Appendix S1

Search strategy

Ovid MEDLINE(R) ALL <1946 to October 06, 2022>

Medline

1	exp smoking/ or Smokers/ 616696
2	(smoking or smoker* or tobacco usage or "tobacco use" or cigarette*).tw,kf. 795947
3	1 or 2 954726
4	(male* and female*).tw,kf. 1422627
5	((sex or gender) adj2 (difference* or disparit*)).tw. 227684
6	sex* dimorph*.tw,kf. 37016
7	(gender difference* or gender disparit*).kw. 12034
8	(sex or gender).ti. 359929
9	(gender based or gender specific).tw,kw. 41479
10	Sex Factors/ 290947
11	Sex Characteristics/ 63305
12	(sex dependent or sex factor* or sex characteristic*).tw,kf. 21801
13	or/4-12 1963943
14	3 and 13 88495
15	exp Renal Insufficiency, Chronic/ 268704
16	(chronic adj2 (renal or kidney) adj3 (disease* or insufficienc* or failure)).tw,kf. 259103
17	ckd.tw,kf. 113025
18	(end stage renal or end stage kidney).tw,kf. 124634
19	esrd.tw,kf. 52605
20	exp Renal Dialysis/ 256083
21	dialysis.tw,kf. 303583
22	Albuminuria/ 38070
23	Albuminuria.tw,kf. 34580
24	Glomerular Filtration Rate/ 121237
25	Creatinine/ 296510
26	Creatinine.tw,kf. 349307
27	Nephrosis/ 8961
28	nephrosis.tw,kf. 10893
29	Nephrotic Syndrome/ 48825
30	Nephrotic Syndrome*.tw,kf. 49833
31	Kidney Glomerulus/ 55903
32	Glomerulonephritis/ 67130
33	glomerulonephritis.tw,kf. 75209
34	eskd.tw,kf. 6216
35	(glomerular filtration rate or egfr).tw,kf. 304654
36	h?emodialysis.tw,kf. 215341
37	or/15-36 1559997

38	14 and 37 5203	
39	exp animals/ not humans/	17337966
40	38 not 39 3972	
41	limit 40 to english language	3727
40	41	

42 41 use medall 1556

Embase

43	smoking/ or "tobacco use"/ or smoking habit/ or cigarette smoke/ or cigarette/ or cigarette/	rette
smokin	ng/ 618541	
44	(smoking or smoker* or tobacco usage or "tobacco use" or cigarette*).tw. 790	476
45	43 or 44 949283	
46	(male* and female*).tw. 1420205	
47	((sex or gender) adj2 (difference* or disparit*)).tw. 227684	
48	sex* dimorph*.tw. 34884	
49	(sex or gender).ti. 359929	
50	(gender based or gender specific).tw. 41422	
51	*sex factor/ or sex difference/ 502184	
52	sex dependent.tw. 13412	
53	*sexual characteristics/ 1371	
54	(sex dependent or sex factor* or sex characteristic*).tw. 17039	
55	or/46-54 2023834	
56	45 and 55 97233	
57	exp chronic kidney failure/ 237973	
58	(chronic adj2 (kidney or renal) adj3 (disease* or failure* or insufficienc*)).tw. 246	6669
59	ckd.tw. 110960	
60	(end stage renal or end stage kidney or esrd or eskd).tw. 133081	
61	exp *hemodialysis/ 148766	
62	(dialysis or h?emodialysis).tw. 418128	
63	*albuminuria/ 12455	
64	Albuminuria.tw. 32599	
65	exp *glomerulus filtration rate/ 16857	
66	(Glomerular Filtration Rate or egfr).tw. 299384	
67	*creatinine/ 22562	
68	Creatinine.tw. 346978	
69	*nephrosis/ 4732	
70	Nephrosis.tw. 9790	
71	*nephrotic syndrome/ 29906	
72	Nephrotic Syndrome*.tw. 48051	
73	*glomerulus/ 9046	
74	*glomerulonephritis/ or *chronic glomerulonephritis/ 40679	
75	Glomerulonephritis.tw. 71356	
76	or/57-75 1353375	
77	56 and 76 5651	

- 78 (exp animals/ or animal experimentation/) not exp humans/
- 11158265

- 79 conference abstract.pt. 4559436
- 80 77 not (78 or 79) 3971
- 81 limit 80 to english language 3658
- 82 81 use emczd 2565
- 83 42 or 82 4121



PRISMA 2020 Checklist

Section and Topic	ltem #	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.	2
INTRODUCTION			
Rationale	3	Describe the rationale for the review in the context of existing knowledge.	4
Objectives	4	Provide an explicit statement of the objective(s) or question(s) the review addresses.	5
METHODS			
Eligibility criteria	5	Specify the inclusion and exclusion criteria for the review and how studies were grouped for the syntheses.	19
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.	5
Search strategy	7	Present the full search strategies for all databases, registers and websites, including any filters and limits used.	41
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.	5
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.	5
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.	6
	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.	6
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.	6
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.	n/a
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)).	n/a
	13b	Describe any methods required to prepare the data for presentation or synthesis, such as handling of missing summary statistics, or data conversions.	n/a
	13c	Describe any methods used to tabulate or visually display results of individual studies and syntheses.	n/a
	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.	n/a
	13e	Describe any methods used to explore possible causes of heterogeneity among study results (e.g. subgroup analysis, meta-regression).	n/a
	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).	6
Certainty assessment	15	Describe any methods used to assess certainty (or confidence) in the body of evidence for an outcome.	6



Section and Topic	ltem #	Checklist item	Location where item is reported
RESULTS	1		
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.	6
	16b	Cite studies that might appear to meet the inclusion criteria, but which were excluded, and explain why they were excluded.	n/a
Study characteristics	17	Cite each included study and present its characteristics.	20
Risk of bias in studies	18	Present assessments of risk of bias for each included study.	39
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.	n/a
Results of	20a	For each synthesis, briefly summarise the characteristics and risk of bias among contributing studies.	6
syntheses	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.	n/a
	20c	Present results of all investigations of possible causes of heterogeneity among study results.	n/a
	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
Certainty of evidence	22	Present assessments of certainty (or confidence) in the body of evidence for each outcome assessed.	n/a
DISCUSSION			
Discussion	23a	Provide a general interpretation of the results in the context of other evidence.	8
	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 INFORMA	TION		
Registration and	24a	Provide registration information for the review, including register name and registration number, or state that the review was not registered.	5
protocol	24b	Indicate where the review protocol can be accessed, or state that a protocol was not prepared.	5
	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.	11
Competing interests	26	26 Declare any competing interests of review authors.	
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.	12

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. BMJ 2021;372:n71. doi: 10.1136/bmj.n71