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1.	Quality	of evidence is a continuum; any discrete categorisation involves	some degr	ree of	Table 5.2: Factors that can reduce the	quality of the evidence	
2	arbitrar	iness.			e .	0	r - P-Let
2					Factor	Consequence	toelichting
	While fa	ctors influencing the quality of evidence are additive - such that the	reduction	or increase in	Limitations in study design or execution	1 1 or 2 levels	
3	each indi	vidual factor is added together with the other factors to reduce or in	crease the	quality of	(risk of bias)	¥ = -0.000	zie linksonder Study Design en volgend tabblad voor Risk of Bias.
	evidence	for an outcome - grading the quality of evidence involves judgeme	ents which	are not			
	exclusive	. Therefore, GRADE is not a quantitative system for grading the quantitative system for grading the quantitative system for grading the quantitative system.	ality of ev	idence, Each			
	factor for	downgrading or upgrading reflects not discrete categories but a co	ntinuum w	thin each	Inconsistency of results	1 1 or 2 levels	Niet toegelichte heterogeniteit van resultaten (vooral bij syst reviews, als er veel
4		and among the categories. When the body of evidence is intermedia					verschillende bevindingen zijn, gemengd bewijs).
	particula	factor, the decision about whether a study falls above or below the	threshold	for up- or			
	downgra	ding the quality (by one or more factors) depends on judgment.					Bijvoorbeeld gemeten met een surrogaat maat (niet gedrag, maar intentie of
					Indirectness of evidence	↓ 1 or 2 levels	zelfgerapporteerd gedrag) Of nt andere interventie (niet thuisblijven bij klachten
5							maar thuisbiljven in het algemeen).
-							and the second s
					Imprecision	↓ 1 or 2 levels	Kleine steekproef of kleine hoeveelheid events, dus wijd confidence interval
ь							resultaten niet zijn gepubliceerd en dus niet zijn opgenomen. Vooral voor syst
1	Table 5.1:	Quality of Evidence Grades	l .		Publication bias	1.1 or 2 levels	
7		*y				¥	reviews relevante factor. Bij losse studies gaat het om reporting bias (zijn er
1	Grade	Definition			Table 5.3: Factors that can increase the	quality of the evidence	
8	uuc					quant, or the criticise	
1	TT: 1	We are very confident that the true effect lies close to that of the	I		r.		
9	High	estimate of the effect.	ı		Factor	Consequence	
-		2000 CO 2000 TO 2000 EAR OWN TO THE TOTAL CO.					
1			l				Als er een groot effect wordt gevonden. For simple regression β is like R. Thus I
1		We are moderately confident in the effect estimate: The true effect is	l				would use R rules of thumb I use the following with my Psychology students:
	Moderate	likely to be close to the estimate of the effect, but there is a	l		1 1 6 60	† 1 or 2 levels	β < 0.1 - Small effect size β ∈ [0.1; 0.5] - Medium effect size β ≥ 0.5 - Large
	Moderate		l		Large magnitude of effect	1 or 2 levels	
		possibility that it is substantially different	l				effect size. For multiple regression these rules are not that straightfoward, but for
10			l				Social Sciences they seem to hold (also following Cohen's d suggestions).
10			-				
			l .		The same states and the same states		
		Our confidence in the effect estimate is limited: The true effect may	l .		All plausible confounding would reduce	Address to the control of the contro	
	Low	be substantially different from the estimate of the effect.	l		the demonstrated effect or increase the	† 1 level	Is er gecontroleerd voor plausibele confounders?
			l .		effect if no effect was observed		
11							
		W. I					
	Very Low	We have very little confidence in the effect estimate: The true effect	l .		Dose-response gradient	† 1 level	
12		is likely to be substantially different from the estimate of effect	l .				
13							
14							
15		Study Design					
16		Study design is critical to judgments about the quality	of evidence	<u>.</u>			
17		For recommendations regarding management strategies			accuracy of diagnostic tests -		
18		randomized trials provide, in general, far stronger evidence				ence than uncontrolled case series.	
19		In the GRADE approach to quality of evidence:			,		
20		randomized trials without important limitations provide h	igh quality	evidence			
21		observational studies without special strengths or importa					
22		minous special strengths of importe		, quanty eridence			
23		Limitations or special strengths can, however, modify the	quality of	he evidence of both randomized	trials and observational studies		
24		Note:			The state of the s		
1							
1		N					
1		Non-randomised experimental trials (quasi-RCT) without important					
1		limitations also provide high quality evidence, but will automatically					
		be downgraded for limitations in design (risk of bias) - such as lack					
		of concealment of allocation and tie with a provider (e.g. chart					
25		number).					
1		Case series and case reports are observational studies that investigate					
1		only patients exposed to the intervention. Source of control group					
		results is implicit or unclear, thus, they will usually warrant					
26		downgrading from low to very low quality evidence.					
1		Expert opinion is not a category of quality of evidence. Expert					
1							
1		opinion represents an interpretation of evidence in the context of					
1		experts' experiences and knowledge. Experts may have opinion about					
		evidence that may be based on interpretation of studies ranging from					
1		uncontrolled case series (e.g. observations in expert's own practice)					
		to randomized trials and systematic reviews known to the expert. It is important to describe what type of evidence (whether published or					
2=							
27		unpublished) is being used as the basis for interpretation.					

	A	В	С	D	E	F	G	н	I
				1	Table 5.4:	Study lim	tations in randomized co	trolled trials Explanation	
					ack of allo		cealment	Those enrolling patients are aware of th (or period in a crossover trial) to which enrolled patient will be allocated (a maj problem in "pseudo" or "quasi" random trials with allocation by day of week, bi chart number, etc.).	the next or nized irth date,
1					ack of blir		g of patients and outcome	Patient, caregivers, those recording outc those adjudicating outcomes, or data an are aware of the arm to which patients a allocated (or the medication currently be received in a crossover trial). Loss to follow-up and failure to adhere	alysts are eing
2					events		, o., p	intention-to-treat principle in superiority or in noninferiority trials, loss to follow failure to conduct both analyses conside only those who adhered to treatment, an patients for whom outcome data are a va The significance of particular rates of lc	y trials; -up, and ering nd all nilable. oss to
3	estimate of the effect and in the followin grecommend limitations are, the more likely it is that the quality of ev	scution may bias the estimates of the treatment effect. Ourconfidence in the atton decreases if studies suffer from major limitations. The more serious the idence will be downgraded. Mumerous tools exist to evaluate the risk of bias in dtrials and observational studies						follow-up, however, varies widely and i dependent on the relation between loss i follow-up and number of events. The hi proportion lost to follow-up in relation i intervention and control group event rat differences between intervention and con-	to igher the to tes, and
3					Selective or		orting	groups, the greater the threat of bias. Incomplete or absent reporting of some outcomes and not others on the basis of results. Stopping trial early for benefit. Substan	the
4	Risk of bias	Uitleg			one mine	uous		overestimates are likely in trials with fe than 500 events and that large overestin likely in trials with fewer than 200 even Empirical evidence suggests that formal	wer nates are nts.
5	None							stopping rules do not reduce this bias. Use of unvalidated outcome measures (patient-reported outcomes) Carryover effects in crossover trial Recruitment bias in cluster-randomized	e.g.
	Due to confounding	Baseline confounding occurs when one or more prognostic variables (factors that predict the outcome of interest) also predicts the intervention received at baseline. RDBINS-I can also address time-environ confounding, which occurs when individuals which between the interventions being compared and when post-baseline prognostic factors affect the interventions being down and the past leaves the linetervention received after baseline.		-					
7	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 interventions and outcome even if the effects of the interventions are districted. This form of selection bias I distinct from conformaling—a specific example is bias due to the inclusion of prevalent users, rather than new users, of an intervention.							
	Due to missing data	Bias that arises when later follow-up is missing for individuals initially included and followed (such as differential loss to follow-up that is affected by prognosist factors): bias due to							
8		exclusion of individuals with missing information about intervention status or other variables such as confounders.							
	In measurement of predic/outcome	Bias introduced by either differential or non-differential errors in measurement of outcome datas. 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							
10	In selection of reported result	errors are related to intervention status or effects Selective reporting of results in a way that depends on the findings and prevents the estimate from being included in a meta-analysis (or other synthesis)							
11	In misclassification of intervention (randomization)	Bias introduced by either differential or non-differential misclassification of intervention status. Non-differential misclassification is unrelated to the outcome and will usually bas the estimated effect of intervertion towards the null Differential misclassification cours when misclassification of intervention status is related to the outcome or the risk of the outcome, and is likely to lead to bias. Bas that arises when there are systematic differences between experimental intervention and							
12	Due to deviation from intended intervention	comparator groups in the care provided, which represent a deviation from the intended intervention(s)							
13 14	Tussen studies								
15 16	 Comparison: bij interventie studies, goed be 	klijken wat de comparison conditie is en of studies vergelijkbaar zijn met elkaar.		!					
17 18	- Outcomes: zijn gebruikte uitkomstmaten ve	rgelijkbaar? (gaat het om intentie van gedrag, zelfgrapporteerde naleving, daac	werkelijke	naleving, e	tc)		Table 5.5: Study limitation	ons in observational studies	
19							Failure to develop and	Under- or over-matching in case-	
20							apply appropriate eligibility criteria (inclusion of control population)	Selection of exposed and unexposed	
21								in cohort studies from different populations Differences in measurement of	
23							Flawed measurement of both exposure and outcome	exposure (e.g. recall bias in case-control Differential surveillance for outcome in exposed and unexposed in cohort studies	
24							Failure to adequately	Failure of accurate measurement of all known prognostic factors	
25							control confounding	Failure to match for prognostic factors and/or adjustment in statistical analysis	
26							Incomplete or inadequately short follow-up	Especially within prospective cohort studies, both groups should be followed for the same amount of time.	

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2						1. Were the criteria for inclusion in the sample clearly defined?	The authors should provide clear inclusion and exclusion criteria that they developed prior to recruitment of the study participants.
3						2. Were the study subjects and the setting described in detail?	The study sample should be described in sufficient detail so that other 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 demographics, location, and time period.
4						3.Was the exposure measured in a valid and reliable way?	The study should clearly describe the method of measurement of exposure. Assessing validity requires that a 'gold 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 epidemiological study to check repeatability of measurements of the exposures. These usually include intra-observer reliability and inter-observer reliability.
5						4. Were objective, standard criteria used for measurement of the condition?	It is useful to determine if patients were included in the study based on 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 definitions should provide evidence on matching by key characteristics.
6						5.Were confounding factors identified?	Typical confounders include baseline characteristics, prognostic factors, or concomitant exposures (e.g. smoking). A confounder is a difference between the comparison groups and in fintlences 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 behavioral, attitudinal or lifestyle factors may impact on the results.
7						6.Were strategies to deal with confounding factors stated?	Strategies to deal with effects of confounding factors may be dealt within the study design or in data analysis. By matching or stratifying sampling of participants, effects of confounding factors can be adjusted for. When dealing with adjustment in data analysis, assess the stalistics used in the study. Most will be some form of multivariate regression analysis to account for the confounding factors measured
8						7.Were the outcomes measured in a valid and reliable way?	Importantly, determine if the measurement tools used were validated instruments as this has a significant impact on outcome assessment validity. Having established the objectivity 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 trained or deucated in the use of the instrument's? (e.g. radiographers). If there was more than one data collector, were they similar in terms of level of education, clinical or research experience, or level of responsibility in the piece of research being appraised?
9						8.Was appropriate statistical analysis used?	As with any consideration of statistical analysis, consideration should be 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, regression or stratification) and how specific confounders were measured. For studies utilizing regression analysis, it is useful to identify if the study identified which variables were included and how they related to the outcome. If statification was the analytical approach used were the strata of analysis defined by the specified variables? Additionally, it is also important to assess the appropriateness of the analytical strategy in terms of the assumptions associated with the approach as differing methods of analysis are basedon differing assumptions about the data and how it will respond.
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Sheet1

	Α	В	С	D	E	F	G	Н	1	J
	Land/ culturele									
	context									
	(vergelijkbaar									
1	met NL?)				VS	3				
2	UK	_				4				
	VS	_			UK					
3					Finland	1	_			
4	Polen				China	1	_			
5	Finland				Italië	1				
6	USA				Japan	1				
7	China				Internationaal	2				
	UK, Ireland. In									
8	Apri 2020.				Noorwegen	1				
9	Italië				Polen	1				
10	Japan				Israel	1				
	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 (38.5%) and Australia or New Zealand (5.5%).								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,	Factoren geasscoeerd met alle nalevings uitkomsten: lage naleving was geassocleerd 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.
									verwachten zij, de naleving verhogen.	us geen symptomen.
11						16				
	Het betreft data									
	uit verschillende									
	landen, veel uit									
	UK, maar ook									
	aantal reviews									
	The state of the s									
	met meerdere									
12	studies.									
	UK in begin mei									
13	2020									
14	UK									
15	nvt									
16	Noorwegen									
17	USA.									
18	Israel									