	Α	В	C	D	E	F	G	Н	1
1	Toegevoegd door	Literatuur (titel, auteur, jaartal)	Type literatuur (review of paper, preprint of published)	Quality of evidence	QoE check	Toelichting QoE s12e indirectness, inconsistency, indirectness, imprecision, effect size, right confounders, dose response)	Risk of bias (study limitations: design, inclusion and sample info, measurement, confounding, folluw-up)	Toelichting bias	Een zo kort mogelijke samenvatting voor in het document
2		Mask-Wearing Increased After a Government Recommendation: A Natural Experiment in the U.S. During the COVID-19 Pandemic	peer reviewed single study observationeel	moderate		512e geen gevalideerde maten, self reports. Groot N verschil in voor- na meting	Estate Estate and and a	onderdeel van bestaand panel	meer kopen en dragen van mondkapjes na aanbeveling

Г	A	J	K	L	M	N	0
	Toegevoegd door	Type studie (zie werkblad hierna)	Land	Verplichting of advies	Steekproef (grootte, populatie)	Recruitment (opvallende in/excl cr., hoe geworven)	Representatief? (is deze studie vergelijkbaar met NL situatie of populatie?)
	5.1.2e		vs	advies	3933		Westers
	5.1.2e						

	Α	P	Q	R	S	T	U	V
1	Toegevoegd door	Sleutelwoorden (gedrag: determinanten/omstandigheden/redenen/mat e van naleving/verschillen/interventies)	Doel studie	Methode (controle groep, etc)	Measures DV en IV (item/schaal/gevalideerd/inte ntie/gedrag/self-report)	Confounders	Belangrijkste bevindingen	Beoordeling effect sizes
2		The study found significant increases in reported mask wearing (+12 percentage points) and mask buying (+7 points).	effect aanbevling op mondkapjes kopen en dragen bekijken					
3	5.1.2e							

	A	W	X
1	Toegevoegd door	Verschillen tussen subpopulaties	Link naar studie
	5.1.2e		
2			
3	5.1.2e		

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	-	evidence als tabblad hiernaast, Bias, zijn		zowel observationeel als	Table 5.2: Factors that can reduce the	quality of the evidence
1		meegenomen)		RCT>		
2	Table 5.1:	Quality of Evidence Grades		Quality of evidence hangt af van volgende factoren, waaronder design (wat je bij Bias bekijkt)	Factor	Consequence
3	Grade	Definition			Limitations in study design or execution (risk of bias)	↓ 1 or 2 levels
4	High	We are very confident that the true effect lies close to that of the estimate of the effect.			Inconsistency of results	↓ 1 or 2 levels
5	Moderate	We are moderately confident in the effect estimate: The true effect is likely to be close to the estimate of the effect, but there is a possibility that it is substantially different			Indirectness of evidence	↓ 1 or 2 levels
6	Low	Our confidence in the effect estimate is limited: The true effect may be substantially different from the estimate of the effect.			Imprecision	↓ 1 or 2 levels
7	Very Low	We have very little confidence in the effect estimate: The true effect is likely to be substantially different from the estimate of effect			Publication bias	↓ 1 or 2 levels
8					Table 5.3: Factors that can increase th	e quality of the evidence
		VOOR REVIEWS, gebruik dit				
		formulier voor een oordeel en sla			Factor	Consequence
9		deze op				
10	3				Large magnitude of effect	† 1 or 2 levels
11		5.1.2h			All plausible confounding would reduce the demonstrated effect or increase the effect if no effect was observed	† 1 level
12					Dose-response gradient	† 1 level
13						
14						
15						

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1		**	is a continuum; any discrete categoris	ation i	nvolves	some o			J
3	toelichting zie linksonder Study Design en volgend tabblad voor Risk of Bias. Observationeel kan hierdoor eigenlijk niet als HIGH beoordeeld worden. Niet toegelichte heterogeniteit van resultaten (vooral bij syst reviews, als er veel verschillende bevindingen zijn, gemengd bewijs).	each individual factor evidence for an outcor exclusive. Therefore, factor for downgradicategory and among particular factor, the	cing the quality of evidence are additive r is added together with the other factors one – grading the quality of evidence inv. GRADE is not a quantitative system for a gor upgrading reflects not discrete cate the categories. When the body of evidendecision about whether a study falls about ity (by one or more factors) depends or	to reduvolves jegrading gories le is inverse ve or be	udgements of the quant to the q	crease ents wh uality o ntinuur ate witl	the quaich are f evider within respec	lity of not nce. Each n each et to a	
5	Bijvoorbeeld gemeten met een surrogaat maat (niet gedrag, maar intentie of zelfgerapporteerd gedrag) Of nt andere interventie (niet thuisblijven bij klachten maar thuisbiljven in het algemeen).	Study Design	inty (by one or more factors) depends of	Ljuagn	nent.				
6 7 8	Kleine steekproef of kleine hoeveelheid events, dus wijd confidence interval Lastig te achterhalen, gaat erom in hoevrre er studies met negatieve of andere resultaten niet zijn gepubliceerd en dus niet zijn opgenomen. Vooral voor syst reviews relevante factor. Bij losse studies gaat het om reporting bias (zijn er	For recommendation the accuracy of diag Randomized trials p	cal to judgments about the quality of evidence. ns regarding management strategies – as opposed to nostic tests – rovide, in general, far stronger evidence than observ nal studies provide stronger evidence than uncontro	ational st	tudies, an				
9		Randomized trials v	oach to quality of evidence: vithout important limitations provide high quality evi es without special strengths or important limitations		low				
10	Als er een groot effect wordt gevonden. For simple regression β is like R. Thus I would use R rules of thumb I use the follwoing with my Psychology students: β < 0.1 - Small effect size β \(\in 0.1; 0.5 [- Medium effect size β \(\in 0.5 - Large effect size. For multiple regression these rules are not that straightfoward, but for Social Sciences they seem to hold (also following Cohen's d suggestions). Is er gecontroleerd voor plausibele confounders?	randomized trials ai Non-randomised ex quality evidence, bu as lack of concealm Case series and cass intervention. Source	al strengths can, however, modify the quality of the on observational studies. perimental trials (quasi-RCT) without important limit it will automatically be downgraded for limitations ir ent of allocation and tie with a provider (e.g. chart nue reports are observational studies that investigate o e of control group results is implicit or unclear, thus, ow to very low quality evidence.	ations al: design (umber). nly patier	so provide risk of bia nts expose	s) – such			
12		Expert opinion is no of evidence in the c evidence that may l	t a category of quality of evidence. Expert opinion re ontext of experts' experiences and knowledge. Exper e based on interpretation of studies ranging from up	ts may h	ave opinio	on about eries			
14 15		the expert. It is imp	expert's own practice) to randomized trials and sysiortant to describe what type of evidence (whether pass for interpretation.						

1	A	RCT's gebruik deze> tabel 5.4	С	D	E Table 5.4:	F Study lim	G itations in	H randomiz	ed contro	J Iled trials Explanatio	K	L	М
2	RISK OF BIAS = Limitations in the study design and ex	recution may bias the estimates of the treatment effect. Our confidence in the			Lack of al	location co	ncealment		0	Those enro (or period enrolled po problem in	olling patients are in a crossover tria atient will be alloc a "pseudo" or "qua	asi" randomized	
	Nisk of Dias	Dutieg Baschie confounding occurs when one or more prognantic variables (factors that prestict the outcome of interest) also predict the intervention received at baseline. ROBRFS can also address time-anymous confounding, within occur when individuals within thereties the interventions being compared and when post-baseline prognantic factors affect the intervention received after baseline.			Lack of bl		a of paties	nts and outc	1 1 2 2 2 2 2 2 2 2 2 2 2 2 2 2 2 2 2 2	rials with chart numl Patient, ca those adju- are aware allocated (received in	allocation by day ber, etc.).	of week, birth date, cording outcomes, , or data analysts th patients are currently being	
5	In participant selection	When exclusion of some eligible participants, or the initial follow-up time of some participants, or some outcome events is related to both intervention and outcome, there will be an association between intervention and outcome event in the effects of the interventions are identificant. This form of selection being is distinct from conflounding—is specific example to bias due to the including selection design of several least, radio that however, and in interventions are identified users, radio that however, and in interventions are identified users, radio that however, and in intervention are supported to the control of the con			events					intention-t or in nonir failure to conly those patients fo The signifi follow-up, dependent follow-up proportion	n superiority trials; ss to follow-up, and reses considering reatment, and all data are available, ir rates of loss to widely and is tween loss to ents. The higher the in relation to		
6	Due to missing data	Bus that arises when later follow-up is missing for inclividuals initially included and followed (such as differential ios to follow-up that is affected by prognessis faction; it has due to exclusion of incliducials with missing information about intervention status or other variables such as de-confounders.			Selective of	outcome rep	porting		3 3 3 5	groups, the Incomplete outcomes a results. Stopping to overestime than 500 e	e greater the threat e or absent reporti and not others on rial early for bene ates are likely in tr	ng of some the basis of the fit. Substantial ials with fewer ge overestimates are	
7	In measurement of predic/outcome	Bas introduced by either differential or non-differential errors in measurement of outcome data. Such bias can arise when outcome assessors are aware of intervention status, if different methods are used to assess outcomes in different intervention groups, or if measurement errors par related to intervention status or effects.		2	_			1	Empirical stopping ro Use of unv patient-rep Carryover	evidence suggests ales do not reduce validated outcome orted outcomes) effects in crossov nt bias in cluster-r	that formal this bias. measures (e.g.		
9	In selection of reported result	Selective reporting of results in a way that depends on the findings and prevents the estimate from being included in a meta-analysis for other swithersis).											
o	In misclassification of intervention (randomization)	from being included in a meta-analysis (or other synthesis) Bas introduced by either differential or non-differential inkclassification of intervention status Non-differential incidenced in the control of the contro											
9	Due to deviation from intended intervention	outcome or the risk of the outcome, and is linky to lead to bias: All the state arises where are systematic differences between engerimental intervention and comparator groups in the care provided, which represent a deviation from the intended intervention(s)											
10													
12													
14 15	 Comparison: bij interventie studies, goed bek 	ijken wat de comparison conditie is en of studies vergelijkbaar zijn met elkaar.											
16 17	- Outcomes: zijn gebruikte uitkomstmaten ver	gelijkbaar? (gaat het om intentie van gedrag, zelfgrapporteerde naleving, daadwer	kelijke nalev	ring, etc)									
18 19													
19 20													
21											studies	mitations in observational	
23												Under- or over- matching in case-control studies	
24											appropriate eligibility criteria (inclusion of	Selection of exposed and unexposed in cohort studies from different populations	
26											Flawed measurement of both exposure and	Differences in measurement of exposure (e.g. recall bias in ease- control studies)	
26											outcome	 Differential surveillance for outcome in exposed and unexposed in cohort studies 	
28											Failure to adequately control confounding	Failure of accurate measurement of all known prognostic factors Failure to match for	
29												prognostic factors and/or adjustment in statistical analysis	
30											Incomplete or inadequately short follow-up	Especially within prospective cohort studies, both groups should be followed for the same amount of time.	

Comparation of the controller section of t	N	0	P	Q R	S T	U V	W	Х	Υ
Service to the company of the control of the contro	1 2	Observationele studie		over factoren	, staat ook in	link	_		
Service of the city of the cit		inclusion in the sample	The authors should provide clear inclusion and exclusion criteria that	1					
Section for excession and excession of a control of the control of	4	and the setting described	researchers can determine if it is comparable to the population of interest to them. The authors should provide a clear description of the population from which the study participants were selected or recruited, including	1					
where containing the containing function sheet and the containing function	5	measured in a valid and	exposure. Assessing validity requires that a 'god standard' is available to which the measure can be compared. The validity of exposure measurement usually relates to whether a current measure is appropriate or whether a measure of past exposure is needed. Reliability refers to the processes included in an epidemiclogical study to check repeatability of measurements of the exposures. These usually include intro-closerver	1					
Actives continued and active act	6	standard criteria used for measurement of the	either a specified diagnosis or definition. This is more likely to decrease the risk of bias. Characteristics are another useful approach to matching groups, and studies that did not use specified diagnostic methods or	0					
Service throughout the date of the controlled by	7		or concomitant exposures (e.g., smoking). A confounder is a difference between the comparison groups and it influences the direction of the study results. A high quality study at the level of cohort design will identify the potential confounders and measure them (where possible). This is difficult for studies where behaviora, altifuturial or lifestyle factors	1					
Processor P. Were the outcomes measured in a valid and relatible way? 3 R. Were appropriate E. West appr	8	with confounding factors	the study design or in data analysis. By matching or stratifying sampling of participants, effects of conflouncing factors can be adjusted for. When dealing with adjustment in data analysis, assess the statistics used in the study. Most will be some form of multivariate regression analysis to						
As with any consideration of statistical analysis, consideration should be given to write the time was announced programs of the control of t		measured in a valid and	instruments as this has a significant impact or outcome assessment validly Harring entablished the opicientity of the outcome measurement (e.g., lung cancer)instrument, it's important to establish how the measurement was conducted. Were those involved in collecting data triands or declarated in the use of the instruments'? (e.g., radiographers), if there was more than one data collector, were they similar in terms of level of education, climical or research experience, or level of	0					
112 12 13 14 15 16 17 17 17 17 17 17 17			given to whether there was a more appropriate alternate statistical method that could have been used. The methods section should be detailed enough for reviewers to identify which analytical techniques were used (in particular, ergression or stratification) and how specific confounders were nessured for studies utilizing regression analysis, it is useful to identify if the study dentified which variables were included approach used where the stead of analysis defined by the specified variables. Additionally, it is also inorprated to assess the appropriations of the enneyted service; in terms of the assumptions associated with the approach and offering methods of analysis are based on differing methods of analysis are based on differing approach of analysis are based on differing methods of analysis are based on differing methods.						
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	Land/ culturele context (vergelijkbaar met					~	-	****		
1	NL?)				vs	3				
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6	USA				Japan	1				
7	China				Internat	2				
	UK, Ireland. In									
8	Apri 2020.				Noorwe					
9	Italië				Polen	1				
11	Japan International: The majority currently lives in North America (48.1%), followed by participants in Europe or transcontinental countries with territory in both Europe and Asia Het betreft data				Israel	16			Als zij waren getraceerd door de nationale gezondheidsdienst omdat zij in contact waren geweest met iemand die COVID-19 bleek te hebben, gaf 10.9% aan dat zij gedurende twee weken hun huis niet uit waren geweest. De enige factor die sterk samenhing met niet-naleving was het hebben van een afhankelijk kind in het huishouden. Zelf gegeven redenen om de quarantaine niet na te leven waren: denken dat het niet nodig is om weg te blijven van mensen buiten je eigen huishouden als je niet kan wegblijven van mensen in je eigen huishouden (14.3%), geen symptomen ontwikkelen (11.9%), om boodschappen te doen (10.9%), omdat je net klaar was met een andere quarantaine periode (10.9%). In het algemeen, voor alle uitkomsten, hing niet-naleving samen met man zijn, jongere leeftijd, een afhankelijk kind in het huishouden hebben, lagere socio economische status, het lastiger hebben tijdens de pandemie en in een belangrijke sector werken. Praktische hulp en financiële vergoedingen zullen, verwachten zij, de naleving verhogen.	Factoren geasscoeerd met alle nalevings uitkomsten: lage naleving was geassocieerd met man zijn, jonger zijn, een afhankelijk kind hebben in het huishouden, het moeilijker hebben, lagere socio economische status, minder geinformeerd zijn over covid 19 en informatie over voorkomen verspreiding virus (zoals key symptomen herkennen, niet overheidsbegeleiding weten als je symptomen ontwikkelt, en het niet eens zijn met kans op besmetting als geen symptomen.
12	uit verschillende landen, veel uit UK, maar ook aantal reviews met meerdere studies.									
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