments of risk of bias for each included study.	11
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), deally using structured tables or plots.	
Results of syntheses	20a	For each synthesis, briefly summarise the characteristics and risk of bias among contributing studies.	
	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.		13-15
	20c	Present results of all investigations of possible causes of heterogeneity among study results.	Not applicable
	Present results of all sensitivity analyses conducted to assess the robustness of the synthesized results.		Not applicable
Reporting biases	21	Present assessments of risk of bias due to missing results (arising from reporting biases) for each synthesis assessed.	Not applicable
Certainty of evidence	22	Present assessments of certainty (or confidence) in the body of evidence for each outcome assessed.	
DISCUSSION			
Discussion	23a	Provide a general interpretation of the results in the context of other evidence.	15-18
	23b	Discuss any limitations of the evidence included in the review.	18
	23c	Discuss any limitations of the review processes used.	18
	23d	Discuss implications of the results for practice, policy, and future research.	19

Topic	No.	Item	Location where item is reported
OTHER INFORMATION			
Registration and protocol			6
	24b Indicate where the review protocol can be accessed, or state that a protocol was not prepared.		6
	24c	Describe and explain any amendments to information provided at registration or in the protocol.	Not applicable
Support	25	Describe sources of financial or non-financial support for the review, and the role of the funders or sponsors in the review.	
Competing interests	26	Declare any competing interests of review authors.	20
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.	20

Table S1b: PRIMSA Abstract Checklist

Topic	No.	Item	Reported?		
TITLE					
Title	1	Identify the report as a systematic review.			
BACKGROUND					
Objectives	2	Provide an explicit statement of the main objective(s) or question(s) the review addresses.	Yes		
METHODS					
Eligibility criteria	3	Specify the inclusion and exclusion criteria for the review.	Yes		
Information sources	4	Specify the information sources (e.g., databases, registers) used to identify studies and the date when each was last searched.			
Risk of bias	5	Specify the methods used to assess risk of bias in the included studies.			
Synthesis of results	6	Specify the methods used to present and synthesize results.			
RESULTS					
Included studies	7	Give the total number of included studies and participants and summarise relevant characteristics of studies.	Yes		
Synthesis of results	8	Present results for main outcomes, preferably indicating the number of included studies and participants for each. If meta-analysis was done, report the summary estimate and confidence/credible interval. If comparing groups, indicate the direction of the effect (i.e. which group is favoured).			
DISCUSSION					
Limitations of evidence	9	Provide a brief summary of the limitations of the evidence included in the review (e.g. study risk of bias, inconsistency and imprecision).			
Interpretation	10	Provide a general interpretation of the results and important implications.	Yes		
OTHER					
Funding	11	Specify the primary source of funding for the review.	No		
Registration	12	Provide the register name and registration number.	No		

From: Page MJ, McKenzie JE, Bossuyt PM, Boutron I, Hoffmann TC, Mulrow CD, et al. The PRISMA 2020 statement: an updated guideline for reporting systematic reviews. MetaArXiv. 2020, September 14. DOI: 10.31222/osf.io/v7gm2. For more information, visit: www.prisma-statement.org

Table S2a: Search methods

		Number of articles in each database			
		Scopus	Cochrane (CENTRAL)	PubMed (MEDLINE)	EMBASE
Domain	Search details	Searched on 21 July 2020	Searched on 21 July 2020	Searched from 1947 to 21 July 2020	Searched from 1947 to 21 July 2020
yoga and PCOS and randomized controlled trial	("yoga"[MeSH Terms] OR "yoga"[All Fields]) AND "PCOS"[All Fields] AND ("randomized controlled trial"[Publication Type] OR "randomized controlled trials as topic"[MeSH Terms] OR "randomized controlled trial"[All Fields] OR "randomized controlled trial"[All Fields])	5	11	3	5
yoga and PCOS	("yoga"[MeSH Terms] OR "yoga"[All Fields]) AND PCOS [All Fields]	8	7	10	16
lifestyle intervention and PCOS	("life style"[MeSH Terms] OR ("life"[All Fields] AND "style"[All Fields]) OR "life style"[All Fields] OR "lifestyle"[All Fields]) AND ("methods"[MeSH Terms] OR "methods"[All Fields] OR "intervention"[All Fields]) AND PCOS [All Fields]	257	170	83	145
lifestyle manageme nt and PCOS	("life style"[MeSH Terms] OR ("life"[All Fields] AND "style"[All Fields]) OR "life style"[All Fields] OR "lifestyle"[All Fields]) AND ("organization and administration"[MeSH Terms] OR ("organization"[All Fields] AND "administration"[All Fields]) OR "organization and administration"[All Fields] OR "management"[All Fields] OR "disease management"[MeSH Terms] OR ("disease"[All Fields] AND "management"[All Fields]) OR "disease management"[All Fields]) AND	256	48	26	160
complemen tary and	complementary [All Fields] AND ("complementary	20	13	13	46

		Number of articles in each database			
Damain		Scopus	Cochrane (CENTRAL)	PubMed (MEDLINE)	EMBASE
Domain	Search details	Searched on 21 July 2020	Searched on 21 July 2020	Searched from 1947 to 21 July 2020	Searched from 1947 to 21 July 2020
alternative	therapies"[MeSH Terms] OR				
therapy and	("complementary"[All Fields]				
PCOS	AND "therapies"[All Fields])				
	OR "complementary				
	therapies"[All Fields] OR				
	("alternative"[All Fields] AND				
	"therapy"[All Fields]) OR				
	"alternative therapy"[All				
	Fields]) AND PCOS[All Fields]				
	Total	546	249	135	372

Table S2b: Search methods

Domain	CTRI
	(Searched on 21 July 2020)
PCOS	101
Yoga	530
Lifestyle intervention	33
Lifestyle management	4
Complementary and alternative therapy	0
Total	668

Table S3: Conversion factors

Parameter	Convert from	Convert to	Conversion factor
FBG	mg/dL	mmol/L	0.056
FI	μIU/mL	pmol/L	6.0

FBG fasting blood glucose; FI fasting insulin