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Measuring multimorbidity in research: Delphi consensus study

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ABSTRACT

OBJECTIVE To develop international consensus on the definition and measurement of multimorbidity in research.

DESIGN Delphi consensus study.

SETTING International consensus; data collected in three online rounds from participants between 30 November 2020 and 18 May 2021.

PARTICIPANTS Professionals interested in multimorbidity and people with long term conditions were recruited to professional and public panels.

RESULTS 150 professional and 25 public participants completed the first survey round.

Response rates for rounds 2/3 were 83%/92% for professionals and 88%/93% in the public panel, respectively. Across both panels, the consensus was that multimorbidity should be defined as two or more long term conditions. Complex multimorbidity was perceived to be a useful concept, but the panels were unable to agree on how to define it. Both

panels agreed that conditions should be included in a multimorbidity measure if they were one or more of the following: currently active; permanent in their effects; requiring current treatment, care, or therapy; requiring surveillance; or relapsing-remitting conditions requiring ongoing care. Consensus was reached for 24 conditions to always include in multimorbidity measures, and 35 conditions to usually include unless a good reason not to existed. Simple counts were preferred for estimating prevalence and examining clustering or trajectories, and weighted measures were preferred for risk adjustment and outcome prediction.

CONCLUSIONS Previous multimorbidity research is limited by inconsistent definitions and approaches to measuring multimorbidity. This Delphi study identifies professional and public panel consensus guidance to facilitate consistency of definition and measurement, and to improve study comparability and reproducibility.

WHAT IS ALREADY KNOWN ON THIS TOPIC

- ⇒ How multimorbidity is defined and measured in research studies varies widely
- ⇒ Previous consensus studies have focused on choice of conditions to include in multimorbidity measures, and have usually involved only local or regional professional panels

WHAT THIS STUDY ADDS

- ⇒ This study provides guidance on how to define and measure multimorbidity in research studies, based on Delphi consensus in professional and public panels; although consensus was reached that multimorbidity should be defined as two or more long term conditions, none was reached on alternative definitions of complex multimorbidity
- ⇒ Panels agreed on which conditions to always include and which to usually include in multimorbidity measurement
- ⇒ Panels also agreed that simple counts of conditions were preferred or considered acceptable for studies estimating prevalence, identifying and counting disease clusters, and exploring trajectories of multimorbidity over time, and that weighted measures were for assessing severity of disease burden, and risk adjustment or outcome prediction

HOW THIS STUDY MIGHT AFFECT RESEARCH, PRACTICE, OR POLICY

- ⇒ The consensus list of conditions to always and usually include in multimorbidity measurement provides a core set for researchers to use to improve comparability and replicability, although researchers can add other conditions relevant to local context and purpose
- ⇒ Consensus about when weighted measures or simple counts were preferred depending on the purpose of an analysis provides a guide to inform researchers choice of methods
- ⇒ Further research is needed to better define and demonstrate the value of concepts such as complex multimorbidity

Introduction

In many regions of the world, a growing proportion of adults has multiple long term conditions or multimorbidity. ^{1–3} Multimorbidity is defined as the coexistence of two or more long term conditions. ⁴ Multimorbidity prevalence increases substantially with age, and is the norm in people aged 65 years or older. ^{5–7} Prevalence is also higher in less affluent and less well educated groups, ^{6–7} with multimorbidity also occurring at younger ages in these groups. ^{1–5} About 30-40% of people with multimorbidity have both a physical and a mental health condition, ^{5–6} with multimorbidity involving a combination of physical and mental health being more common in women, and less affluent and less well educated individuals. ^{5–6}

Despite broad agreement that multimorbidity should be defined as the presence of two or more chronic conditions, no international consensus exists on how to operationalise this broad definition in measures used in research. Multimorbidity measures vary widely in terms of the number, labelling, type, and severity of included conditions or groups of conditions. Without common definitions, many different tools have been developed and used to measure multimorbidity. The tools commonly used in research and clinical practice include: simple (unweighted) disease counts, weighted disease counts, and weighted medication counts. In



addition, many different weighting schemes have been applied to serve different purposes.

Consequently, comparing and reproducing studies is difficult, with for example, large variation in estimates of the prevalence of multimorbidity in different studies, ranging from 3.5% to 100%. The high level of heterogeneity in multimorbidity prevalence has been found to be mainly attributed to age and inconsistent multimorbidity measurement. The estimated pooled prevalence was 68.7% for an oldest population (aged \geq 74 years), 26.3% for a younger population (aged \geq 55 years), 29.3% for a measure including fewer than nine conditions, and 87.6% for a measure including 44 or more conditions.

Previous studies have synthesised existing evidence on multimorbidity measures, 81112 compared the performance of different measures in predicting selected outcomes, 13 14 and adapted existing measures to meet the professionally perceived needs of specific regions or populations. 15 16 These studies have identified heterogeneity in the definition and measurement of multimorbidity as a key issue or limitation, and demonstrate the need for shared approaches to definition to improve comparability and reproducibility. In addition, little attention has been given to directly involving patients and the public in the discussion of multimorbidity definition and measurement. Therefore, this study aimed to explore views and develop consensus on how to measure multimorbidity using a modified Delphi study with an international panel of professionals and the public.

Methods

The overall study design was a modified Delphi method with two international panels of professionals and of members of the public.¹⁷ We used this method as a group consensus strategy to systematically and iteratively explore opinions of professionals and public contributors, and develop consensus on methods of defining and measuring multimorbidity. The study protocol is provided in online supplemental appendix 1.

Data collection methods

Data were collected in three rounds of online questionnaires sent to each individual member of the panels between 30 November 2020 and 18 May 2021. Core questions were the same for both panels, but some more technical questions were only asked of one panel (eg, questions about the acceptability of simple counts or weighted measures for different research purposes were only asked of the professional panel). In the second and third rounds, participants were fed back a summary of all responses to inform their judgments. ^{17 18}

Round 1 questions were informed by the findings of a recent systematic review, ¹⁹ which identified the

characteristics of multimorbidity measures used in research in relation to the study purposes. Each questionnaire included both closed (Likert scaled) questions and open ended questions. Depending on the question, participants were asked to rate (from strongly agree to strongly disagree) or rank (the importance of statements on a scale of 1-5) items or statements using Likert scales.¹⁷ The open ended responses were triangulated with close ended responses, and the results were used to develop new items in the following rounds. Second and third round items were a mix of those scored in the previous round that did not achieve consensus, and new items based on open ended responses in previous rounds. The interactive and repetitive survey rounds, as part of standard Delphi methods, were to improve the framing of the statements for panellists, attest their responses through the iterative process, and achieve consensus. All questionnaires are provided in online supplemental appendix 2.

To conceptualise multimorbidity, eight aspects were explored in the Delphi surveys (online supplemental appendices 2 and 3): the cut-off number of conditions for defining multimorbidity (and complex multimorbidity), duration of a condition for it to be defined as long term, types of conditions to include (eg, medical diagnoses, risk factors, and health behaviours), categorisation of conditions, choice of conditions based on their impact, data sources, which conditions to include (eg, name of individual conditions), and choice of simple counts versus weighted measures for different purposes.

Participants

Participants recruited to the professional panel were clinicians with experience of caring for patients with multiple long term conditions; and researchers and policy makers with an interest in multimorbidity. Participants recruited to the public panel were members of the public with multiple long term conditions or an interest in multimorbidity.

We identified participants using a range of methods: publicly available information including published work, publicly available websites, reports, and policy documents (to identify healthcare professionals, policy makers, or public participants for example, in guideline development). For the public panel, we asked conveners of patient and public involvement groups to forward the invite to their members, and asked participants (and potential participants) to forward study information to others who might meet the criteria, directly or via social media (snowball sampling). No direction on the number of participants is required for a Delphi survey.¹⁷ To provide representative information, some studies have involved more than 60 experts, while others involved as few as 15.18 In this Delphi study, we aimed to recruit a minimum number of experts and public contributors of 25-30, but we had no maximum limit.

Minimising bias and data analysis

We used several techniques to minimise sampling and non-response bias.²⁰ These techniques included sampling expert panellists with different study interests in the field of multimorbidity, using multiple survey distribution methods to increase response rates, highlighting the match between the survey and participant interests, identifying any differences in personal characteristics of those who did or did not complete the surveys, collecting multiple waves of data, and ensuring anonymity among panellists to facilitate open and truthful discussion about their views.

Descriptive statistics were used to describe participants' personal characteristics and responses to statements in three rounds of surveys (including frequency, percentage, median, and interquartile range). Before any data collection, we prespecified consensus as $\geq 70\%$ of panellists providing the same response. ^{17 21}

For items relating to multimorbidity definition, any statements that reached consensus (to "strongly agree," "strongly disagree," "very important," and "not important at all"; rated on a scale of 1-5) in the initial round would not be asked again in the next rounds. If no consensus was reached, then guestions were asked again in the following rounds. If statements did not reach consensus in all rounds, we examined for any consensus in terms of "agree" (the sum of strongly agree and agree), "disagree" (the sum of strongly disagree and disagree), "sufficiently important" (the sum of very important and sufficiently important), or "not important" (the sum of not important at all and slightly important) in the final round (online supplemental figure S1). "Don't know" responses were excluded from the denominator when calculating percentages.

For questions related to the choice of conditions to include in multimorbidity measures, we first identified whether consensus was reached to always include a condition (≥70% agreeing) in multimorbidity measurement. If no consensus was reached, we identified any agreement (≥70%) to usually include unless a good reason to exclude in a particular context (referred to here as "usually include"), defined as the sum of responses to "always include" and "usually include."

For the choice of conditions to include in measures, we included all conditions as "always include" if either panel rated it as "always" and the other rated it as "usually." If one panel rated a condition as "usually include" and the other did not, we used the Rasch dichotomised model as a sensitivity analysis to examine items (conditions) being endorsed (rated always or usually include) and unendorsed (not rated always or usually include) by all participants (online supplemental box 1; this analysis was not prespecified).²² The level of endorsement was estimated on the basis of the item difficulty

parameter in the Rasch model, with negative values representing more frequently endorsed and positive values representing less frequently endorsed.²³ Conditional maximum likelihood estimation in the Rasch analysis was used to produce consistent item parameter estimates without assuming a specific population distribution for the latent trait.²⁴ In the face of disagreement between panels (ie, one panel saying "usually include," the other not), we rated conditions as "usually include" if the item difficulty parameter was ≤0.5.²⁵ All statistical analyses were conducted using R version 4.0.4.

Patient and public involvement

A member of our research team (SS) organised an online meeting with a public reference group in September 2020 to discuss the development and design of the first Delphi questionnaires. Feedback provided by the public reference group included use of simple terms to describe medical diagnosis, and clarity about the difference between multimorbidity and comorbidity and questions relating to weighting. Based on the feedback, we therefore incorporated a short description explaining each medical diagnosis and inserted a two page document introducing the study topic in the online questionnaires. With the support of Health Data Research UK and our colleagues, several members of the public took part in the Delphi study to provide their views on how multimorbidity should be defined and measured. Subsequent round questionnaires were modified in response to comments and suggestions from all panellists including the public. All participants were sent a summary of the findings after completion of data analysis.

Results

In round 1, 150 professional panellists and 25 public panellists took part in the survey (figure 1). Owing to the use of multiple sampling strategies, the response rate in round 1 could not be estimated. The response rates for rounds 2 and 3 in the professional panel were 83% (112/135) and 92% (97/105), respectively, and 86% (n=31/36) and 93% (25/27) in the public panel, respectively. The number of participants in round 2 increased because of snowballing sampling (figure 1). Characteristics of respondents and non-respondents were similar across the three rounds in the professional panel and the public panel (table 1 and online supplemental table S1).

In the professional panel in round 1 (table 1), 53.3% of panellists were from Europe and 20.7% from North America with smaller proportions from Australasia (8.7%), Asia (13.3%), South America (3.3%), and Africa (0.6%). Most professional panellists were interested in multimorbidity in the general population or in middle aged or older adults, but only 12.7% were interested in multimorbidity in children.

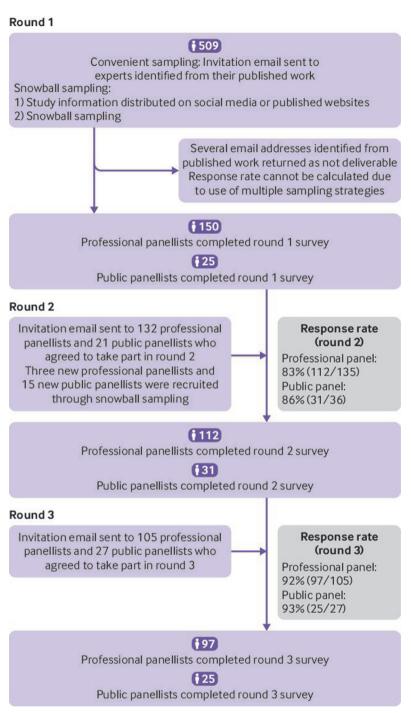


Figure 1 | Process of participant recruitment

More than half of professional panellists were interested in multimorbidity in socially deprived populations (56.7%), and 38.0% in multimorbidity in ethnic minority and indigenous groups. In the public panel, most panellists were from Europe, with fewer than 4% from Asia, North America, or South America. Just over half of public panellists were women (56.0%), and 48.0% of the public panellists were aged 65 years and older. The proportions of participant characteristics were similar across rounds.

Both panels agreed that multimorbidity should be defined as the co-occurrence of two or more long term

conditions. Defining complex multimorbidity was considered useful by more than 80% of both panels, with consensus in the public panel that complex multimorbidity could be defined as the co-occurrence of three or more long term conditions. However, no consensus in the professional panel was reached on how to define complex multimorbidity with variation in whether three or more conditions had to come from any, at least two, or at least three body systems. Neither panel agreed on the value of any other patterns of complex multimorbidity, with physical-mental comorbidity chosen by 33% of professional

	easurement			
Professional panellists Public panellists				
Round 2 (n=31)	Round 3 (n=25)			
30 (96.8)	24 (96.0)			
0	0			
0	0			
0	0			
1 (3.2)	1 (4.0)			
0	0			
0	0			
30 (96.8)	24 (96.0)			
1 (3.2)	1 (4.0)			
0	0			
19 (61.3)	17 (68.0)			
12 (38.7)	8 (32.0)			
0	0			
24 (77.4)	20 (80.0)			
7 (22.6)	5 (20.0)			
0	0			
_	_			
_	_			
_	_			
_	_			
_	_			
_	_			
_	_			
_	_			
_	_			
_	_			
_	_			
_	_			
_	_			
_	_			
_	_			
_	_			
_				
_	_			
1 (3.2)	1 (4.0)			
	5 (20.0)			
	6 (24.0)			
	13 (52.0)			
10 (71.0)	17 (72.0)			
19 (61 3)	14 (56.0)			
	14 (56.0)			
	- (3.2) 7 (22.6) 7 (22.6) 16 (51.6) 19 (61.3) 12 (38.7)			

*Participants could choose more than one response so percentages can sum to >100%.

†Female and male refers to the sex of the public panellists; women and men refer to the populations that the professional panellists have research interests

panellists and 44% of public panellists, physical functional limitations by 30.9% of professional panellists and 32% of public panellists, difficulties

in managing illness due to social factors by 26.8% of professional panellists and 28% of public panellists, and frailty by 25.8% of professional panellists and

Data are number (%) of participants.

12% of public panellists (online supplemental table S2).

Conditions were considered to be long term if they persisted for six months or more in the professional panel (70.5%); conditions were considered long term if they lasted 12 months or more in the public panel (76.0%). More than 95% of panellists from both panels would include formal medical diagnoses in multimorbidity measurement. While the public panel agreed that clinical risk factors were important for multimorbidity measurement (74.2%) (online supplemental table S2), the professional panel did not reach a consensus. Symptoms, health behaviour, health impacts, social deprivation, and consequences of treatment did not reach consensus in both panels as conditions to include for measurement. Both panels agreed that conditions should be included in a multimorbidity measure if they were any of the following: currently active; permanent in their effects; requiring current treatment, care, or therapy; requiring surveillance (including treated cancers that require surveillance); or relapsing-remitting conditions that require ongoing treatment, care, or therapy (online supplemental table S3). On the other hand, no consensus was reached on the conditions that might recur or remit but happen rarely and that usually require treatment or therapy at some point in the future even if not currently treated. Both panels reached consensus that studies should count individual conditions rather than categories defined by body system, and that disease complications should be counted separately from diseases (eg, peripheral neuropathy and diabetes). The public panel (but not the professional panel) agreed that individual cancers should be counted separately (table 2 and online supplemental table S2).

In respect to criteria for selecting conditions based on impact, more than 70% of both panels agreed that conditions were appropriate to include in multimorbidity measurement if they were any of the following: significantly reduce quality of life, significantly worsen mental health, significantly increase risk of death, cause frailty, cause physical disability, or significantly increase treatment burden. The professional panel (but not the public panel) reached consensus on including conditions that significantly worsen self-perceived health status. The public panel (but not the professional panel) reached consensus on including conditions that are affected by social deprivation and poverty (table 2 and online supplemental table S4). Both panels agreed that conditions included for measurement should be similar in self-report, administrative databases, and medical records.

Technical questions about the use of simple counts versus weighted measures based on study purposes were only asked in the professional panel. In round 1, no consensus was reached on whether simple counts or weighted measures were

generally preferable (online supplemental table S5). In rounds 2 and 3, for a range of different purposes, professionals were asked if they preferred simple counts or weighted measures or if either was acceptable. There was no consensus that one or other type of measure was preferred for any of the purposes asked, but for all but one purpose, there was clear consensus that one type of measure was preferred or acceptable (table 2 and online supplemental table S6). Simple counts were preferred or acceptable for estimating the prevalence of multimorbidity, identifying and counting disease clusters, and exploring trajectories of multimorbidity. Weighted measures were preferred or acceptable for assessing the severity of disease burden, risk adjustment, and outcome prediction (in general) and for every specific outcome asked about (online supplemental table S7). No consensus was reached on the best type of measure for exploring or identifying predictors of multimorbidity. In round 2, 21.7% (n=20) of panellists preferred to use weighted indices, 46.7% (n=43) preferred to empirically derive weights based on the individual impact of diseases on outcome (eg, regression models to calculate weights), and 26.1% (n=24) preferred to set rules based on level of severity to grade each condition (eg, having presence of a condition=1 point, treatment=additional 1 points, functional limitation=additional 1 point). In both professional and public panels, mortality, healthcare use, health related quality of life, physical disability, and frailty were rated as sufficiently important or very important to weight against by ≥70% panellists if weighted measures were preferred.

Of the 107 individual conditions asked about in the Delphi questionnaires (online supplemental file 2), 24 were rated as "always include" in multimorbidity measurement (the 107 conditions were defined on the basis of results of a recent systematic review19 and panellists' suggestions in initial rounds). This "always include" list consisted of 16 conditions (table 3) that reached consensus in both professional and public panels (end stage kidney disease, heart failure, dementia, chronic liver disease, chronic kidney disease, stroke, solid organ cancers, metastatic cancers, haematological cancers, multiple sclerosis, Parkinson's disease, coronary artery disease, cystic fibrosis, epilepsy, diabetes, and HIV/AIDS), seven conditions reaching consensus in the professional (but not public) panel (chronic obstructive pulmonary disease, inflammatory bowel disease, connective tissue disease, paralysis, schizophrenia, peripheral artery disease, and asthma), and one condition reaching consensus in the public (but not professional) panel (Addison's disease; online supplemental tables S8 and S9).

Table 2 | Responses to questions relevant to definitions of multimorbidity and complex multimorbidity. Data are percentage of panellists agreeing (and Delphi survey round (R))

Question or statement	Professional panellists	Public panellists
Definition of multimorbidity		
Multimorbidity is two or more long term conditions	84.8 (R2)	88.0 (R1)
Complex multimorbidity is a useful idea	87.5 (R2)	84.0 (R2)
Complex multimorbidity is three or more long term conditions	No consensus	76.0 (R3)
Types of conditions to include		
Long term means present for six months or more	70.5 (R2)	No consensus
Long term means present for 12 months or more	No consensus	76.0 (R1)
Medical diagnoses	99.1 (R2)	96.8 (R1)
Clinical risk factors	No consensus	74.2 (R2)
Currently active	98.7 (R1)	93.5 (R2)
Permanent in their effects	98.6 (R1)	96.0 (R1)
Requiring current treatment, care, or therapy	100.0 (R2)	96.8 (R2)
Requiring surveillance	74.7 (R3)	88.0 (R3)
Remitting-relapsing conditions requiring ongoing treatment or care	93.8 (R3)	92.0 (R3)
Counting or categorisation		
Count individual conditions not broad disease categories	72.0 (R1)	88.0 (R1)
Count individual cancers separately	No consensus	76.0 (R1)
Criteria for selecting conditions relating to impact		
Significantly increase risk of death	94.6 (R2)	100 (R1)
Significantly reduce quality of life	96.6 (R1)	93.5 (R2)
Cause frailty	89.9 (R2)	90.3 (R2)
Cause physical disability	93.3 (R1)	96.8 (R2)
Significantly worsen mental health	92.6 (R1)	87.1 (R2)
Significantly worsen self-perceived health status	77.4 (R2)	No consensus
Significantly increase treatment burden	87.4 (R2)	87.1 (R2)
Impacted by social deprivation and poverty	No consensus	74.2 (R2)
Data source		
Conditions should be the same/similar in both self-report and clinical/administrative database studies	71.8 (R2)	96.0 (R2)
Purposes where a simple count preferred or acceptable*		
Estimating prevalence	83.7 (R3)	Not asked
Identifying and counting disease clusters	80.2 (R3)	Not asked
Exploring trajectories of multimorbidity	72.7 (R3)	Not asked
Purposes where weighted measure preferred or acceptable*		
Assessing severity of disease burden	94.5 (R3)	Not asked
Risk adjustment or outcome prediction	91.2 (R3)	Not asked
Outcomes important to weight against		
Death	92.8 (R1)	96.8 (R2)
Healthcare use	83.7 (R1)	90.3 (R2)
Health related quality of life	92.3 (R1)	90.3 (R2)
Physical disability	87.8 (R1)	86.7 (R2)
Frailty	86.3 (R1)	76.7 (R2)

Absolute numbers for percentage data are as follows: professional panel, round 1 n=150, round 2 n=112, round 3 n=97; public panel, round 1 n=25, round 2 n=31, round 3 n=25.

*Panellists could either state that they preferred a simple or weighted measure for the listed options, or that a simple or weighted measure were both acceptable; values are the sum of "preferred" or "acceptable."

Of 37 conditions rated to usually include unless a good reason to exclude in a particular context, 34 reached consensus in both panels (table 3, online supplemental tables S10 and S11). Of the 22 conditions that reached consensus to usually include in only one panel, three conditions (treated hypertension, gout, and anxiety) had an estimated difficulty parameter <0.5, and were therefore considered to be in the "usually include"

list (online supplemental table S12). Twenty seven conditions did not reach consensus to include in either panel, but no condition was rated as "usually exclude" or "always exclude" (online supplemental table S12).

Endorsement did not vary by participant characteristics apart from attention deficit hyperactivity disorder, which did not reach consensus in both panels, but was substantially more endorsed by

Table 3 | Conditions with consensus to always include and usually include unless there is a specific reason not to in a multimorbidity measure, by panel, based on Delphi surveys. Data are percentage of panellists agreeing (and Delphi survey round (R)) unless stated otherwise

Always include condition		Usually include condition*		Difficulty	
Condition	Professional panellists	Public panellists	Professional panellists	Public panellists	parameter estimate in both panels (logit)†
Heart failure	90.0 (R1)	83.9 (R2)	_	_	-3.1
Chronic liver disease	88.5 (R1)	80.6 (R2)	_	_	-3.5
Diabetes	87.3 (R1)	71.0 (R2)	_	_	-3.5
Parkinson's disease	86.6 (R1)	77.4 (R2)			-2.8
End stage kidney disease	86.4 (R1)	90.3 (R2)	_	_	-2.0
Coronary artery disease	82.7 (R1)	74.2 (R2)	_	_	-2.6
Dementia	82.6 (R1)	83.3 (R2)	_	_	-2.3
Multiple sclerosis	80.7 (R1)	77.4 (R2)		_	-1.9
Stroke	80.0 (R1)	80.6 (R2)		_	-2.6
Chronic kidney disease	79.3 (R1)	80.6 (R2)		_	-2.8
HIV/AIDS	78.5 (R1)	71.0 (R2)			-1.5
Metastatic cancers	77.4 (R1)	70.8 (R1)			-1.3
Haematological cancers	77.2 (R1)	70.8 (R1)			-1.9
Solid organ cancers	76.5 (R1)	70.8 (R1)	_	_	-2.0
Cystic fibrosis	75.8 (R1)	74.2 (R2)		_	-1.3
Epilepsy	73.0 (R1)	74.2 (R2) 71.0 (R2)		_	-1.3 -2.2
Chronic obstructive pulmo-		No consensus	-	96.8 (R2)	
nary disease					-3.1
Inflammatory bowel disease	82.6 (R1)	No consensus	-	100 (R2)	-1.9
Connective tissue disease	79.7 (R1)	No consensus	_	93.3 (R2)	-2.3
Paralysis (other than stroke)	76.0 (R1)	No consensus	_	93.3 (R2)	-0.9
Schizophrenia	75.2 (R1)	No consensus	_	93.5 (R2)	-1.6
Peripheral arterial disease	71.1 (R1)	No consensus	_	96.8 (R2)	-1.6
Asthma	70.7 (R1)	No consensus	_	80.6 (R2)	-1.1
Addison's disease	No consensus	70.8 (R2)	86.9 (R2)	_	-0.8
Depression	_	_	92.9 (R2)	77.4 (R2)	-0.9
Heart valve disorders	_	_	92.0 (R2)	100 (R2)	-1.6
Bipolar disorder	_	_	90.0 (R2)	93.5 (R2)	-1.1
Melanoma	_	_	88.2 (R2)	100 (R2)	-1.1
Bronchiectasis	_	_	86.7 (R3)	88.0 (R3)	-0.7
Osteoarthritis	_	_	84.7 (R2)	87.1 (R2)	-0.5
Pancreatic disease	-	_	84.4 (R2)	96.7 (R2)	-0.7
Arrhythmia	_	_	83.9 (R2)	85.7 (R2)	-0.4
Thyroid disorders	_	_	82.7 (R2)	87.1 (R2)	-0.3
Venous thrombotic disease	_	_	82.4 (R2)	96.8 (R2)	-0.5
Drug or alcohol misuse	_	_	81.8 (R2)	74.2 (R2)	-0.02
Anaemia	-	_	81.7 (R2)	96.7 (R2)	-0.4
Chronic Lyme disease	=	-	81.3 (R3)	79.2 (R3)	-0.03
Transient ischaemic attack	_	=	80.4 (R2)	96.8 (R2)	-0.4
Treated cancer requiring surveillance	_	_	79.3 (R3)	80.0 (R3)	0.01
Eating disorders	-	-	79.1 (R2)	74.2 (R2)	0.2
Vision impairment that cannot be corrected	_	=	78.6 (R2)	74.2 (R2)	0.1
Long term musculoskeletal problems due to injury	-	-	78.4 (R3)	70.8 (R3)	0.2
Tuberculosis	-	_	82.4 (R2)	90.3 (R2)	0.07
Endometriosis	=	-	75.7 (R2)	89.3 (R2)	0.1
Chronic primary pain	_	_	75.3 (R3)	80.0 (R3)	0.2
Hearing impairment that cannot be corrected	_	=	73.9 (R2)	74.2 (R2)	0.4

Continued

Table 3 Continued					
	Always include conditio	n	Usually include condition	n*	Difficulty
Condition	Professional panellists	Public panellists	Professional panellists	Public panellists	parameter estimate in both panels (logit)†
Peptic ulcer	_	_	73.9 (R2)	83.9 (R2)	0.3
Post-traumatic stress disorder	_	_	73.4 (R2)	74.2 (R2)	0.5
Post-acute covid-19	_	_	73.4 (R3)	92.0 (R3)	0.2
Benign cerebral tumours	_	_	73.3 (R3)	76.0 (R3)	0.4
Peripheral neuropathy	_	_	73.1 (R2)	96.7 (R2)	0.1
Hypertension (untreated)	_	_	73.0 (R2)	71.0 (R2)	0.4
Congenital disease and chromosomal abnormalities	_	_	72.6 (R2)	90.0 (R2)	0.2
Chronic urinary tract infection	_	_	71.8 (R2)	86.7 (R2)	0.3
Aneurysm	_	_	71.6 (R3)	96.8 (R2)	0.5
Meniere's disease	_	_	71.3 (R2)	71.0 (R2)	0.5
Osteoporosis	_	_	70.3 (R2)	80.6 (R2)	0.5
Autism	_	_	70.1 (R2)	87.1 (R2)	0.5
Hypertension (treated)	_	_	80.4 (R2)	No consensus	0.3
Anxiety	_	_	80.0 (R2)	No consensus	0.4
Gout	_	_	76.1 (R2)	No consensus	0.4

*Consensus of conditions to usually include is defined as more than 70% of panellists rated conditions as always include or usually include. †Used to examine endorsement (see also online supplemental table S14).

professional panellists interested in multimorbidity in children than those who were not (online supplemental table S13).

Discussion

Principal findings

Figure 2 and figure 3 summarise the research and reporting recommendations, and table 4 lists the conditions recommended for inclusion in multimorbidity measures. This consensus study found that more than 70% of professional and public panellists defined multimorbidity as the co-occurrence of two or more long term conditions. Despite consensus that complex multimorbidity was a useful concept in addition to this, no consensus was reached on how best to define it. Twenty four conditions were rated as ones to "always include," and 37 to "usually include (unless a good reason to exclude in a particular context)." Of the 37 conditions to usually include, untreated and treated hypertension were combined, and conditions that require surveillance has been generally agreed to be included for multimorbidity measurement (criteria for types of conditions to include) and thus treated cancer requiring surveillance was not particularly included in the recommended list of conditions, leading to 35 conditions recommended to usually include in multimorbidity measurement (table 4).

No conditions were rated by either panel to always exclude or usually exclude, consistent with allowing researchers to choose to additionally include other conditions of particular importance in their context.

General criteria reaching consensus in both panels on reasons to select and include conditions in multimorbidity measurement (which could inform such choices) were that a condition was one or more of the following: medical diagnosis; conditions that are currently active; conditions that are permanent in their effects; conditions that require current treatment, care, or therapy; conditions that require surveillance; and remitting-relapsing conditions that require ongoing treatment or care.

Professional and public panels disagreed on how long a condition should persist to be defined as long term, with consensus in the professional panel on ≥6 months versus consensus in the public panel on ≥12 months. Our judgment was to recommend the 12 month cut-off period, but the discrepancy means that other researchers might decide to use a six month cutoff period. Health impacts agreed by both panels as important consideration in the choice of conditions included risk of death, quality of life, frailty, mental health, and treatment burden. As data could be collected from different sources, the consensus was that a consistent approach to multimorbidity measurement should be adopted, irrespective of whether the study used routine data (from patient records or insurance claims databases) or patient self-report. In this study, we found that panellists chose the type of multimorbidity measures depending on study purposes.

Simple counts of conditions were preferred or considered acceptable for estimating prevalence, identifying disease clusters, and exploring

Types of conditions to include Multimorbidity measure design Definition If including other conditions, If including other conditions, then How to count conditions then consider if they are: consider those with high impact Multimorbidity Count individual conditions 1) Lasting ≥12 months 1) Increase risk of death Co-occurrence of two or rather than categories 2) Medical diagnoses 2) Reduce quality of life more long term conditions defined by body system 3) Currently active 3) Cause frailty Complex multimorbidity 4) Permanent in effect 4) Cause physical disability Consensus that it was a useful 5) Requiring current treatment, 5) Worsen mental health concept, but no clear consensus 6) Increase treatment burden care, or therapy on how to measure it 6) Requiring surveillance Choice of simple counts or weighted measures Data source 7) Remitting-relapsing conditions requiring ongoing treatment or Conditions chosen for measurement Simple counts preferred or acceptable care should be consistent in both self report 1) Estimating prevalence and clinical or administrative studies 2) Identifying and counting clusters 3) Exploring trajectories Weighted measures preferred or acceptable 1) Assessing severity of disease burden 2) Risk adjustment 3) Outcome prediction

Figure 2 | Summary of findings and recommendations on multimorbidity definition. Professional panel consensus was >6 months; patient panel consensus was >12 months

trajectories of multimorbidity, whereas weighted measures were preferred or considered acceptable for assessing disease severity and predicting outcomes. No consensus was reached on how to weight measures, consistent with this depending on study purpose, but researchers should therefore explicitly state and justify their choice of how to weight (eg, in relation to severity of disease or in relation to a particular outcome). Stirland et al²⁶ provide guidance on which weighted measures to use for a particular purpose for those researchers who judge that a weighted measure is appropriate.²⁶

Strengths and limitations of the study

Strengths of this study include that the surveys were designed on the basis of results of a systematic review and in response to panellists' input, and that participants were recruited to both professional and public panels with good retention. Limitations include that less than 20% of panellists were from low or middle income countries, meaning that long term conditions prevalent in low or middle income countries might not have been prioritised. The professional panel was also larger than the public panel, meaning that where panels disagreed in which conditions to include, analysis could have favoured the professional perspective. An implication is that the conditions recommended for inclusion are probably best seen as a core list, and that researchers should carefully consider any additional conditions in their context to be included, and ensure public and patient involvement in their choice. However, if reporting prevalence of multimorbidity, then reporting the prevalence using the core list is recommended to improve comparability as well as reporting prevalence using the study specific set of conditions.

Secondly, owing to the difficulty of navigating experts in this relatively new research specialty

of multimorbidity, the study results might have differed if those interested in multimorbidity but never involved in multimorbidity research had been included. Finally, the professional and public panels disagreed on a small number of areas, meaning that findings should be interpreted with caution. Future studies could explore these areas of disagreement in more depth than is possible in a Delphi study. More in-depth studies could also explore more technical questions that were not asked of the public panel in this study (eg, relating to the construction of weighted measures).

$Comparison\ of\ results\ with\ previous\ studies$

Several previous consensus studies and group developed position papers have focused on the definition of multimorbidity, but these typically do not consider how to apply these definitions in measurement.²⁷ ²⁸ Other studies have highlighted variable measurement of multimorbidity, with large variation in the number and nature of conditions included in measures. 19 29 30 Prior consensus studies have examined which conditions to include. N'Goran et al³¹ used a modified RAND consensus method with a Swiss family practitioner panel to identify 75 International Classification of Primary Care diagnoses pertinent to the clinical consideration of people with multimorbidity. The main differences with this study were their inclusion of a more heterogeneous set of conditions in the psychological domain (including tobacco abuse and memory disturbance that is not dementia).31

Hafezparast et al³² aimed to identify local consensus on the choice of conditions to include in a measure relevant to inner city London.³² Unspecified participants were asked to rate 86 conditions identified in a scoping review, considering them in terms of their prevalence, impact, preventability and

All multimorbidity studies should report

- Their core multimorbidity definition and whether measured by simple count or weighted index
- 2) Which conditions are included in the measure
- 3) Why these conditions were selected in relation to purpose of study, including justifying exclusion of recommended conditions, and inclusion of other conditions
- 4) How each condition has been defined, including any clinical code sets used

Report where appropriate

- If measuring complex multimorbidity, then report definition used and justify it
- 2) If measuring prevalence, then report prevalence of "always include" or "always and usually" conditions as well as study specific measure prevalence to allow direct comparison
- If measuring prevalence, then report prevalence stratified by age, sex, ethnicity group and socioeconomic status to improve comparability
- 4) If using weighted measures to predict outcomes, then ideally compare predictive performance with that of simple count of same conditions

Figure 3 | Reporting recommendations on multimorbidity

modifiability, treatment burden, disease progression, and data quality. Thirty two conditions were rated as locally important to include in multimorbidity measurement, of which only two were not rated as always or usually include in our study (learning difficulties and morbid obesity). In addition, a qualitative study by Drye et al³³ identified 10 chronic conditions for quality care measurement (based on their adverse

effects on health status, function, and quality of life), all of which were included in the core list of this study. However, several conditions rated as "always or usually include" in our study were not in Drye's recommendations, such as cancers, schizophrenia, and chronic liver disease. ³³

As previous review has shown that more than half of existing studies did not include mental health conditions in measurement, 19 the nine mental health conditions rated as "always or usually include" could provide more comprehensive quality measurement for individuals with multimorbidity. Others have noted that the exact choice of conditions is likely to vary by study purpose, that episodic conditions should be included, and that there might be patient characteristics which are very important in clinical care (eg, smoking or socioeconomic status).^{29 30} In line with previous studies, we found consensus on the inclusion of episodic conditions only if they are active, permanent in their effects, or require ongoing treatment or surveillance; but we found no consensus on patient characteristics and social factors in both panels.

Implications of results

This study has several implications. Firstly, while we recognise that the choice of conditions to include in measurement should be sensitive to purpose and local context, ³⁰ research in the field would be

Body system (based on ICD-10 chapters)	Always include (n=24)	Usually include (unless a good reason not to in a particular context) (n=35)*
Cardiovascular disease	Stroke, coronary artery disease, heart failure, peripheral artery disease	Heart valve disorders, arrhythmia, venous thromboembolic disease, aneurysm, nypertension (treated and untreated)
Metabolic and endocrine disease	Diabetes, Addison's disease, cystic fibrosis	Thyroid disorders
Respiratory disease	Chronic obstructive pulmonary disease, asthma	Bronchiectasis
Neurological disease	Parkinson's disease, epilepsy, multiple sclerosis, paralysis	Transient ischaemic attack, peripheral neuropathy, chronic primary pain
Cancer	Solid organ cancers, haematological cancers, metastatic cancers	Melanoma, benign cerebral tumours that can cause disability
Mental and behavioural disorder	Dementia, schizophrenia	Depression, anxiety, bipolar disorder, drug or alcohol misuse eating disorder, autism, post-traumatic stress disorder
Musculoskeletal disease	Connective tissue disease	Osteoarthritis, long term musculoskeletal problems due to injury, osteoporosis, gout
Digestive disease	Chronic liver disease, inflammatory bowel disease	Chronic pancreatic disease, peptic ulcer
Urogenital disorder	Chronic kidney disease, end stage kidney disease	Endometriosis, chronic urinary tract infection
Haematological disorder	_	Anaemia (including pernicious anaemia, sickle cell anaemia)
Eye disease	_	Vision impairment that cannot be corrected
Ear disease	_	Hearing impairment that cannot be corrected, Meniere's disease
Infectious disease	HIV/AIDS	Chronic Lyme disease, tuberculosis, post-acute covid-19
Congenital disease	_	Congenital disease and chromosomal abnormalities

ICD-10=international classification of diseases, 10th revision.

*Untreated and treated hypertension were combined. Conditions that require surveillance (including cancers) were agreed to be included by both sets of panellists, and thus "cancers that require surveillance" was not stated separately in the list.

improved if researchers used a common set of conditions as core, which is provided in the list of conditions to always and usually include be identified in this study (table 4). For studies of prevalence, we recommend that researchers also report age and sex stratified prevalence based on the "always include" and "always or usually include" lists to improve comparability of studies. 10 More generally, although not the focus of this study, multimorbidity measures are often poorly reported, and clarity about choices made and their rationale is critical (figure 3).¹⁹ We recommend that selection of other long term conditions in measures should take account of the criteria agreed as important by panellists in this study (figure 2), and that researchers explicitly report why and how they make decisions on condition and measurement choice (figure 3).

Secondly, this study has identified a need for consistent use of validated clinical code lists, but did not seek to identify them. Others have published lists of such codes for use in this context, ³⁴ and with several initiatives set up to standardise identification of conditions in healthcare data (eg, the Health Data Research UK Phenotype Library ³⁵).

Thirdly, although others have said that weighted measures are generally preferred over simple counts, ³⁶ this study provides professional consensus about the particular purposes where simple counts or weighted measures were preferred or considered acceptable (figure 2). However, we need research that considers the relative performance of simple counts and weighted measures (eg, in predicting outcomes), and for wider public discussion about the relevance of weighted measures to patients (eg, in relation to which outcomes measures are weighted against).

Finally, our study found consensus that complex multimorbidity was a useful concept but no clear consensus on how to define it. Researchers who adopt definitions of multimorbidity beyond two or more conditions should therefore clearly justify their choice (figure 3). Research is needed to better understand the experience of complex multimorbidity from a patient perspective, and to examine whether different definitions of complex multimorbidity have better predictive performance than existing measures. We recommend that complex multimorbidity definitions should be co-developed with patients to ensure that these are relevant to their illness experience.

In conclusion, existing measurement of multimorbidity is highly inconsistent. The findings of this Delphi study provide guidance on multimorbidity measurement that will help bring greater consistency to the field, facilitating replication, comparison between studies, and evidence synthesis.

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Ethics approval This study involves human participants and was approved by the University of Edinburgh Usher Institute's research ethics committee (reference 2113_A2). Participants gave informed consent before participating in the study.

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Measuring multimorbidity in research: a Delphi consensus study

Appendix 1: Study protocol





Non-CTIMP Study Protocol

A Delphi study to explore an international consensus on the definition and measurement of multimorbidity

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Protocol authors	Iris Ho and Bruce Guthrie
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Chief Investigator	Professor Bruce Guthrie
Sponsor number	CFC 0110
REC Number	
Project registration	If applicable trials should be registered on a publically accessible database. ACCORD can provide log-in credentials for clinicaltrials.gov. Please email resgov@accord.scot to arrange
Version Number and Date	Version 1: 24/02/2020





Version 1: 24/February/2020

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LIST OF ABBREVIATIONS

ACCORD	Academic and Clinical Central Office for Research & Development - Joint office for The University of Edinburgh and Lothian Health Board	
CI	Chief Investigator	
CRF	Case Report Form	
GCP Good Clinical Practice		
ICH	International Conference on Harmonisation	
PI	PI Principal Investigator	
QA	Quality Assurance	
REC Research Ethics Committee		
SOP Standard Operating Procedure		
СоСоРор	Condition, Context, Population	
BOS	Bristol Online Survey	

1 INTRODUCTION

1.1 BACKGROUND

In many regions of the world, a growing proportion of the adult population is affected by more than one chronic condition [1-3]. Evidence from several studies indicates that the prevalence of multimorbidity increases substantially with age, and commonly occurs in people aged 65 or older [4-6]. Prevalence is also inversely related to socio-economic status and educational attainment [5, 6]. People living in deprived areas and with lower education tend to be multimorbid at a younger age [1, 4]. Of the population with multimorbidity, approximately 30% to 40% have both a physical and a mental health condition [4, 5]. Women and people with lower education and living in deprived areas are more likely to have physical and mental comorbidity [4, 5].

Multimorbidity is defined as the co-existence of multiple chronic conditions [7].. Unlike comorbidity that focuses on the effects of additional conditions in reference to an index chronic condition, multimorbidity addresses the total effects of chronic conditions without giving priority to any one of the co-existing conditions [8]. In reference to their difference in definition, multimorbidity has, therefore, been placed as a separate Medical Subject Heading (MeSH) since January 2018, distinct from comorbidity [8]. Notwithstanding some agreement on the broad definition of multimorbidity, there remains no international consensus on its

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operational definition regarding how to measure it, with the measures varying widely in terms of the number, labelling, type, and severity of included conditions [7].

Without a clear and agreed guide, many measurement tools have been developed and used to measure multimorbidity. The measurement tools used in research and practice include: unweighted disease counts; weighted disease counts; weighted medication counts [9]. In addition, different weighting schemes have been applied to serve different purposes. Consequently, there is a wide discrepancy in prevalence estimates, ranging from 12.9%-95.1% in the general population [3].

Existing research recognises the importance of establishing an agreed approach to the definition and measurement of multimorbidity. Several attempts have been made to synthesise existing evidence on multimorbidity measures [9-11], to compare different measures of multimorbidity to predict certain outcomes [12], and to adapt existing measures to meet the needs of specific regions or populations [13, 14]. Due to the heterogeneity of current approaches, continued efforts are needed to develop an agreed approach as to what constitutes multimorbidity, what should be included in multimorbidity measures, and how to measure it. The aim of this study is to use the Delphi technique to explore international experts' views on the definition and measurement of multimorbidity, and to further provide a comprehensive guide on the use of multimorbidity measures to meet different purposes.

1.2 RATIONALE FOR STUDY

The Delphi technique is an iterative and participatory method to explore experts' opinions, discuss issues and build consensus through a structured group communication process [15]. In the current evidence base, this method has been used in health research for needs assessment [16], policy determination [17], and guideline development [18]. Given the wide range of applications of multimorbidity definition and measures, the Delphi method is considered as a suitable method for this study to collaboratively engage international experts to address the complex issues and support decision-making on how to measure multimorbidity.

2 STUDY OBJECTIVES

2.1 OBJECTIVES

- Objective 1: To identify and summarise published multimorbidity definitions and measures used in the existing literature, and to explore how these vary by the stated purpose of each study through a systematic scoping review
- Objective 2: To develop consensus construction to provide a comprehensive guidance on multimorbidity definition and measurement

3 STUDY DESIGN

This study involves two phases. The first phase is a systematic scoping review to address objective 1, and the second phase is a Delphi study to address objective 2.

3.1 PHASE ONE: A SYSTEMATIC SCOPING REVIEW

We will use a systematic scoping review to examine the broad area of multimorbidity to map key concepts and identify gaps in the evidence [19]. In this review, we will follow the

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CoCoPop framework (Condition, Context and Population) to define eligibility criteria and conduct searches. This framework was developed by Munn et al. (2015) for systematic reviews of observational studies [20]. "Condition" is referred to as the variable of interest, which is multimorbidity in this review. In terms of context and population, we will look at studies conducted in hospitals, primary care and community that a population sample of people is being studied. Please see table 1 for the inclusion and exclusion criteria.

Table 1. Inclusion and exclusion criteria

Inclusion criteria:

- Quantitative studies that measured multimorbidity in a defined population
- In any setting, including hospitals, primary care, community
- Studies reported in English

Exclusion criteria:

- Studies that used measures with the presence of an index disease integral to the measures
- Conference proceedings or articles that do not provide full text
- Qualitative research and case series

3.1.1 Search strategy

The search strategy for this review will be developed in collaboration with a medical librarian (Appendix 1). Two sets of key terms (multimorbidity and measure) will be combined to search relevant literature by a means of Boolean logic. Medical subject headings will be used to capture concepts and to see if it yields additional studies in comparison with the results from keyword search. Searches will be conducted in Ovid interface (PsycINFO, Embase, Global Health, Medline), Scopus, Web of Science, Cochrane Library, EBSCO interface (CINAHL Plus), and ProQuest Dissertations & Theses Global, from inception to 21 January, 2020. Following the database searches, reference lists of retrieved articles will be hand-searched and citations will be tracked using Scopus/Google Scholar to ensure the inclusion of all relevant literature.

Conference proceedings and literature that did not provide full text will not be included in the review due to the limited information available for data extraction and analysis.

3.1.2 Study screening

References identified by the search strategy will be exported to EndNote X9 bibliographic software and Excel for deduplication, and then will be imported to Covidence for screening. Titles, abstracts, and full-texts of retrieved articles will be screened against the eligibility criteria by two reviewers (SH and PH). The first reviewer is from a nursing background and trained in systematic review methods and with experience of conducting reviews. The second reviewer is a clinician with experience of providing care for people with multimorbidity. Throughout the review process, any disagreement that arises will be resolved through discussion between the two reviewers or the chief investigator's arbitration.

3.1.3 Data extraction

Pre-designed data extraction tables will be created to organise data (Appendix 2). Where any relevant or important data that do not fit in the tables, the tables will be revised to CFC 0110

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facilitate data organisation and capture the holistic picture of the study topic. The types of data that will be extracted include:

- author's name
- year of publication
- study title
- purpose
- methodology (cross sectional/prospective)
- country
- study participants
- number of participants
- · reference definition of multimorbidity
- tool used to measure multimorbidity (disease counts, weighted indices, medicationbased measures)
- data collection method/data source (self-reports: survey or interview; administrative data, clinical routine data, research data)
- conditions, severity and other elements (e.g. age, gender, sociodemographic information) included in each measure
- rationale for selecting the items
- · weighting scheme
- testing of reliability or validity
- prevalence of multimorbidity in the whole population and by sub-groups
- outcome variable
- follow-up years
- results
- confounding factors.

These data will be entered into Microsoft Excel 2016 and will be checked for completeness by SH, PH and BG.

3.1.4 Data analysis

Narrative synthesis will be used to synthesise the evidence. Data will be categorised by collapsing those that are similar and dissimilar into broader and higher order of categories. The relationships and interactions between and within data, categories and observations will be explored (including patterns and processes). Data visualisation tools will be used during data analysis to help make sense of the data. Thereafter, a questionnaire for the Delphi survey will be developed based on review findings and gaps identified from the review.

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3.2 PHASE TWO: DELPHI STUDY

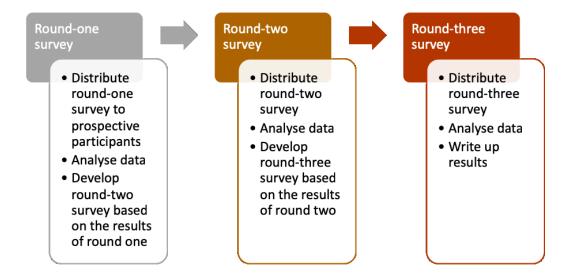


Figure 1: A Map of the project plan

The Delphi technique is designed as a communication process which allows for the inclusion of expert panellists from different sectors to participate in a detailed examination and discussion of a specific problem [15, 21, 22]. Therefore, this method is well suited for consensus building by using a series of questionnaires to collect data from the expert panel. Consensus here is referred to as a statement that is agreed by the majority [15, 23]. As a part of the Delphi process, each individual's responses from each questionnaire will be fed back to the panel alongside a summary of all other responses [15, 24]. The use of successive questionnaires can inform the panel members of the current status of their collective opinions [24]. Typically, more than two rounds of data collection are required to collect needed information and to reach consensus [22, 24, 25].

3.2.1 Study population

Unlike a traditional survey that aims to generalise results, the Delphi is a group decision mechanism requiring individuals who have knowledge, experience or deep understanding of the topic of interest [26]. In that sense, the main focus of this approach is the selection of experts and patients needed for inclusion of all the relevant perspectives [15]. In this proposed Delphi, we aim to recruit international experts and the public who have knowledge and experience of multimorbidity. Individuals who meet one of the following criteria will be included in this study:

Inclusion criteria





- Clinicians with experience of treating patients with more than one chronic condition
- Individuals who have been involved in health policy-making or research where multimorbidity was the focus or relevant
- Members of the public who are interested in multimorbidity or have experience of being multimorbid

3.2.2 Number of participants

There is no direction on the number of participants required for a Delphi survey [15]. The size of expert panel in existing Delphi studies varied considerably from under 10 to 500 [15]. In this Delphi study, the minimum number of experts we plan to recruit is approximately 25-30 experts.

3.2.3 Identifying participants

The expert panellists will be identified by SH (Szu-Szu Ho) and BG (Bruce Guthrie) using publicly available information, including:

- 1) published work (to identify researchers and policy makers)
- publicly available websites, reports, and policy documents (to identify healthcare professionals, policy makers or public participants e.g. in guideline development)
- 3) social media, such as twitter, where we will distribute this study information

or experts will be identified by research participants where they forward the study information to colleagues who meet the criteria.

3.2.4 Consenting participants

Participants will be invited to take part in the study through email along with an attachment of the participant information sheet (PIS) (Appendix 3 and 4). Participants will be asked to read the PIS before deciding to take part, and the survey link will be provided in the PIS to ensure that they have read the study information. Each participant will be given two to three weeks to consider and respond to each round.

3.2.5 Withdrawal of Study Participants

Participants will be informed that they can withdraw their participation at any time by not completing the survey and informing our researcher of their decision. If withdrawal occurs their personally-identifiable information will be permanently deleted, and they will also be allowed to request withdrawal of the data they submitted if they wish.

4 DATA COLLECTION

The Delphi process is iterated until consensus or the consistency of responses between successive rounds of survey is achieved [23]. In this proposed Delphi, we plan to use online questionnaires to collect data. Members of the panel will be anonymous to one another but the research team will know their identity in order that they can be sent individual feedback and invitations to participate in subsequent rounds of survey. The promise of anonymity among panellists is to facilitate participants to be open and truthful about their views on multimorbidity without feeling pressured by more influential panel members [15, 25]. We expect to have three rounds of data collection but will terminate early if consensus is

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reached and consider a fourth round if required for elements where consensus is not reached.

4.1 DEFINING CONSENSUS OR STABILITY

Consensus is pre-defined as 70% of participants agreeing/strongly agreeing or disagreeing/strongly disagreeing with a statement. However, not all statements in a Delphi study will gain consensus irrespective of the number of rounds [15, 25]. von der Gracht [23] therefore proposes that the level of consensus should not be the only criterion for stopping the Delphi process, but rather that sufficient clarification of the different viewpoints can also be a desirable goal. We will therefore not only examine consensus, but also stability of responses between rounds. The stability will be measured as the percentage change in agreement with a statement from round to round, and a change lower than 15% will be considered a stable answer [23]. If in round 3, all responses either achieve consensus or stability, then we will terminate the Delphi after round 3. If there are sufficient responses which lack consensus and are unstable, then we will take those items to a round 4.

4.1.1 Round 1

We will use the findings of the scoping review described above to identify the range of multimorbidity measures used in research and practice, their characteristics and use of standards, and purposes for measuring multimorbidity. The existing evidence and gaps identified from the review will serve as a point of reference for developing the round-one Delphi survey. The design of the questionnaire will be structured in a way that includes closed-ended/likert-scaled questions and some open-ended questions. The combination of both types of questions allows each participant an opportunity to generate additional insights, as well as formally scoring items derived by the research team from the literature review. Prospective participants will be required to rate (from strongly agree to strongly disagree or from very important to not important at all) and rank (rank importance of statements on a scale of 1-7) items or statements using Likert scales [15]. The open-ended responses will be triangulated with close-ended responses, and the results will be used to develop new items.

4.1.2 Round 2

In the second round, participants will receive a structured questionnaire and be asked to review their previous response and the items summarised based on the aggregated results of the first round. Second round items will be a mix of those scored in round 1 which did not achieve consensus, and new items based on round 1 open-ended responses. In this round, consensus may begin to form and areas of disagreement and agreement are likely to be identified.

4.1.3 Round 3

Round three will be designed using the results from round two. Prospective participants will receive a questionnaire that includes items or ratings summarised in the previous round and will be asked to review the results and provide their judgements. This round offers an opportunity for panellists to reflect and make further clarifications on the information developed by previous iterations and make their final judgements. In an attempt to keep the panel motivated, a summary sheet of statements that have achieved consensus will be provided and serve as feedback of completed work to the panel.

4.2 SOURCE DATA DOCUMENTATION

The documents that contain source data are participant-completed questionnaires stored on the Qualtrics or Jisc online survey tools, and in the downloaded csv file. The anonymous CFC 0110

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research data will be stored separately from personally-identifiable information, including email address and name (preferred name given by participants). These will be stored securely in the university server with password protection.

4.3 CASE REPORT FORMS

Not applicable

5 DATA MANAGEMENT

5.1 PERSONAL DATA

The personal information that will be collected in this study are: 1) name and email address; 2) country; 3) type of work they are doing; 4) their expertise (Appendix 6). All data will be stored securely on password-protected servers at the University of Edinburgh. The name and email address of participants will be kept separately from the main dataset, and will only be accessible to the research team. Once this study is completed, the file containing participants' names and email addresses will be deleted permanently. Data, other than names and email addresses, will not contain information that can identify participants, and thus will be stored securely and indefinitely on the university server and will not be transferred to external individuals or organisations outside of the sponsoring organisation.

Participants will be assured that their information will be treated confidentially and that their responses would remain anonymous to other panel members throughout the Delphi process. Nonetheless, the researchers will be able to identify participants through their name and email address, which allows for distributing subsequent rounds of survey to the participants and providing them with individual responses and a summary of results from the previous round. They will be informed that they can withdraw their participation at any time. Ethics approval for the study will be sought and obtained from the Usher Research Ethics Committee at the University of Edinburgh.

5.2 DATA INFORMATION FLOW

Data will be collected using Qualtrics or Jisc survey tools and stored in a downloaded csv file for analysis. After the study is completed, participants' personally-identifiable information, name and email address, will be deleted permanently.

5.3 TRANSFER OF DATA

Data collected by the study (including personal data) will not be transferred to any external individuals or organisations outside of the Sponsoring organisation.

5.4 DATA CONTROLLER

The University of Edinburgh and the research team involved in delivering the study will be the data controllers.

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5.5 DATA BREACHES

As described above, we will take care to ensure that identifiable information is kept separate from study data. Given the nature of the study data, the potential harm of any data breach is low. We therefore consider the risk of a data breach to be remote and low risk of harm if occurred. All data breaches will be reported to the University of Edinburgh and HDRUK.

6 STATISTICS AND DATA ANALYSIS

6.1 SAMPLE SIZE CALCULATION

There is no one sample size that has been advocated for Delphi studies. It has been suggested that a smaller sample size would be sufficient for a Delphi study, such as 10-15 participants, although others argue that larger sample sizes are likely to produce more generalizable results. Concerned over the lower response rate in online surveys and no guideline for estimating a Delphi sample size, we aim to recruit 25-30 participants for each round.

6.2 PROPOSED ANALYSES

A database will be set up for the analysis of quantitative data (including demographic data) of rounds 1, 2 and 3 using Rstudio (version 4.0.1). Frequencies and descriptive statistics will be used to provide information on the level of agreement with each statement across rounds and expert groups. If the distributions of the responses show less than 15% change from round to round, the responses will be considered to have reached stability [22]. Central tendencies (medians) and levels of dispersion (inter-quartile range) will be provided to see the probability distribution of responses. Bar charts will be created to visually depict the study results. For qualitative data in round one, similar responses will be collapsed and used to modify existing or construct additional unique statements for round 2. New statements created from round 1 will be used for the design of round 2 survey. Response rate for each round of the survey will be calculated.

7 ADVERSE EVENTS

The only plausible potential harm could occur in this study is breach of participants' confidentiality. Precautions will be seriously taken to prevent their identity being disclosed to other panellists and people outside of the research team. Firstly, the file that contains their private information will be kept on the university server with password protection, and separated from the main dataset. Secondly, an invitation email will be sent to participants individually and for the final report, we will use the BCC field to send an email to the participants who would like to receive the report. Once the study is completed, their name and email address will be deleted permanently. Together, these measures are to protect participants' confidentiality.

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8 GOOD CLINICAL PRACTICE

8.1 ETHICAL CONDUCT

The study will be conducted in accordance with the principles of the International Conference on Harmonisation Tripartite Guideline for Good Clinical Practice (ICH GCP).

Before the study can commence, all required approvals will be obtained and any conditions of approvals will be met.

8.2 INVESTIGATOR RESPONSIBILITIES

The Investigator is responsible for the overall conduct of the study at the site and compliance with the protocol and any protocol amendments. In accordance with the principles of ICH GCP, the following areas listed in this section are also the responsibility of the Investigator. Responsibilities may be delegated to an appropriate member of study site staff.

8.2.1 Informed Consent

The Investigator is responsible for ensuring informed consent is obtained before any protocol specific procedures are carried out. The decision of a participant to participate in clinical research is voluntary and should be based on a clear understanding of what is involved.

Participants must receive adequate oral and written information – appropriate Participant Information and Informed Consent Forms will be provided. The oral explanation to the participant will be performed by the Investigator or qualified delegated person, and must cover all the elements specified in the Participant Information Sheet and Consent Form.

The participant must be given every opportunity to clarify any points they do not understand and, if necessary, ask for more information. The participant must be given sufficient time to consider the information provided. It should be emphasised that the participant may withdraw their consent to participate at any time without loss of benefits to which they otherwise would be entitled.

The participant will be informed and agree to their medical records being inspected by regulatory authorities and representatives of the sponsor(s).

The Investigator or delegated member of the trial team and the participant will sign and date the Informed Consent Form(s) to confirm that consent has been obtained. The participant will receive a copy of this document and a copy filed in the Investigator Site File (ISF) and participant's medical notes (if applicable).

8.2.2 Study Site Staff

The Investigator must be familiar with the protocol and the study requirements. It is the Investigator's responsibility to ensure that all staff assisting with the study are adequately informed about the protocol and their trial related duties.

8.2.3 Data Recording

Not applicable

8.2.4 Investigator Documentation

Not applicable

8.2.5 GCP Training

For non-CTIMP (i.e. non-drug) studies all researchers are encouraged to undertake GCP training in order to understand the principles of GCP. However, this is not a mandatory requirement unless deemed so by the sponsor. GCP training status for all investigators should be indicated in their respective CVs.

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8.2.6 Confidentiality

All laboratory specimens, evaluation forms, reports, and other records must be identified in a manner designed to maintain participant confidentiality. All records must be kept in a secure storage area with limited access. Clinical information will not be released without the written permission of the participant. The Investigator and study site staff involved with this study may not disclose or use for any purpose other than performance of the study, any data, record, or other unpublished information, which is confidential or identifiable, and has been disclosed to those individuals for the purpose of the study. Prior written agreement from the sponsor or its designee must be obtained for the disclosure of any said confidential information to other parties.

8.2.7 Data Protection

All Investigators and study site staff involved with this study must comply with the requirements of the appropriate data protection legislation (including the General Data Protection Regulation and Data Protection Act) with regard to the collection, storage, processing and disclosure of personal information.

Computers used to collate the data will have limited access measures via user names and passwords.

Published results will not contain any personal data and be of a form where individuals are not identified and re-identification is not likely to take place.

9 STUDY CONDUCT RESPONSIBILITIES

9.1 PROTOCOL AMENDMENTS

Any changes in research activity, except those necessary to remove an apparent, immediate hazard to the participant in the case of an urgent safety measure, must be reviewed and approved by the Chief Investigator.

Amendments will be submitted to a sponsor representative for review and authorisation before being submitted in writing to the appropriate REC, and local R&D for approval prior to participants being enrolled into an amended protocol.

9.2 MANAGEMENT OF PROTOCOL NON COMPLIANCE

Not applicable

9.3 SERIOUS BREACH REQUIREMENTS

A serious breach is a breach which is likely to effect to a significant degree:

- (a) the safety or physical or mental integrity of the participants of the trial; or
- (b) the scientific value of the trial.

If a potential serious breach is identified by the Chief investigator, Principal Investigator or delegates, the co-sponsors (seriousbreach@accord.scot) must be notified within 24 hours. It is the responsibility of the co-sponsors to assess the impact of the breach on the scientific value of the trial, to determine whether the incident constitutes a serious breach and report to research ethics committees as necessary.

9.4 STUDY RECORD RETENTION

All study documentation will be kept for a minimum of 3 years from the protocol defined end of study point. When the minimum retention period has elapsed, study documentation will not be destroyed without permission from the sponsor.

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9.5 END OF STUDY

The end of study is defined as the last participant's last visit.

The Investigators or the co-sponsor(s) have the right at any time to terminate the study for clinical or administrative reasons.

The end of the study will be reported to the REC, and R+D Office(s) and co-sponsors within 90 days, or 15 days if the study is terminated prematurely. The Investigators will inform participants of the premature study closure and ensure that the appropriate follow up is arranged for all participants involved. End of study notification will be reported to the co-sponsors via email to response-reported to the co-sponsors via email to <a href="responted-response-reported-r

A summary report of the study will be provided to the REC within 1 year of the end of the study.

10 REPORTING, PUBLICATIONS AND NOTIFICATION OF RESULTS

10.1 AUTHORSHIP POLICY

Ownership of the data arising from this study resides with the study team.

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12 APPENDICES

APPENDIX 1 – SEARCH STRATEGY

Database	Search strategy
Ovid Interface PsycINFO Embase Global Health Ovid MEDLINE	 (multimorbidit\$ or multi-morbidit\$ or comorbidit\$ or co-morbidit\$ or polymorbidit\$ or poly-morbidit\$ or multicondition\$ or multicondition\$ or multiple chronic condition\$" or "morbidity burden" or ((multiple or coexisting or co-existing or concurrent or con-current or comorbid or co-morbid) adj2 (disease\$ or illness\$ or condition\$ or diagnos\$ or morbid\$))).m_titl. (measure\$ or index or indices or instrument\$ or scale\$ or "disease count\$").mp. 1 and 2
EBSCO Interface	Limit 3 to human MM (multimorbidit* or multi-morbidit* or comorbidit*
EBSCO Interface CINAHL Plus	or co-morbidit" or multi-morbidit" or comorbidit" or co-morbidit" or polymorbidit* or multicondition* or multicondition* or "multiple chronic condition*" or "morbidity burden" or ((multiple or coexisting or co-existing or concurrent or con-current or comorbid or co-morbid) N2 (disease* or illness* or condition* or diagnos* or morbid*))) 2. AB (measure* or index or indices or instrument* or scale*) 3. 1 AND 2
0	Limiters – Full Text; Human; Language: English
Scopus	TITLE (multimorbidit* or multi-morbidit* or comorbidit* or co-morbidit* or polymorbidit* or poly-morbidit* or multicondition* or multicondition* or "multiple chronic condition*" or "morbidity burden" or ((multiple or coexisting or co-existing or concurrent or con-current or morbid or co-morbid) W/2 (disease* or illness* or condition* or diagnos?s or morbid*)) AND TITLE (measure* or index or indices or instrument* or scale* or "disease counts")
Web of Science	(TI=(measure* or index or indices or instrument* or scale*))AND (TI=(multimorbidit* or multi-morbidit* or comorbidit* or co-morbidit* or polymorbidit* or polymorbidit* or multicondition* or multicondition* or 'multiple chronic condition*' or 'morbidity burden' or ((multiple or coexisting or co-existing or concurrent or comorbid or co-morbid) NEAR/2 (disease* or illness* or condition* or diagnos* or morbid*)))) AND LANGUAGE: (English)

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Cochrane library	(multimorbidity or multi-morbidity or comorbidity or comorbidity or polymorbidity or poly-morbidity or multicondition or multicondition or 'multiple chronic conditions' or 'morbidity burden' or ((multiple or coexisting or co-existing or concurrent or con-current or comorbid or co-morbid) NEAR/2 (disease or illness or condition or diagnosis or morbid))) AND (measure or index or indices or instrument or scale or "disease count*"):ti
	ti((multimorbidit* OR multi-morbidit* OR comorbidit* OR
Theses Global	co-morbidit* OR polymorbidit* OR poly-morbidit* OR
	multicondition* OR multicondition* OR 'multiple chronic condition*' OR 'morbidity burden' OR ((multiple OR
	coexisting OR co-existing OR concurrent OR con-current
	OR morbid OR co-morbid) NEAR/2 (disease* OR illness* OR condition* OR diagnos?s OR morbid*)))) AND
	noft((measure* OR index OR indices OR instrument* OR scale*))
	Limited by: Manuscript type: Doctoral dissertations,
	Master's theses
	Language: English





APPENDIX 2 - DATA EXTRACTION FORMS FOR THE SCOPING REVIEW

Characteristics of multimorbidity measures

Study	Purpose	Participants	Country/Region	Definition of	Measurement	Data	Data	Elements in	Rationale	Weighting	Reliability	Prevalence of	Outcome
	-	-		Multimorbidity	tool	collection	source	the	for		/validity	multimorbidity	variable
						method		measure	selecting				
									the items				
								Condition,					
								severity,					
								frailty					

Confounding factors	Key results

Items of multimorbidity measures

Rome of Hidden of Market Marke									
Measure	Condition name	Severity scale	Age						
Name of		0	0	0	0				
measure					_				
Name of)			0	0	0	0		
measure						_			
Name of									
measure									

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APPENDIX 3 – INVITATION EMAIL

Subject: Multimorbidity definition and measurement study

Dear [to be addressed individually to each participant],

We are a research team working on a project aiming to explore an international consensus on the definition and measurement of multimorbidity. This study is funded by the Health Data Research UK (HDR UK — CFC0110).

We send you this email to invite you to participate in a Delphi survey. Your name and email address were identified from one of your published articles, reports, policy documents, or websites relevant to multimorbidity through our scoping review conducted between January and September, 2020. We seek your participation in this survey as we appreciate your valuable input on multimorbidity definition and measurement. International multimorbidity researchers, clinicians, policy makers and members of the public are all welcome to take part in the study. Please share this survey with your colleagues, if you can, to ensure we reach as many experts as possible. This research experience allows us to share, debate and scrutinise anonymously and move forward through consensus.

The Delphi process consists of **two to four** surveys— one that needs to be completed by [date] and the other rounds that will be sent out subsequently in the next few months. The first survey will take about 22 minutes to complete. Taking part in the survey is voluntary and please read the attached Participant Information Sheet (PIS) carefully before you decide to take part. The online survey can be accessed via the link provided in the attached PIS (this is to ensure your acknowledgement of the study information prior to participation).

The data you provide us will be treated confidentially and will not be accessed by anyone outside of the research team.

The results are envisioned to support and inform research, practice and policy around multimorbidity definition and measurement thanks to your contribution. A summary of the results will be available to you at the end of the study.

We hope that you are able to support this important work. If you have any questions, please contact the researcher, Iris Ho or the principal investigator, Bruce Guthrie.

Yours Sincerely,

Professor Bruce Guthrie, Chief investigator, Centre for Population and Health Sciences, Usher Institute, University of Edinburgh, Email: bruce.guthrie@ed.ac.uk

Dr Iris S.S. Ho, Centre for Population and Health Sciences, Usher Institute, University of Edinburgh, Email: iris.s.ho@ed.ac.uk

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APPENDIX 4 - FOLLOW-UP EMAIL REMINDER

Subject: Multimorbidity definition and measurement study

Dear [to be addressed individually to each participant],

You were recently invited to participate in a Delphi survey aimed at exploring a consensus on multimorbidity definition and measurement (see the email below). If you have not already completed the survey, we encourage you to take a few munities to do so before [date]. If you have already completed the survey we thank you for your support and ask you to ignore this email. Your valuable opinions can help to shape the future of multimorbidity research and management. If possible, please share this study information with your colleagues in the field.

The survey can be accessed via the link provided in the attached Participant Information Sheet. Please complete the survey by [date].

Thank you in advance.

Yours sincerely,

Professor Bruce Guthrie, Chief investigator, Centre for Population and Health Sciences, Usher Institute, University of Edinburgh, Email: bruce.guthrie@ed.ac.uk

Dr Iris S.S. Ho, Centre for Population and Health Sciences, Usher Institute, University of Edinburgh, Email: iris.s.ho@ed.ac.uk

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APPENDIX 5 – PARTICIPANT INFORMATION SHEET

Principal investigator: Professor Bruce Guthrie

Postdoctoral researcher: Dr Iris Ho

Institute: Usher Institute, University of Edinburgh

Research Ethics Committee: Usher Institute Research Ethics Committee at the University of

Edinburgh (ref:).
Survey link:

• If you are **clinicians**, **academics**, **researchers or policy makers** who have read the PIS and agree to take part, please access the survey via this link:

 If you are members of the public who have read the PIS and agree to take part, please access the survey via this link:

You are being invited to take part in this Delphi study aimed at exploring international consensus and guide on the definition and measurement of multimorbidity. Before you decide to take part, it is important that you understand why this research is being conducted and what it involves. Please take time to read this study information carefully.

Thank you for taking the time to read this.

Who will conduct the research?

This research is being conducted by Iris Ho, a researcher at the University of Edinburgh, and Professor Bruce Guthrie (the Chief Investigator). Our names and contact details are given at the end of this information sheet.

Aim of this study

The aim of this study is to explore a consensus on the definition of multimorbidity and on the design of quantitative measures of multimorbidity.

Why have I been invited to take part?

You are invited to participate in this study because we recognised your expertise relevant to multimorbidity through your published work, publicly-available reports, websites, or policy documents following our scoping review. You may also have been invited to take part in this study by a colleague who has forwarded the survey to you.

Do I have to take part?

No – participation is entirely voluntary. You can ask questions about the study before deciding whether or not to participate. If you do agree to participate, please keep this study information sheet and check the "Yes" boxes at the first page of the online survey to indicate that you agree with the listed statements and consent to take part.

Even if you agree to participate, you may withdraw yourself from the study at any time and without giving a reason by advising the researchers of this decision. We will respect your decision to withdraw and there will be no loss of good feeling/will. If withdrawal occurs, your personal and contact information we collected for this project will be permanently deleted. Participant survey responses collected prior to your withdrawal will be kept for analyses, as they will be completely anonymous.

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measures>
<Version 1: 24/February/2020>





What will happen if I decide to take part?

We are inviting you to participate as a panel member in this Delphi study. The Delphi method is used in this study to obtain your opinions and ideas on the definition of multimorbidity, and what should be included in multimorbidity measures and how to use them for different purposes. Two to four rounds of survey will be carried out to ensure that the convergence of opinion on this topic is achieved. This survey takes approximately 22 minutes to complete. A second round of survey will be sent to you via email in a few months following data analysis. Thank you for your contribution.

What are the possible benefits of taking part?

Your valuable input in this Delphi process will allow researchers, practitioners, funders and policy makers in the field to better understand what constitutes multimorbidity, and how to measure it to meet different purposes.

Are there any risks associated with taking part?

There are no foreseeable risks associated with participation in this study.

What happens to the data provided?

We will not tell anyone you have taken part in this study. The research data you give us will be anonymised and stored confidentially at our university server with password protection. Only Bruce Guthrie and Iris Ho have the permission to use the data. The name and email address we collected from you is for sending the subsequent rounds of survey. This information will be permanently deleted after this study is completed. The anonymous research data will be stored securely and indefinitely and will not be transferred to external individuals or organisations outside of the sponsoring organisation.

What will happen with the results of this study?

The results of this study will be summarised in articles and reports, and will be disseminated through peer-reviewed journals and presented at conferences. The results will be made anonymous in any formal outputs, and be available to you all.

Who do I contact if I have a concern about the study or I wish to complain?

If you have any further questions about the study, please contact Dr Iris Ho or the Principal Investigator Professor Bruce Guthrie, who will do their best to answer your query. If you remain unhappy or wish to make a formal complaint, please contact Professor Sarah Cunningham Burley, Dean SMGPHS [Email: sarah.c.burley@ed.ac.uk; Tel: 01316503217]

Further information and contact details

Principal investigator: Professor Bruce Guthrie [bruce.guthrie@ed.ac.uk] Postdoctoral researcher: Dr Iris S.S. Ho [iris.s.ho@ed.ac.uk]

Survey link:

- If you are clinicians, academics, researchers or policy makers who have read the PIS and agree to take part, please access the survey via this link:
- If you are members of the public who have read the PIS and agree to take part, please access the survey via this link:

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Measuring multimorbidity in research: a Delphi consensus study

Appendix 2: Surveys

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 \square No

choose all that apply)?

Round-one survey in the professional panel

Socio-demographic information
Q1: Please describe the country where you are currently living or working.
Q2: Please check the category that best describes the type of work you do most of the time (choose one).
☐ Research ☐ Public policy or other public sector work ☐ Clinical Practice ☐ Other ☐ Other ☐ Q3: Please check the category that best describes the setting in which you work most of the time (choose one).
☐ Government ☐ Academia ☐ Hospital ☐ Primary care/General practice ☐ Social care sector ☐ Other
Q4: Do you have multiple chronic conditions?
□ Yes □ No
Q5: Do you have family or friends who have multiple chronic conditions?
□ Yes

Q6: To understand multimorbidity, what population would you be more interested in (please

Round one professional panel
☐ General population ☐ Older people ☐ Middle-aged and older ☐ Socially-deprived population (including homeless people or drug users) ☐ Women ☐ Men ☐ Children ☐ Other
Operational definition of multimorbidity
Q7: Researchers have defined multimorbidity in many ways. Some define it as '2 or more long-term conditions', but others define it as '3 or more', '4 or more' or '5 or more' long-term conditions. How many long-term conditions do you think someone has to have in order to have multimorbidity? Please choose one.
✓ 2 or more long-term conditions☐ 3 or more long-term conditions
□ 4 or more long-term conditions
☐ Other, X or more long-term conditions (please only type a number in the box)
Q8: How would you define "condition" for the concept of multimorbidity? (please choose al that apply)
□ Formal medical diagnoses (e.g. coronary heart disease, alcohol dependence) □ Clinical risk factors (e.g. obesity, high cholesterol) □ Symptoms that are not formal medical diagnoses (e.g. dizziness or fatigue) □ Health behaviours (e.g. smoking or exercise level) □ Health impact (e.g. disability or frailty) □ Social deprivation and poverty □ Consequences of treatment and care (e.g. side effects of medications or the overal burden of treatment) □ Other (please specify)

Q9: Researchers justify their choice of conditions in many different ways. Please answer the following questions on the principles for selection of chronic conditions in multimorbidity measures.										
 How long-term a condition is: Researchers vary in what they mean by "long-term". How long does a condition have to be to count as long-term? Please choose one. 										
☐ Conditions lasting for three ☐ Conditions lasting for six r ☐ Conditions lasting for twel ☐ Other, conditions lasting for box) 2) Whether a condition is disagree with each of the	nonths or move months or X months	nore or more s or more (ple								
Statement	Strongly disagree	Disagree	Neither agree nor disagree	Agree	Strongly agree	Don't know				
Include conditions which are permanent in their effects (e.g. Parkinson's disease)										
Include conditions which are currently active or currently treated (e.g. asthma with intermittent wheeze; asthma using regular inhalers)										
Include conditions which may recur but happen rarely (e.g. people with a history of asthma or depression with no current symptoms and not currently on treatment)										
Other										
3) Whether condition is treated in healthcare: Please rate the degree to which you agree or disagree with each of the following statements										
Statement	Strongly disagree	Disagree	Neither agree nor disagree	Agree	Strongly agree	Don't know				

Round	l one	prof	fessi	ional	pane	ı
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Only include conditions which usually require formal treatment or care (e.g. hypertension, diabetes, schizophrenia)			
Other			

4) The impact of the condition on a range of outcomes: Please rate the degree to which you agree or disagree with each of the following statements

Statement	Strongly disagree	Disagree	Neither agree nor disagree	Agree	Strongly agree	Don't know
Include conditions which significantly increase risk of death						
Include conditions which significantly reduce health-related quality of life						
Include conditions which cause significant physical disability						
Include conditions which cause frailty (general physical and/or mental weakness and vulnerability)						
Include conditions which significantly worsen mental health						
Include conditions which significantly worsen self-perceived health status						
Impact: Other						

Q10: Researchers vary in how detailed their definitions of 'condition' are. For example, they might be very broad – one category for "lung disease". Or they might be more detailed – separately count "asthma", "chronic obstructive pulmonary disease" and so on. Please choose one option and explain your choice if you like.

Round	one	prof	fessional	l panel
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☐ Broad disease category based on body and mind system (e.g. cardiovascular disease, mental health problems, skin conditions). Please explain:
☐ Individual conditions (e.g. coronary heart disease, hypertension, depression, schizophrenia, eczema, psoriasis). Please explain:
☑ Other. Please explain:
☑ Don't know
Q11: Some conditions included in multimorbidity measures are closely linked to each other. In particular, some conditions can be caused by other conditions. For example, heart attacks can lead to heart failure. Diabetes can lead to kidney failure. How do you think researchers should count conditions in this situation? Please choose one option and explain your choice it you like.
☐ Count all of the conditions that are currently active. Please explain:
☐ Only count the complications (e.g. if people with heart attack develop heart failure, we would count only heart failure). Please explain:
☐ Count the primary health condition (e.g. if people with diabetes develop kidney failure, we would only count diabetes). Please explain:
☐ Other. Please explain:

if

☑ Don't know

What conditions should be included?

In our review of over 500 research studies, we identified all the conditions which researchers have counted when measuring multimorbidity. There is very large variation in which conditions researchers choose. Only seven conditions were counted by more than half of studies. Many conditions were rarely counted. We would like you to rate conditions that you think are more or less important to include in multimorbidity measures. There are now a set of questions organised by body system.

Q12: We are starting with body systems/disease domains. Please rate how important you think it is to include the following systems/domains in a multimorbidity measure.

Condition	Exclude (not importan t)	Usually exclude (unless a good reason to include in a particular context)	Could include or exclude	Usually include (unless a good reason to exclude in particular context)	Always include (extremely important)	Don't know
Mental health						
Cancer						
Cardiovascular						
Haematological disease						
Neurological disease						
Metabolic and Endocrine						
Musculoskeletal						
Digestive						
Urogenital						
Skin						
Ear, Nose and Throat (ENT)						
Oral						
Ophthalmology						
Chronic infections						

Q13: Please rate how important you think it is to include the listed mental health conditions in a multimorbidity measure. If you are not sure, just tick 'Don't know'.

Condition	Exclude (not important)	Usually exclude (unless a good reason to include in a particular context)	Could include or exclude	Usually include (unless a good reason to exclude in particular context)	Always include (extremely important)	Don't know
Anxiety						
Depression						
Dementia						

Schizophrenia				
Substance misuse				
(including				
alcohol and drug				
dependence)				
Nicotine				
dependence				
Post-traumatic				
stress disorder				
(PTSD)				
Bipolar disorder				
Chronic insomnia				
Dissociative or				
personality				
disorders				
Eating disorders				
(including				
bulimia or				
anorexia)				
Learning				
disability				
Autism				
Obsessive				
compulsive				
disorder (OCD)				
Somatoform				
disorders				
(psychological				
disorder where a				
person				
experiences				
physical				
symptoms that cannot be				
explained by				
medical doctors)				
Attention deficit				
hyperactivity				
disorder (ADHD)				
Other (please				
specify either one				
specify chiler one		ı		
or more				
or more conditions):				

Q14: Please rate how important you think it is to include the listed cancers in a multimorbidity measure.

Condition	Exclude (not important)	Usually exclude (unless a good reason to include in a particular context)	Could include or exclude	Usually include (unless a good reason to exclude in particular context)	Always include (extremely important)	Don't know
Solid organ cancers (e.g. lung,						

Round	one	professional	panel
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colon, prostate,			
breast etc)			
Haematological			
cancers (e.g.			
leukaemia,			
lymphoma,			
myeloma)			
Melanoma			
(malignant skin			
cancer)			
Non-melanoma			
skin cancer			
Benign cancers			
(excluding benign			
skin lumps and			
bumps)			
Metastatic cancers			
Other (please			
specify either one			
or more			
conditions):			

Q15: For cancers, some researchers count every cancer a person has had. Others only count 'cancer' once even if someone has had more than one type of cancer. Which methods would you recommend. Please choose one:

Count all cancers as one
Count individual cancers separately.
Other

Q16: Please rate how important you think it is to include the listed cardiovascular conditions in a multimorbidity measure.

Condition	Exclude (not important)	Usually exclude (unless a good reason to include in a particular context)	Could include or exclude	Usually include (unless a good reason to exclude in particular context)	Always include (extremely important)	Don't know
Hypertension						
Stroke						
Transient Ischaemic Attack (mini stroke)						
Dyslipidaemia/ Lipid disorder (e.g. high cholesterol)						

Coronary artery disease (heart attack or angina)			
Heart failure			
Peripheral artery disease			
Arrhythmia			
Heart valves problem			
Other (please specify either one or more conditions):			

Q17: Please rate how important you think it is to include the listed haematological conditions in a multimorbidity measure.

Condition	Exclude (not important)	Usually exclude (unless a good reason to include in a particular context)	Could include or exclude	Usually include (unless a good reason to exclude in particular context)	Always include (extremely important)	Don't know
Anaemia (including Iron deficiency anaemia, pernicious anaemia, sickle cell anaemia, aplastic anaemia)						
Venous thrombotic disease						
Other (please specify either one or more conditions):						

Q18: Please rate how important you think it is to include the listed neurological conditions in a multimorbidity measure.

Condition	Exclude (not important)	Usually exclude (unless a good reason to include in a particular context)	Could include or exclude	Usually include (unless a good reason to exclude in particular context)	Always include (extremel y important)	Don't know
Parkinson's disease						
Epilepsy (fits/ seizures)						
Chronic pain (including chronic low back pain fibromyalgia,						

trigeminal neuralgia			
and other chronic pain)			
Migraine or other			
regular headache			
Multiple sclerosis			
Peripheral neuropathy			
Paralysis/ Hemiplegia/			
Paraplegia (not			
including those caused			
by stroke)			
Other (please specify			
either one or more			
conditions):			

Q19: Please rate how important you think it is to include the listed metabolic/endocrine/nutritional conditions in a multimorbidity measure.

Condition	Exclude (not important)	Usually exclude (unless a good reason to include in a particular context)	Could include or exclude	Usually include (unless a good reason to exclude in particular context)	Always include (extremely important)	Don't know
Diabetes (any type)						
Thyroid problems (including hypothyroidism and hyperthyroidism)						
Malnutrition						
Other (please specify either one or more conditions):						

Q20: Please rate how important you think it is to include the listed musculoskeletal conditions in a multimorbidity measure.

Condition	Exclude (not important)	Usually exclude (unless a good reason to include in a particular context)	Could include or exclude	Usually include (unless a good reason to exclude in particular context)	Always include (extremely important)	Don't know
Osteoporosis (thinning of the bones)						
Osteoarthritis						
Connective tissue disease						
Gout						
Long-term musculoskeletal						

Round	l one	prof	fessiona	l panel
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problems due to injury (including hip fracture)			
Other (please specify either one or more conditions):			

Q21: Please rate how important you think it is to include the listed respiratory conditions in a multimorbidity measure.

Condition	Exclude (not important)	Usually exclude (unless a good reason to include in a particular context)	Could include or exclude	Usually include (unless a good reason to exclude in particular context)	Always include (extremely important)	Don't' know
Chronic obstructive pulmonary disease						
Asthma						
Cystic fibrosis						
Chronic/Allergic rhinitis						
Other (please specify either one or more conditions):						

Q22: Please rate how important you think it is to include the listed gastrointestinal conditions in a multimorbidity measure.

Condition	Exclude (not important)	Usually exclude (unless a good reason to include in a particular context)	Could include or exclude	Usually include (unless a good reason to exclude in particular context)	Always include (extremely important)	Don't know
Chronic liver disease (including liver						
cirrhosis, liver failure and chronic hepatitis)						
Inflammatory bowel disease (including ulcerative colitis and Crohn's disease)						
Irritable bowel syndrome						
Diverticular disease						
Pancreatic disease						

Round	l one	prof	fessi	ional	pane	ı
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Peptic ulcer (including gastric/ stomach ulcers and duodenal ulcers)			
Gastroesophageal			
reflux (acid reflux and			
heartburn)			
Gall bladder problems			
(including gallstones)			
Other (please specify			
either one or more			
conditions):			

Q23: Please rate how important you think it is to include the listed urogenital conditions in a multimorbidity measure.

Condition	Exclude (not important)	Usually exclude (unless a good reason to include in a particular context)	Could include or exclude	Usually include (unless a good reason to exclude in particular context)	Always include (extremely important)	Don't know
Chronic kidney disease						
End-stage kidney disease (including kidney dialysis and transplant)						
Kidney or bladder stones						
Chronic urinary tract infections (including chronic bladder infections)						
Urinary incontinence						
Uterus (womb) problems (including prolapse and fibroid)						
Polycystic ovary syndrome						
Prostatic hypertrophy						
Endometriosis						
Infertility						
Sexual dysfunction						
Other (please specify either one or more conditions):						

Q24: Please rate how important you think it is to include the listed chronic infectious conditions in a multimorbidity measure.

Condition	Exclude (not important)	Expect to usually exclude (unless a good reason to include in a particular context)	Could include or exclude	Expect to usually include (unless a good reason to exclude in particular context)	Always include (extremely important)
HIV/AIDS					
Tuberculosis					
Other (please specify either one or more conditions):					

Q25: Please rate how important you think it is to include the listed skin conditions in a multimorbidity measure.

Condition	Exclude (not important)	Usually exclude (unless a good reason to include in a particular context)	Could include or exclude	Usually include (unless a good reason to exclude in particular context)	Always include (extremely important)	Don't know
Psoriasis						
Eczema						
Chronic urticarial (chronic hives)						
Other (please specify either one or more conditions):						

Q26: Please rate how important you think it is to include the listed ENT, eye & oral conditions in a multimorbidity measure.

Condition	Exclude (not	Usually	Could	Usually	Always	Don't
	important)	exclude	include or	include	include	know
		(unless a	exclude	(unless a	(extremely	
		good reason		good reason	important)	
		to include in		to exclude in		
		a particular		particular		
		context)		context)		
Hearing						
impairment or						
deafness						

Meniere's disease			
Ear, nose, throat disease (including chronic sinusitis)			
Vision impairment or blindness			
Cataract			
Glaucoma			
Edentulism (having no teeth)			
Chronic gum disease			
Other (please specify either one or more conditions):			

Q27: Please rate how important you think it is to include the listed congenital conditions in a multimorbidity measure.

Condition	Exclude (not important)	Usually exclude (unless a good reason to include in a particular context)	Could include or exclude	Usually include (unless a good reason to exclude in particular context)	Always include (extremely important)	Don't know
Congenital disease (conditions that babies are born with, including congenial heart disease, genital anomaly)						
Other (please specify either one or more conditions):						

Q28: Please rate how important you think it is to include the listed risk factors/health behaviour/symptoms/syndromes in a multimorbidity measure.

Condition	Exclude (not important)	Usually exclude (unless a good reason to include in a particular context)	Could include or exclude	Usually include (unless a good reason to exclude in particular context)	Always include (extremely important)	Don't know
Obesity (body mass index ≥30)						
Smoking						

Round	l one	prof	fessi	ional	pane	ı
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High blood		Ш	Ш	Ш	
pressure					
(untreated)					
High cholesterol					
(untreated)					
Sedentary lifestyle					
Physical disability					
Dizziness (without					
a specific					
diagnosis)					
Chronic cough					
(without a specific					
diagnosis)					
Post-sepsis					
syndrome					
Side effects of					
medications					
Treatment burden					
(the sum of all the					
hassles of taking					
medicines or					
attending					
appointments)					
Social deprivation					
and poverty					
Other (please					
specify either one					
or more					
conditions):					

Weighting

Most researchers define multimorbidity by just counting how many conditions someone has (a "simple count" of conditions). Others use a "weighted" count to estimate multimorbidity burden and predict outcomes (e.g. death, hospitalisation, quality of life). For example, a simple count would say that hay fever + heart attack + back pain = 3 conditions. A weighted measure that gave more weight to risk of death might count hay fever = 0.5, high blood pressure = 3, back pain = 0.5, total score = 4 weighted for risk of death. However a weighted measure that gave more weight to quality of life might count hay fever = 1, high blood pressure = 0.5, and back pain = 2, total score = 3.5 weighted for quality of life.

In practice, weighted counts have been commonly used for predicting outcomes despite the fact that a simple count is the most common type of measures used for various purposes.

Q29: Please describe what type of measures you would use to measure multimorbidity and for what purposes.
☐ Weighted morbidity measures for the purpose of (please describe):

Round one professional panel
☐ A simple count of conditions for the purpose of (please describe):
☐ Other (please describe)
Q30: There is much debate over whether 'weighted morbidity measures' or 'simple counts or conditions' are better at predicting outcomes (e.g. mortality, healthcare utilisation etc). Please describe what type of measures you would use to understand the impact of multimorbidity on outcomes.
☐ Simple counts of conditions

Note: A key issue for weighted measures, is which outcomes to focus on. Researchers vary in which outcomes they think are most important. Professionals and patients also vary in which outcomes they think are important.

☐ Weighted morbidity measures

☐ Other (please specify)

Q31: Please rate the degree to which the outcomes listed below are important to be weighted against.

Outcome	Not at all important	Slightly important	Important	Fairly important	Very important	No opinion
Mortality						
Healthcare use (e.g. number of emergency admissions to hospital; outpatient appointments)						
Health-related quality of life						
Physical disability						
Frailty (general physical and/or						

mental weakness			
and vulnerability)			
Mental health			
Treatment burden			
(the sum of all the			
hassles of taking			
medicines or			
attending			
appointments)			
Healthcare costs			
(how much			
treatment and care			
for each individual			
costs)			
Self-perceived			
health status			
Other (please			
specify)			
		l l	

Q32: What weighting methods would you use in a multimorbidity measure?	
☐ Use existing weighted indices.	
☐ Empirically derive weights based on the individual impact of disea outcome (e.g. use regression models to calculate weights)	ses on an
☐ Set rules based on level of severity to grade each condition/disease rate each condition/category based on the rules on if having— presence condition: 1 point; treatment: additional 1 point; functional limitation: point)	e of a
☐ No opinion	
□ Other	

choose all that apply)?

☐ General population

Round-two survey in the professional panel

Socio-demographic information
Q1: Please describe the country where you are currently living or working.
Q2: Please check the category that best describes the type of work you do most of the time (choose one).
 □ Research □ Public policy or other public sector work □ Clinical Practice □ Teaching □ Other
Q3: Please check the category that best describes the setting in which you work most of the time (choose one).
☐ Government ☐ Academia ☐ Hospital ☐ Primary care/General practice ☐ Social care sector ☐ Other
Q4: Do you have multiple chronic conditions?
□ Yes □ No
Q5: Do you have family or friends who have multiple chronic conditions?
□ Yes □ No

Q6: To understand multimorbidity, what population would you be more interested in (please

Round two professional panel
 □ Older people □ Middle-aged and older □ Socially-deprived population (including homeless people or drug users) □ Women □ Men □ Children □ Ethnic minority groups or indigenous populations □ People with disabilities □ Other
Operational definition of multimorbidity
Note inserted: In round one, there was no consensus on the number of conditions someone have to have in order to have multimorbidity. There were also numerous free text comments including some which suggested that there should be distinctions made between 'simple' an 'complex' multimorbidity. In this section, we are therefore asking you the round 1 question about 'number of conditions' and some new questions about 'simple' and 'complex' multimorbidity.
Q7-1: How many long-term conditions do you think someone has to have in order to have multimorbidity? Please choose one.*
 □ 2 or more long-term conditions (chosen by 68.5% of panellists in round one) □ 3 or more long-term conditions (chosen by 28.9% of panellists in round one) □ 4 or more long-term conditions (chosen by 1.3% of panellists in round one) □ 5 or more long-term conditions (chosen by 0.7% of panellists in round one) □ Other, X or more long-term conditions
Q7-2-1: Some studies differentiated "complex multimorbidity" from "basic multimorbidity" and used "complex multimorbidity" to identify people with higher care needs (e.g. older people). Do you agree that defining complex multimorbidity in addition to a core definition of simple multimorbidity is useful?
Strongly disagree/Disagree/Neither disagree nor agree/Agree/Strongly agree/Don't know
Q7-2-2: Irrespective of whether you agree, how would you define "complex multimorbidity based on number of conditions. (please choose one)
☐ 3+ conditions irrespective of how many body systems

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3+ conditions from 3+ body systems 4+ conditions irrespective of how many body systems 4+ conditions from 4+ body systems 5+ conditions irrespective of how many body systems 5+ conditions from 5+ body systems Other+ conditions from+ body systems
p-question: In addition to the above option you choose, please describe if there are nents that, you think, should be included in the definition of "complex iity".
Any combination of 2+ conditions which includes both physical and mental th conditions Other
ed: Based on the results from round one, only medical diagnoses reached ≥70% Having read a summary of the results from round one, we would like you to question again.
you define "condition" for the concept of multimorbidity? (please choose all that
Formal medical diagnoses (e.g. coronary heart disease, alcohol dependence) Clinical risk factors (e.g. obesity, high cholesterol) Symptoms that are not formal medical diagnoses (e.g. dizziness or fatigue) Health behaviours (e.g. smoking or exercise level) Health impact (e.g. disability or frailty) Social deprivation and poverty Consequences of treatment and care (e.g. side effects of medications or the overall en of treatment) Environmental factors (e.g. polluted areas or busy roads) Other (please specify)

Q9-1: Definition of chronicity or long-term

Note inserted: None of the statements in Q9-1 reached \geq 70% consensus in round one. Most professionals defined long-term conditions as conditions that last for 6 months or more, whereas more than 70% of members of the public considered conditions lasting for 12

months or more as "long term". Having read a summary of responses, please answer Q9-1 again so that we could know if your responses have changed between two rounds.

How long does a condition have to be to count as long-term? Please choose one.*

Conditions lasting for three months or more
Conditions lasting for six months or more
Conditions lasting for twelve months or more
Conditions lasting for eighteen months or more

Q9-2: Recurrence or remission

Note inserted: In round one, more than 70% of panellists strongly agreed that conditions can be included in a multimorbidity measure if they are permanent in their effects, or are currently active or currently treated. There was no consensus for categories related to remitting/relapsing conditions. The first question below was asked in round 1 (reworded in response to feedback), and the second is based on suggestions in feedback. Please rate the degree to which you agree or disagree with the following statements.

Statement	Strongly disagree	Disagree	Neither agree nor disagree	Agree	Strongly agree	Don't know
Include conditions which may recur						
Include remitting-relapsing conditions which have happened during the last five years						
Other						

Q9-3: Treatment, care or surveillance

Note inserted: Some panellists suggested that some conditions that do not necessarily require treatment or care should be included because of significant impact on the individual. For example, people with arthritis might not be receiving current treatment/care but their quality of life could still be significantly affected. In addition, around 53.7% of panellists would include clinical risk factors many of which do not require treatment. Due to widely divergent views on Q9-3, we would like to ask you to rate the degree to which you agree or disagree with each of the following statements.

Statement	Strongly disagree	Disagree	Neither agree nor disagree	Agree	Strongly agree	Don't know
Include conditions which usually require current treatment, care or therapy (e.g. hypertension, diabetes, schizophrenia)						
Include conditions which usually require treatment, care or therapy at some point in the future even if not currently treated						
Include conditions which usually require surveillance (e.g. treated cancer or depression)						
Other						

Q9-4: Principles of selecting conditions based on impacts

Note inserted: In the professional version of survey, more than 70% of panellists strongly agreed that quality of life, physical disability and mental health should be taken into account when selecting conditions, and we are not asking you about these again. On the other hand, only risk of death was strongly agreed by more than 70% of members of the public.

Please rate the degree to which you agree or disagree with each of the following statements

Statement	Strongly disagree	Disagree	Neither agree nor disagree	Agree	Strongly agree
Include conditions which significantly increase risk of death					
Include conditions which cause frailty (general physical and/or mental weakness and vulnerability)					
Include conditions which significantly worsen self-perceived health status					
Include conditions which significantly increase treatment burden					

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Include conditions which could be impacted by or impact social deprivation and poverty			
Impact: Other			

Q10: Categorisation of conditions/counting methods

Note inserted: In round 1, we asked about how to account for complications of conditions and there was consensus (>80%) that all conditions that are currently active should be counted even if some are complications of other conditions. We are not asking this again. There was no consensus about whether to count broader categories of conditions or individual conditions. There was considerable free-text comment, including about the challenges of identifying specific conditions in electronic medical records. We have therefore revised the options for this question.

Please choose one option that you would prefer to use.

\square Broad disease category based on body systems (e.g. cardiovascular disease, mental health problems, skin conditions).					
☐ Individual conditions (e.g. myocardial infarction, angina, hypertension, depression, schizophrenia, eczema, psoriasis).					
☐ Grouping together similar conditions that are in the same body system and treated similarly (e.g. angina and myocardial infarction, or aplastic anaemia and sickle cell anaemia)					
☐ Other. Please explain:					
□ Don't know					

Q11: Data source

Multimorbidity has been measured by existing studies using either self-report or databases (e.g. medical records and administrative databases). Please rate the degree to which you agree or disagree the following statements.

Statement	Strongly disagree	Disagree	Neither agree nor disagree	Agree	Strongly agree
Conditions included for multimorbidity measurement should be the same in both self-report and databases (e.g. identifying a core set of conditions or condition framework).					
Conditions included in multimorbidity measures should differ between self-report and databases					
Other					

What conditions should be included?

Q12: System/disease domain

Note inserted: There was consensus that six body systems should always be included in a multimorbidity measure (mental health, cancer, cardiovascular, neurological, metabolic/endocrine, and musculoskeletal). We are not asking about these again in round 2. Please rate the degree to which the following systems/disease domains, you think, are important.

Condition	Exclude (not important)	Usually exclude (unless a good reason to include in a particular context)	Could include or exclude	Usually include (unless a good reason to exclude in particular context)	Always include (extremely important)	Don't know
Respiratory disease						
Haematological						
disease						
Digestive						
Urogenital						
Skin						
Ear, Nose and Throat						
(ENT)						
Oral						
Ophthalmology						
Chronic infections						

Note inserted: The number of conditions included in multimorbidity measurement varies substantially, which has led to the heterogeneity of multimorbidity prevalence. In the following section, we would like to ask you to rate a number of conditions that have been used for multimorbidity measurement, with the aim to identify a set of core conditions and conditions likely included for particular populations or regions.

Q13: Mental health

Note inserted: There was consensus that dementia and schizophrenia should always be included in multimorbidity measurement and we are not asking about these again. Please rate

the following mental health conditions again which allows us to see if your responses have changed between rounds. If you are not sure, please tick 'Don't know'.

Condition	Exclude (not important)	Usually exclude (unless a good reason to include in a particular context)	Could include or exclude	Usually include (unless a good reason to exclude in particular context)	Always include (extremely important)	Don't know
Anxiety						
Depression						
Substance misuse (including alcohol and drug dependence)						
Nicotine dependence						
Post-traumatic stress disorder (PTSD)						
Bipolar disorder						
Chronic insomnia						
Dissociative or personality disorders						
Eating disorders (including bulimia or anorexia)						
Learning disability						
Autism						
Obsessive compulsive disorder (OCD)						
Somatoform disorders (psychological disorder where a person experiences physical symptoms that cannot be explained by medical doctors)						
Attention deficit hyperactivity disorder (ADHD)						
Other (please specify either one or more conditions):						

Q14: Note inserted: There was consensus that solid organ cancers, haematological cancers and metastatic cancers should always be included in multimorbidity measurement and we are not asking about these again. Please rate the following cancer conditions again which allows us to see if your responses have changed between rounds. If you are not sure, please tick 'Don't know'.

Condition	Exclude (not important)	Usually exclude (unless a good reason to include in a particular context)	Could include or exclude	Usually include (unless a good reason to exclude in particular context)	Always include (extremely important)	Don't know
Melanoma						
Non-melanoma skin cancer						
Benign cancers (excluding benign skin lumps and bumps)						
Other (please specify either one or more conditions):						

Q15: Based on the panellists' responses, we have revised the options and would like to ask you to choose the method you would recommend (there were other comments for this question relating to recurrence and remission, but Q9 that you have already answered covers these).

	Count all cancers as one						
· 		magnetive of which avatoms that offer					
	Count individual cancers separately irrespective of which systems they affect						
	count gastric cancer and liver cancer	1 27					
	Count individual cancers separately i	f they affect different systems (e.g.					
panc	reatic cancer and lung cancer).						
	Other						

Q16: Note inserted: There was consensus that stroke, coronary heart disease, heart failure and peripheral artery disease should always be included in multimorbidity measurement and we are not asking about these again. Please rate the following cardiovascular conditions again which allows us to see if your responses have changed between rounds. If you are not sure, please tick 'Don't know'.

Condition	Exclude (not	Usually	Could	Usually	Always	Don't
	important)	exclude	include or	include	include	know
		(unless a	exclude	(unless a	(extremely	
		good reason		good reason	important)	

	to include in a particular context)	to exclude in particular context)	
Hypertension (treated)			
High blood pressure (untreated)			
Transient Ischaemic Attack (mini stroke)			
Dyslipidaemia/ Lipid disorder (treated)			
High cholesterol (untreated)			
Arrhythmia			
Heart valve problems (Including Rheumatic heart disease)			
Other (please specify either one or more conditions):			

Q17: Haematological conditions

Consistent with the results in Q12 that less than 70% of panellists would always include haematological system in multimorbidity measurement, none of the individual haematological conditions was rated by more than 70% of panellists as 'always include'.

Please rate the following haematological conditions as to whether they should be included or excluded. If you are not sure, please tick 'Don't know'.

Condition	Exclude (not important)	Usually exclude (unless a good reason to include in a particular context)	Could include or exclude	Usually include (unless a good reason to exclude in particular context)	Always include (extremely important)	Don't know
Anaemia (including pernicious anaemia, sickle cell anaemia, aplastic anaemia)						
Venous thrombotic disease						
Other (please specify either one or more conditions):						

Q18: Note inserted: There was consensus that Parkinson's disease, epilepsy, multiple sclerosis, and paralysis/ hemiplegia/ paraplegia (not including those caused by stroke) should always be included in multimorbidity measurement and we are not asking about these again. Please rate the following neurological conditions again which allows us to see if your responses have changed between rounds. If you are not sure, please tick 'Don't know'.

Condition	Exclude (not important)	Usually exclude (unless a good reason to include in a particular context)	Could include or exclude	Usually include (unless a good reason to exclude in particular context)	Always include (extremel y important)	Don't know
Migraine or other regular headache						
Peripheral neuropathy						
Other (please specify either one or more conditions):						

Q19: Note inserted: There was consensus that diabetes (any types) should always be included in multimorbidity measurement and we are not asking about these again. Please rate the following metabolic and endocrine conditions again which allows us to see if your responses have changed between rounds. If you are not sure, please tick 'Don't know'.

Condition	Exclude (not important)	Usually exclude (unless a good reason to include in a particular context)	Could include or exclude	Usually include (unless a good reason to exclude in particular context)	Always include (extremely important)	Don't know
Thyroid problems (including hypothyroidism and hyperthyroidism)						
Hyper and hypoparathyroidism						
Malnutrition (including protein energy deficiency)						
Addison's disease						
Other (please specify either one or more conditions):						

Q20: Note inserted: There was consensus that connective tissue disease (including rheumatoid arthritis or lupus) should always be included in multimorbidity measurement and we are not asking about these again. Please rate the following musculoskeletal conditions again which allows us to see if your responses have changed between rounds. If you are not sure, please tick 'Don't know'.

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Condition	Exclude (not important)	Usually exclude (unless a good reason to include in a particular context)	Could include or exclude	Usually include (unless a good reason to exclude in particular context)	Always include (extremely important)	Don't know
Osteoporosis						
Osteoarthritis						
Gout						
Long-term musculoskeletal problems due to injury (including low back pain, complex regional pain syndrome, neuropraxia)						
Other (please specify either one or more conditions):						

Q21: Note inserted: There was consensus that COPD, asthma and cystic fibrosis should always be included in multimorbidity measurement and we are not asking about these again. Please rate the following respiratory conditions again which allows us to see if your responses have changed between rounds.

Condition	Exclude (not important)	Usually exclude (unless a good reason to include in a particular context)	Could include or exclude	Usually include (unless a good reason to exclude in particular context)	Always include (extremely important)	Don't' know
Chronic/Allergic rhinitis (Nose becomes inflamed by allergens, such as dust, pollen, animal fur)						
Sleep apnoea						
Bronchiectasis						
Post-acute COVID- 19						
Other (please specify either one or more conditions):						

Q22: Note inserted: There was consensus that chronic liver disease and inflammatory bowel disease should always be included in multimorbidity measurement and we are not asking about these again. Please rate the following digestive conditions again which allows us to see if your responses have changed between rounds.

Condition	Exclude (not important)	Usually exclude (unless a good reason to include in a particular context)	Could include or exclude	Usually include (unless a good reason to exclude in particular context)	Always include (extremely important)	Don't know
Irritable bowel syndrome						
Diverticulosis (small pockets in the lining of the intestine)						
Pancreatic disease						
Peptic ulcer						
Gastroesophageal reflux						
Gall bladder problems (including gallstones)						
Other (please specify either one or more conditions):						

Q23: Note inserted: There was consensus that chronic kidney disease and end-stage kidney disease should always be included in multimorbidity measurement and we are not asking about these again. Please rate the following urogenital conditions again which allows us to see if your responses have changed between rounds.

Condition	Exclude (not important)	Usually exclude (unless a good reason to include in a particular context)	Could include or exclude	Usually include (unless a good reason to exclude in particular context)	Always include (extremely important)	Don't know
Kidney or bladder stones						
Chronic urinary tract infections (including chronic bladder infections)						
Urinary incontinence (loss of control over passing urine)						
Uterus (womb) problems (including prolapse and fibroid)						
Polycystic ovary syndrome						

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Prostatic			
hypertrophy (large			
prostate glands)			
Endometriosis			
Infertility			
Sexual dysfunction			
Other (please specify either one or more conditions):			

Q24: Note inserted: There was consensus that HIV/AIDS should always be included in multimorbidity measurement and we are not asking about these again. Please rate the following chronic infection conditions again which allows us to see if your responses have changed between rounds.

Condition	Exclude (not important)	Expect to usually exclude (unless a good reason to include in a particular context)	Could include or exclude	Expect to usually include (unless a good reason to exclude in particular context)	Always include (extremely important)
Tuberculosis					
Lyme disease					
Other (please specify either one or more conditions):					

Q25: Note inserted: None of the individual skin conditions was rated by more than 70% of panellists as 'always include'. Please rate the following skin conditions as to whether they should be included or excluded. If you are not sure, please tick 'Don't know'.

Condition	Exclude (not important)	Usually exclude (unless a good reason to include in a particular context)	Could include or exclude	Usually include (unless a good reason to exclude in particular context)	Always include (extremely important)	Don't know
Psoriasis						
Eczema						
Chronic urticarial (chronic hives)						
Other (please specify either one or more conditions):						

Q26: Note inserted: None of the individual ENT, eye and oral conditions was rated by more than 70% of panellists as 'always include'. Please rate the following ENT, eye and oral conditions as to whether they should be included or excluded. If you are not sure, please tick 'Don't know'.

Condition	Exclude (not important)	Usually exclude (unless a good reason to include in a particular context)	Could include or exclude	Usually include (unless a good reason to exclude in particular context)	Always include (extremely important)	Don't know
Hearing impairment or Deafness (that cannot be easily corrected with hearing aids)						
Meniere's disease (an ear condition that causes sudden attacks of vertigo)						
Ear, nose, throat disease (including chronic sinusitis)						
Vision impairment or Blindness (that cannot be easily corrected with glasses)						
Cataract						
Glaucoma						
Edentulism (having no teeth)						
Chronic gum disease						
Other (please specify either one or more conditions):						

Q27: Note inserted: Congenital condition was rated by less than 70% of panellists as 'always include'. Please rate the following congenital conditions as to whether they should be included or excluded. If you are not sure, please tick 'Don't know'.

Condition	Exclude (not important)	Usually exclude (unless a good reason to include in a particular context)	Could include or exclude	Usually include (unless a good reason to exclude in particular context)	Always include (extremely important)	Don't know
Congenital disease (conditions that						

babies are born with, including congenial heart disease, chromosomal aberrations)			
Other (please specify either one or more conditions):			

Q28: Note inserted: The following risk factors/symptoms have been included in multimorbidity measures by existing studies, but none of them was rated by more than 70% of panellists as 'always include'. Some suggested that those should be measured separately rather than as part of a multimorbdiity measure.

Having read a summary of the results from round one, please rate the following statement as to whether this type of conditions should be included or excluded again. If you are not sure, please tick 'Don't know'.

Condition	Exclude (not important)	Usually exclude (unless a good reason to include in a particular context)	Could include or exclude	Usually include (unless a good reason to exclude in particular context)	Always include (extremely important)	Don't know
Obesity (body mass index ≥30)						
Smoking						
Sedentary lifestyle						
Physical disability						
Dizziness or vertigo (without a specific diagnosis)						
Chronic cough (without a specific diagnosis)						
Post-sepsis syndrome						
Side effects of medications						
Treatment burden (the sum of all the hassles of taking medicines or attending appointments)						
Social deprivation and poverty						
Other (please specify either one or more conditions):						

Weighting

Q29: Weighted count versus simple count

For the following purposes, please choose which type of multimorbidity measure you would prefer to use.

Statement	Simple counts	Weighted measures	Both types of measures	Don't know
For the purpose of estimating the prevalence of multimorbidity, I prefer to use				
For the purpose of identifying and counting disease clusters, I prefer to use				
For the purpose of exploring/identifying predictors of multimorbidity (e.g. sociodemographic information), , I prefer to use				
For the purpose of exploring trajectories of multimorbidity (e.g. trends of multimorbidity prevalence and the number of conditions an individual has had), I prefer to use				
For the purpose of assessing the severity of disease burden, I prefer to use				
For the purpose of risk adjustment or outcome prediction, I prefer to use				

Q31: Outcomes

Note inserted: In round one, you were invited to rate which outcomes are important to be weighted against. Mortality and quality of life were considered very important in round one. However, adapting a measure that fits all purposes (predicting all outcomes) may not be feasible. It has also been suggested that healthcare use and costs should be separated from patient-centred outcomes, and distinct measures for these two outcomes are necessary. Another discrepancy identified from round one is that some preferred to use simple counts for predicting outcomes, but others preferred to use weighted measures. Instead of rating levels of importance, we would like to ask you to choose which type of multimorbidity measures you would prefer to use for predicting the following outcomes.

Outcome	Simple counts	Weighted	Both types of	No
		measures	measures	opinion/don't
				know

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Mortality				
Healthcare use (e.g. number of emergency admissions to hospital; outpatient appointments)				
Health-related quality of life				
Physical disability				
Frailty (general physical and/or mental weakness and vulnerability)				
Treatment burden (the sum of all the hassles of taking medicines or attending appointments)				
Healthcare costs (how much treatment and care for each individual costs)				
Self-perceived health status				
Other (please specify)				
Q32: Having reflected re please choose which wei (which allows to see if th	ghting method you	would prefer to us	se in a multimorbi	

; (choo	ing reflected results from round one and your re loose which weighting method you would prefer lows to see if the responses have changed over the	to use in a multimorbidity measu
		Use existing weighted indices.	
	□ out	Empirically derive weights based on the indiv tcome (e.g. use regression models to calculate w	•
	□ (e.g	Apply weights by defining specific criteria to g. set rules based on level of severity to grade ea	
		No opinion	
		Other	

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□ No

☐ Yes☐ No

Round-three survey in the professional panel

Socio-demographic information
Q1: Please describe the country where you are currently living or working.
Q2: Please check the category that best describes the type of work you do most of the time (choose one).
☐ Research
☐ Public policy or other public sector work
☐ Clinical Practice
☐ Teaching
□ Other
Q3: Please check the category that best describes the setting in which you work most of the time (choose one).
☐ Government
☐ Academia
☐ Hospital
☐ Primary care/General practice
☐ Social care sector
□ Other
Q4: Do you have multiple chronic conditions?
☐ Yes

Q5: Do you have family or friends who have multiple chronic conditions?

Round	three	professional	panel

Q6: To und choose all th	erstand multimorbidity, what population would you be more interested in (please hat apply)?
	General population
	Older people
	Middle-aged and older
	Socially-deprived population (including homeless people or drug users)
	Women
	Men
	Children
	Ethnic minority groups or indigenous populations
	People with disabilities
	Other

What is multimorbidity?

Note: In both the professional panel and public panel, there was consensus (\geq 70%) that someone had to have two or more chronic conditions in order to have multimorbidity. In addition, more than 70% of panellists also strongly agreed that conditions included in multimorbidity measurement should have the following characteristics: 1) permanent in their effects, 2) currently active, 3) lasting six months or longer, 4) requiring current treatment, care or therapy. However, less than 70% of professional panellists agreed that it was useful to conceptualise complex multimorbidity in addition to basic multimorbidity. It is therefore not entirely clear whether and how we should define complex multimorbidity. Please answer the following questions again.

Q7: Please rate the degree to which you agree or disagree with the following statement again so we can see if there is a change between round 2 and round 3.

	Strongly disagree	Disagree	Neither disagree nor agree	Agree	Strongly agree	Don't know
Do you agree that defining complex multimorbidity in addition to a core definition of simple multimorbidity is useful?						

Q8: Irrespective of whether you agree that complex morbidity is a useful concept, we wou	ıld
still like to invite you to define "complex multimorbidity" based on number of conditions	
(please choose one)	

	3+ 0	conditions	irrecn	ective	of how	many	hody	custems
l	$\rightarrow \pm c$	onamons	irresp	ecuve	OT HOV	v manv	DOUV	systems

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 □ 3+ conditions from 2+ body systems □ 3+ conditions from 3+ body systems □ 4+ conditions irrespective of how many body systems □ No opinion □ Other
Note: In the professional and public panel, there was no consensus on what additional patterns of conditions should be included in the definition of complex multimorbidity. We have added a few more statements based on panellists' comments.
Q8-1: Sub-question: Please choose the following statements that, you think, should be included in the definition of "complex multimorbidity". No other patterns that I would like to include Any combination of two or more conditions which includes both physical and mental health conditions Any combination of two or more conditions with significantly physical functional limitation Difficulty in managing illnesses due to social factors/social determinants of health (e.g. poverty). Any combination of two or more conditions and frailty Other
O9 Recurrence or remission

Note inserted: More than 70% of panellists in the professional and public panels agreed that conditions that may recur or remit could be included in the measurement of multimorbidity. However, it remains uncertain as to how this type of conditions should be defined in the concept of multimorbidity. Some suggested suggested take 'treatment' into account when deciding which remitting-relapsing conditions to include, and thus we have revised the statements.

Please rate the degree to which you agree or disagree with the following statements.

Statement	Strongly disagree	Disagree	Neither disagree nor agree	Agree	Strongly agree	Don't know
Include remitting-relapsing conditions that require ongoing treatment/therapy/care (e.g. depression, epilepsy)—newly added						
Include remitting-relapsing conditions which have relapsed during the last five years						

Round three professional panel

Q10: Treatment, care or surveillance

Note inserted: More than 70% of panellists strongly agreed to include conditions that require current treatment, care, or therapy, and this question will not be asked again in round three. Please rate the degree to which you agree or disagree with the following statements.

Statement	Strongly disagree	Disagree	Neither disagree nor agree	Agree	Strongly agree	Don't know
Include conditions which usually require treatment, care or therapy 'at some point in the future' even if not currently treated (one panellist suggested to conceptualise this as 'current risk to future health outcomes')						
Include conditions which usually require surveillance (one panellist suggested to conceptualise this as 'current health needs or complexity of providing care')						

Q11: Categorisation of conditions/counting methods

Please choose one option that you would prefer to use.	
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☐ Broad disease category based on body and mind system (e.g. cardiovascular disease, mental health problems, skin conditions).					
\square Individual conditions (e.g. TIA (mini stroke) and stroke are counted separately).					
☐ Grouping together similar conditions that are in the same category and treated similarly (e.g. group together 'Angina and Myocardial Infarction' or 'Aplastic anaemia and Sickle cell anaemia')					
☐ Other. Please explain:					
☑ Don't know					

Q12: Data source

Please rate the following statement again.

Round	three	professional	panel
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Statement	Strongly disagree	Disagree	Neither disagree nor agree	Agree	Strongly agree	Don't know
Conditions included for multimorbidity measurement should be the same/similar in both self-report and databases (e.g. identifying a core set of conditions or condition framework).						
Other comments						

What conditions should be included?

Q13: Note inserted: In cancer conditions, we added two new statements based on panellists' comments. Please rate the following cancer conditions as to whether they should be included or excluded. If you are not sure, please tick 'Don't know'.

Condition	Exclude (not important)	Usually exclude (unless a good reason to include in a particular context)	Could include or exclude	Usually include (unless a good reason to exclude in particular context)	Always include (extremely important)	Don't know
Treated cancer that requires surveillance						
Treated cancer that did not recur over the past 5 years and does not require surveillance						
Benign cerebral tumours (brain tumours that can cause functional limitations)						

Q14: Please choose the categorisation method you would prefer again.

☐ Count all primary cancers as one	
☐ Count individual primary cancers sep	parately irrespective of which systems they
affect (e.g. count gastric cancer and liver	cancer separately).
☐ Count individual primary cancers sep	parately only if they affect different systems
(e.g. pancreatic cancer and lung cancer).	
□ Other	
I I	

Round three professional panel

Q15: Note inserted: Conditions relevant to chronic pain and consequences of injury were revised based on panellists' comments. Please rate the following conditions as to whether they should be included or excluded. If you are not sure, please tick 'Don't know'.

Condition	Exclude (not important)	Usually exclude (unless a good reason to include in a particular context)	Could include or exclude	Usually include (unless a good reason to exclude in particular context)	Always include (extremel y important)	Don't know
Chronic or recurrent low back pain						
Chronic primary pain (defined as pain with no clear underlying condition but signflicant impact on the person)						
Long-term musculoskeletal problems due to injury (e.g. consequences of accidental injuries)						

Q16: Conditions newly added in round two

Please rate again the following conditions so we can see if your responses have changed between round two and round three

Condition	Exclude (not important)	Usually exclude (unless a good reason to include in a particular context)	Could include or exclude	Usually include (unless a good reason to exclude in particular context)	Always include (extremel y important)	Don't know
Addison's disease						
Bronchiectasis						
Post-acute covid 19 ("long COVID")						
Chronic Lyme disease						
Aneurysm						

Type of measures to use for a particular purpose

Q17: Which type of measures you would use for a particular purpose? (Please tick one or both - if you choose both a simple count and weighted measures, we would count either as being acceptable/usable for that purpose)

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Type of measures	Estimating the prevalence of multimorbidity	Identifying and counting disease clusters	Exploring trajectories of multimorbidity	Exploring/identifying predictors of multimorbidity	Assessing the severity of disease burden	Risk adjustment or outcome prediction
Prefer simple count of conditions						
Prefer weighted measure						
Either is acceptable/ useable						
Don't know						

Q18: Which type of measures you would use for prediction of a particular outcome? (Please tick one or both - if you choose both a simple count and weighted measures, we would count it as either being acceptable/usable for that purpose)

Type of measures	Risk of deat h	Health - related quality of life	Physical disabilit y	Frailt y	Treatmen t burden	Healthcar e use	Healthcar e costs	Self- perceive d health	Menta l health
Prefer simple count of conditions									
Prefer weighted measure									
Either is acceptable / useable									
Don't know									

Round-one survey in the public panel

Socio-demographic information
Q1: Please describe the country where you are currently living.
Q2: What is your age?
□ 18-34
□ 35-54
□ 55-64
$\square \geq 65$
☐ Prefer not to answer
Q3: What gender do you identify as?
☐ Female
☐ Male
☐ Other
☐ Prefer not to answer
Q4: We are seeking your views as a member of the public, but we know that some of you will have worked in healthcare or academia. Do you currently or did you previously work in academia, health and social care practice or healthcare policy?
□ Yes □ No
Q5: Do you have multiple chronic conditions?
☐ Yes ☐ No

What is multimorbidity?

☐ Yes ☐ No

Q7: Researchers have defined multimorbidity in many ways. Some define it as '2 or more long-term conditions', but others define it as '3 or more', '4 or more' or '5 or more' long-term conditions. How many long-term conditions do you think someone has to have in order to have multimorbidity? Please choose one.

Q6: Do you have family or friends who have multiple chronic conditions?

Round one public panel
□ 2 or more long-term conditions □ 3 or more long-term conditions □ 4 or more long-term conditions □ 5 or more long-term conditions □ Other, X or more long-term conditions (please only type a number in the box) □ OS: How would you define "condition" for the concent of multimorbidity? (please choose of
Q8: How would you define "condition" for the concept of multimorbidity? (please choose althat apply)
 □ Formal medical diagnoses (e.g. coronary heart disease, alcohol dependence) □ Clinical risk factors (e.g. obesity, high cholesterol) □ Symptoms that are not formal medical diagnoses (e.g. dizziness or fatigue) □ Health behaviours (e.g. smoking or exercise level) □ Health impact (e.g. disability or frailty) □ Social deprivation and poverty □ Consequences of treatment and care (e.g. side effects of medications or the overal burden of treatment) □ Other (please specify)
Q9: Researchers justify their choice of conditions in many different ways. Please answer the following questions on the principles for selection of chronic conditions in multimorbidity
 How long-term a condition is: Researchers vary in what they mean by "long-term". How long does a condition hav to be to count as long-term? Please choose one. □ Conditions lasting for three months or more □ Conditions lasting for six months or more □ Conditions lasting for twelve months or more □ Other, conditions lasting for X months or more (please only type a number for X in the box)
2) Whether a condition is currently active: Please rate the degree to which you agree or disagree with each of the following statements

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Statement	Strongly disagree	Disagree	Neither disagree nor agree	Agree	Strongly agree	Don't know		
Include conditions which are permanent in their effects (e.g. Parkinson's disease)								
Include conditions which are currently active or currently treated (e.g. asthma with intermittent wheeze; asthma using regular inhalers)								
Include conditions which may recur but happen rarely (e.g. people with a history of asthma or depression with no current symptoms and not currently on treatment)	⊠							
Other								
3) Whether condition is treated in healthcare: Please rate the degree to which you agree or disagree with each of the following statements								
Statement	Strongly disagree	Disagree	Neither disagree nor agree	Agree	Strongly agree	Don't know		

Statement	Strongly disagree	Disagree	Neither disagree nor agree	Agree	Strongly agree	Don't know
Only include conditions which usually require formal treatment or care (e.g. hypertension, diabetes, schizophrenia)						
Other						

4) The impact of the condition on a range of outcomes: Please rate the degree to which you agree or disagree with each of the following statements

Statement	Strongly agree	Agree	Neither agree nor disagree	Disagree	Strongly disagree	Don't know

Round	one	public	c pane

Include conditions which significantly increase risk of death										
Include conditions which significantly reduce quality of life										
Include conditions which cause significant physical disability										
Include conditions which cause frailty (general physical and/or mental weakness and vulnerability)										
Include conditions which significantly worsen mental health										
Include conditions which significantly worsen how people perceive their general health										
Other impact										
Q10: Researchers vary in how detailed their definitions of 'condition' are. For example, they might be very broad – one category for "lung disease". Or they might be more detailed – separately count "asthma", "chronic obstructive pulmonary disease" and so on. Please choose one option and explain your choice if you like. □ Broad disease category based on body and mind system (e.g. cardiovascular disease, mental health problems, skin conditions). Please explain:										
☐ Individual condition schizophrenia, eczema,	. •	•		sion, depress	sion,					
	n:									

Round one public panel
☐ Don't know
Q11: Some conditions included in multimorbidity measures are closely linked to each other. In particular, some conditions can be caused by other conditions. For example, heart attacks can lead to heart failure. Diabetes can lead to kidney failure. How do you think researchers should count conditions in this situation? Please choose one option and explain your choice if you like.
☐ Count all of the conditions that are currently active. Please explain:
☐ Only count the complications (e.g. if people with heart attack develop heart failure, we would count only heart failure). Please explain:
☐ Count the primary health condition (e.g. if people with diabetes develop kidney failure, we would only count diabetes). Please explain:
☐ Other. Please explain:
⊠ Don't know

What conditions should be included?

In our review of over 500 research studies, we identified all the conditions which researchers have counted when measuring multimorbidity. There is very large variation in which conditions researchers choose. Only seven conditions were counted by more than half of studies. Many conditions were rarely counted. We would like you to rate conditions that you think are more or less important to include in multimorbidity measures. There are now a set of questions organised by body system.

Q12: We are starting with body systems/disease domains. Please rate how important you think it is to include the following systems/domains in a multimorbidity measure.

Condition	Exclude (not importan t)	Usually exclude (unless a good reason to include in a particular context)	Could include or exclude	Usually include (unless a good reason to exclude in particular context)	Always include (extremely important)	Don't know
Mental health						
Cancer						
Cardiovascular						
Blood disease (which affects formation of blood cells and blood clots)						
Neurological disease (which affects the nervous system)						
Metabolic and Endocrine (which affects metabolic regulation)						
Musculoskeletal (which affects joints, bones and muscles)						
Digestive (which affects stomach, bowels, liver, pancreas and gallbladder)						
Urogenital (which affects urinary and genital organs)						
Skin						
Ear, Nose and Throat (ENT)						
Oral						
Ophthalmology (which affects eyes)						
Chronic infections	П	П	П	П	П	П

Q13: Please rate how important you think it is to include the listed mental health conditions in a multimorbidity measure. If you are not sure, just tick 'Don't know'.

Condition	Exclude (not important)	Usually exclude (unless a good reason to include in a particular context)	Could include or exclude	Usually include (unless a good reason to exclude in particular context)	Always include (extremely important)	Don't know
Anxiety						
Depression						
Dementia						
Schizophrenia						

Substance misuse (including alcohol and drug dependence)			
Nicotine dependence			
Post-traumatic stress disorder (PTSD)			
Bipolar disorder			
Chronic insomnia			
Dissociative or			
personality disorders (altering to be a person with different			
personality and the alters are imaginary)			
Eating disorders (including bulimia or anorexia)			
Learning disability			
Autism			
Obsessive compulsive disorder (OCD)			
Somatoform disorders (psychological disorder where a person experiences physical symptoms that cannot be explained by medical doctors)			
Attention deficit hyperactivity disorder (ADHD)			
Other (please specify either one or more conditions):			

Q14: Please rate how important you think it is to include the listed cancers in a multimorbidity measure.

Condition	Exclude (not	Usually	Could	Usually	Always	Don't
	important)	exclude	include or	include	include	know
		(unless a	exclude	(unless a		

	good reason to include in a particular context)	good reason to exclude in particular context)	(extremely important)	
Solid organ cancers (e.g. lung, colon, prostate, breast etc)				
Haematological cancers (cancer that affects blood, e.g. leukaemia, lymphoma, myeloma)				
Melanoma (malignant skin cancer)				
Non-melanoma skin cancer				
Benign cancers (excluding benign skin lumps and bumps)				
Metastatic cancers (which have spread to other parts of the body)				
Other (please specify either one or more conditions):				

Q15: For cancers, some researchers count every cancer a person has had. Others only count 'cancer' once even if someone has had more than one type of cancer. Which methods would you recommend. Please choose one:

Count an eancers as one
Count individual cancers separately.
Other

Q16: Please rate how important you think it is to include the listed cardiovascular conditions in a multimorbidity measure.

Condition	Exclude (not	Usually	Could	Usually	Always	Don't
	important)	exclude	include or	include	include	know
		(unless a	exclude	(unless a	(extremely	
		good reason		good reason	important)	
		to include in		to exclude in		
		a particular		particular		
		context)		context)		

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Hypertension			
Stroke			
Transient Ischaemic Attack (mini stroke)			
Lipid disorder (e.g. high cholesterol)			
Coronary artery disease (heart attack or angina)			
Heart failure			
Peripheral artery disease (circulation problems in the legs)			
Heart rhythm problem (irregular or very fast heart beats)			
Heart valves problem (leaking or tight heart valves)			
Aneurysm (a weakening or bulging of an artery wall)			
Other (please specify either one or more conditions):			

Q17: Please rate how important you think it is to include the listed haematological conditions in a multimorbidity measure.

Condition	Exclude (not important)	Usually exclude (unless a good reason to include in a particular context)	Could include or exclude	Usually include (unless a good reason to exclude in particular context)	Always include (extremely important)	Don't know
Anaemia						
Venous thrombotic disease (blood clots formed inside a blood vessel that can block circulation)						
Other (please specify either one or more conditions):						

Q18: Please rate how important you think it is to include the listed neurological conditions in a multimorbidity measure.

Condition	Exclude (not important)	Usually exclude (unless a good reason to include in a particular context)	Could include or exclude	Usually include (unless a good reason to exclude in particular context)	Always include (extremel y important)	Don't know
Parkinson's disease						
Epilepsy (fits/ seizures)						
Chronic pain						
(including chronic low						
back pain and other						
neurological pain)						
Migraine or other						
regular headache						
Multiple sclerosis						
Peripheral neuropathy						
(damage to the nerves						
going to legs or arms)						
Paralysis/ Hemiplegia/						
Paraplegia (not						
including those caused						
by stroke)						
Other (please specify						
either one or more						
conditions):						

Q19: Please rate how important you think it is to include the listed metabolic/endocrine/nutritional conditions in a multimorbidity measure.

Condition	Exclude (not important)	Usually exclude (unless a good reason to include in a particular context)	Could include or exclude	Usually include (unless a good reason to exclude in particular context)	Always include (extremely important)	Don't know
Diabetes (any type)						
Thyroid problems						
Malnutrition						
Other (please specify either one or more conditions):						

Q20: Please rate how important you think it is to include the listed musculoskeletal conditions in a multimorbidity measure.

Condition	Exclude (not important)	Usually exclude (unless a good reason to include in a particular context)	Could include or exclude	Usually include (unless a good reason to exclude in particular context)	Always include (extremely important)	Don't know
Osteoporosis (thinning of the bones)						
Osteoarthritis (wearing and tearing of joints that leads to inflammation and degeneration						
Connective tissue disease (autoimmune disease that can affect our bones, cartilages or joints, such as rheumatoid arthritis or lupus)						
Gout (crystals that formed inside or around joints, causing pain)						
Long-term musculoskeletal problems due to injury (including hip fracture)						
Other (please specify either one or more conditions):						

Q21: Please rate how important you think it is to include the listed respiratory conditions in a multimorbidity measure.

Condition	Exclude (not important)	Usually exclude (unless a good reason to include in a particular context)	Could include or exclude	Usually include (unless a good reason to exclude in particular context)	Always include (extremely important)	Don't' know
Chronic obstructive pulmonary disease (COPD – smoking related lung damage)						
Asthma						
Sleep apnoea (breathing						

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problems when			
asleep)			
Cystic fibrosis			
(inherited condition			
that damages lungs)			
Chronic/Allergic			
rhinitis (long-term			
nasal			
inflammation)			
Other (please			
specify either one			
or more			
conditions):			

Q22: Please rate how important you think it is to include the listed digestive conditions in a multimorbidity measure.

Condition	Exclude (not important)	Usually exclude (unless a good reason to include in a particular context)	Could include or exclude	Usually include (unless a good reason to exclude in particular context)	Always include (extremely important)	Don't know
Chronic liver disease (including liver cirrhosis, liver failure and chronic hepatitis)						
Inflammatory bowel disease (including ulcerative colitis and Crohn's disease)						
Irritable bowel syndrome						
Diverticular disease (small pockets in the lining of the intestine)						
Pancreatic disease						
Peptic ulcer (including gastric/ stomach ulcers and duodenal ulcers)						
Gastroesophageal reflux (acid reflux and heartburn)						
Gall bladder problems (including gallstones)						
Other (please specify either one or more conditions):						

Q23: Please rate how important you think it is to include the listed urogenital conditions in a multimorbidity measure.

Condition	Exclude (not important)	Usually exclude (unless a good reason to include in a particular context)	Could include or exclude	Usually include (unless a good reason to exclude in particular context)	Always include (extremely important)	Don't know
Chronic kidney disease						
End-stage kidney disease (including kidney dialysis and transplant)						
Kidney or bladder stones						
Chronic urinary tract infections (including chronic bladder infections)						
Urinary incontinence (loss of control over passing urine)						
Uterus (womb) problems (including prolapse and fibroid)						
Polycystic ovary syndrome						
Prostatic hypertrophy (large prostate glands)						
Endometriosis						
Infertility						
Sexual dysfunction						
Other (please specify either one or more conditions):						

Q24: Please rate how important you think it is to include the listed chronic infectious conditions in a multimorbidity measure.

Condition	Exclude (not	Expect to	Could	Expect to	Always
	important)	usually	include or	usually	include
		exclude	exclude	include	(extremely
		(unless a		(unless a	important)
		good reason		good reason	
		to include in a		to exclude in	
		particular		particular	
		context)		context)	

HIV/AIDS			
Tuberculosis			
Other (please specify			
either one or more conditions):			

Q25: Please rate how important you think it is to include the listed skin conditions in a multimorbidity measure.

Condition	Exclude (not important)	Usually exclude (unless a good reason to include in a particular context)	Could include or exclude	Usually include (unless a good reason to exclude in particular context)	Always include (extremely important)	Don't know
Psoriasis						
Eczema						
Chronic urticarial (chronic hives)						
Other (please specify either one or more conditions):						

Q26: Please rate how important you think it is to include the listed ENT, eye & oral conditions in a multimorbidity measure.

Condition	Exclude (not important)	Usually exclude (unless a good reason to include in a particular context)	Could include or exclude	Usually include (unless a good reason to exclude in particular context)	Always include (extremely important)	Don't know
Hearing impairment or deafness						
Meniere's disease (an ear condition that causes sudden attacks of vertigo)						
Ear, nose, throat disease (including chronic sinusitis)						
Vision impairment or blindness						
Cataract						
Glaucoma						
Edentulism (having no teeth)						

Chronic gum			
disease			
Other (please			
specify either one			
or more			
conditions):			

Q27: Please rate how important you think it is to include the listed congenital conditions in a multimorbidity measure.

Condition	Exclude (not important)	Usually exclude (unless a good reason to include in a particular context)	Could include or exclude	Usually include (unless a good reason to exclude in particular context)	Always include (extremely important)	Don't know
Congenital disease (conditions that babies are born with, including congenial heart disease, genital anomaly)						
Other (please specify either one or more conditions):						

Q28: Please rate how important you think it is to include the listed risk factors/health behaviour/symptoms/syndromes in a multimorbidity measure.

Condition	Exclude (not important)	Usually exclude (unless a good reason to include in a particular context)	Could include or exclude	Usually include (unless a good reason to exclude in particular context)	Always include (extremely important)	Don't know
Obesity (body mass index ≥30)						
Smoking						
High blood pressure (untreated)						
High cholesterol (untreated)						
Sedentary lifestyle (spending most of the day sitting)						
Physical disability						
Dizziness (without a specific diagnosis)						

(without a specific				Ц	Ц	Ц
diagnosis)						
Post-sepsis						
syndrome						
Side effects of						
medications						
Treatment burden						
(the sum of all the						
hassles of taking						
medicines or						
attending appointments)						
Social deprivation	П	П	П	П	П	П
and poverty						
Other (please						
specify either one						
or more						
conditions):						

Weighting

Q29: Most researchers define multimorbidity by just counting how many conditions someone has (a "simple count" of conditions). Others use a "weighted" count to estimate multimorbidity burden and predict outcomes (e.g. death, hospitalisation, quality of life). For example, a simple count would say that hay fever + heart attack + back pain = 3 conditions. A weighted measure that gave more weight to risk of death might count hay fever = 0.5, high blood pressure = 3, back pain = 0.5, total score = 4 weighted for risk of death. However a weighted measure that gave more weight to quality of life might count hay fever = 1, high blood pressure = 0.5, and back pain = 2, total score = 3.5 weighted for quality of life. In theory, weighted scores are better than simple counts at predicting if someone is going to have the outcome focused on.

A key issue for weighted measures, is which outcomes to focus on. Researchers vary in which outcomes they think are most important. Professionals and patients also vary in which outcomes they think are important.

We would like to know which outcomes you think are most important.

Outcome	Not at all important	Slightly important	Important	Sufficiently important	Very important	No opinion
Death						
Healthcare use (e.g. number of emergency admissions to hospital; outpatient						
appointments)						
Quality of life						
Physical disability						
Frailty (general physical and/or mental weakness and vulnerability)						

Mental nealth	Ш		Ш	Ш
Treatment burden				
(the sum of all the				
hassles of taking				
medicines or				
attending				
appointments)				
Healthcare costs				
(how much				
treatment and care				
for each individual				
costs)				
How people				
perceive their				
general health				
overall (e.g.				
whether it is				
excellent, very				
good, fair, poor,				
very poor)				
Other (please				
specify)				
Other (please				
specify)				

Round-two survey in the public panel

Socio-demographic information Q1: Please describe the country where you are currently living. Q2: What is your age? □ 18-34 □ 35-54 □ 55-64 $\square \geq 65$ ☐ Prefer not to answer Q3: What gender do you identify as? ☐ Female ☐ Male \Box Other ☐ Prefer not to answer Q4: We are seeking your views as a member of the public, but we know that some of you will have worked in healthcare or academia. Do you currently or did you previously work in academia, health and social care practice or healthcare policy? \square Yes □ No Q5: Do you have multiple chronic conditions? ☐ Yes ☐ No

What is multimorbidity?

☐ Yes ☐ No

Note inserted: In round one, more than 80% of panellists defined multimorbidity as the co-occurrence of two or more long-term conditions. This question has been agreed by more than 70% of panellists and thus will not be asked again in round two.

Q6: Do you have family or friends who have multiple chronic conditions?

There were also numerous free text comments, including some which suggested that there should be distinctions made between 'simple' and 'complex' multimorbidity. In this section, we are therefore asking you some new questions about 'simple' and 'complex' multimorbidity.

Q7-1: Do you agree that defining complex multimorbidity in addition to a core definition of simple multimorbidity is useful?							
Strongly disagree/Disagree/Neither disagree nor agree/Agree/Strongly agree/Don't know							
Q7-2: Irrespective of whether you agree, how would you define "complex multimorbidity" based on number of conditions. (please choose one)							
☐ 3+ conditions irrespective of how many body systems ☐ 3+ conditions from 3+ body systems ☐ 4+ conditions irrespective of how many body systems ☐ 4+ conditions from 4+ body systems ☐ 5+ conditions irrespective of how many body systems ☐ 5+ conditions from 5+ body systems ☐ Other+ conditions from+ body systems ☐ Other+ conditions from+ body systems Q7-3: Sub-question: In addition to the above option you choose, please describe if there are other statements that, you think, should be included in the definition of "complex"							
multimorbdiity". Any combination of 2+ conditions which includes both physical and mental health conditions Other							
Q8: Defining 'condition' for multimorbidity measurement							
Note inserted: Based on the results from round one, only medical diagnoses reached \geq 70% consensus. Having read a summary of the results from round one, we would like you to answer the question again.							
How would you define "condition" for the concept of multimorbidity? (please choose all that apply)							
 □ Formal medical diagnoses (e.g. coronary heart disease, alcohol dependence) □ Clinical risk factors (e.g. obesity, high cholesterol) □ Symptoms that are not formal medical diagnoses (e.g. dizziness or fatigue) □ Health behaviours (e.g. smoking or exercise level) 							

☐ Health impact (e.g. disability or frailty)
☐ Social deprivation and poverty
☐ Consequences of treatment and care (e.g. side effects of medications or the overall
burden of treatment)
☐ Other (please specify)

Q9: Recurrence or remission

Round two public panel

Note-inserted: In round one, more than 70% of members of the public defined long-term conditions as conditions that last for 6 months or more, and strongly agreed that conditions can be included in a multimorbidity measure if they are permanent in their effects. There was no consensus for categories related to active conditions and remitting/relapsing conditions. The first two questions below were asked in round 1 (reworded in response to feedback), and the third is based on suggestions in feedback. Please rate the degree to which you agree or disagree with the following statements.

Statement	Strongly disagree	Disagree	Neither agree nor disagree	Agree	Strongly agree	Don't know
Include conditions which are currently active						
Include conditions which may recur						
Include remitting-relapsing conditions which have happened during the last five years						
Other						

Q10: Treatment, care or surveillance

Note inserted: Some panellists suggested that some conditions that do not necessarily require treatment or care should be included because of significant impact on the individual. For example, people with arthritis might not be receiving current treatment/care but their quality of life could still be significantly affected. In addition, around 60% of panellists would

include clinical risk factors many of which do not require treatment. Due to widely divergent views on Q10, we would like to ask you to rate the degree to which you agree or disagree with each of the following statements.

Statement	Strongly disagree	Disagree	Neither agree nor disagree	Agree	Strongly agree	Don't know
Include conditions which usually require current treatment, care or therapy (e.g. hypertension, diabetes, schizophrenia), diabetes, schizophrenia)						
Include conditions which usually require treatment, care or therapy at some point in the future even if not currently treated						
Include conditions which usually require surveillance (e.g. treated cancer or depression)						
Other						

Q11: Principles of selecting conditions based on impacts

Note inserted: In the public version of survey, more than 70% of panellists strongly agreed that death should be taken into account when selecting conditions, and we are not asking you about this again. On the other hand, quality of life, physical disability and mental health were strongly agreed by more than 70% of professionals.

In addition, social deprivation factor is listed here to explore your views on whether it should be one of the determinants of condition selection for multimorbidity measurement. In other parts of the survey, a third of panellists voted to exclude social deprivation as a "condition" but there was considerable comment about its importance, including that it could be taken into account when selecting conditions for multimorbidity measurement.

Please rate the degree to which you agree or disagree with each of the following statements

Statement	Strongly disagree	Disagree	Neither agree nor disagree	Agree	Strongly agree	Don't know
Include conditions which significantly reduce quality of life						

Include conditions which cause significant physical disability			
Include conditions which cause frailty (general physical and/or mental weakness and vulnerability)			
Include conditions which significantly worsen mental health			
Include conditions which significantly worsen how people perceive their general health			
Include conditions which significantly increase treatment burden (the sum of all the hassles of taking medicines or attending appointments			
Include conditions which could be impacted by or impact social deprivation and poverty			
Other impact			

Q12: Categorisation of conditions/counting methods

In round 1, we asked about how to account for complications of conditions and there was consensus (>70%) that all conditions that are currently active should be counted even if some are complications of other conditions. We are not asking this again. There was considerable free-text comment, including about the challenges of identifying specific conditions in electronic medical records. We have therefore revised the options for this question.

Please choose one option that you would prefer to use.

☐ Broad disease category based on body systems (e.g. cardiovascular disease, mental health problems, skin conditions).
☐ Individual conditions (e.g. myocardial infarction, angina, hypertension, depression, schizophrenia, eczema, psoriasis).
☐ Grouping together similar conditions that are in the same body system and treated similarly (e.g. angina and myocardial infarction, or aplastic anaemia and sickle cell anaemia)

Round two public panel						
☐ Other. Please explain:						
□ Don't know						

Q13: Multimorbidity has been measured by existing studies using either medical records or public self-reported surveys. Please rate the degree to which you agree or disagree the following statements.

Statement	Strongly disagree	Disagree	Neither agree nor disagree	Agree	Strongly agree	Don't know
Conditions included for multimorbidity measurement should be the same in both self-report and databases (e.g. identifying a core set of conditions or condition framework).						
Conditions included in multimorbidity measures should differ between self- report and databases						
Other						

What conditions should be included?

Q14: System/disease domain

Note inserted: There was no consensus that which body systems should always be included in a multimorbidity measure.

Please rate the degree to which the following systems/domains, you think, are important.

Condition	Exclude (not importan t)	Usually exclude (unless a good reason to include in a particular context)	Could include or exclude	Usually include (unless a good reason to exclude in particular context)	Always include (extremely important)	Don't know
Mental health						
Cancer						
Cardiovascular						
Blood disease (which affects formation of blood cells and blood clots)						
Respiratory disease						

Neurological disease				
(which affects the				
nervous system)				
Metabolic and				
Endocrine (which				
affects metabolic				
regulation. E.g.				
diabetes, thyroid				
problem)				
Musculoskeletal				
(which affects joints,				
bones and muscles)				
Digestive (which				
affects stomach,				
bowels, liver,				
pancreas and				
gallbladder)				
Urogenital (which				
affects urinary and				
genital organs)				
Skin				
Ear, Nose and Throat				
(ENT)				
Oral				
Ophthalmology				
(which affects eyes)	 			
Chronic infections				

Note inserted: The number of conditions included in multimorbidity measurement varies substantially, which has led to the heterogeneity of multimorbidity prevalence. In the following section, we would like to ask you to rate a number of conditions that have been used for multimorbidity measurement, with the aim to identify a set of core conditions and conditions likely included for particular populations or regions.

Q15: Mental health

Note inserted: In mental health domain, none of the individual mental health conditions was rated by more than 70% of panellists as 'always include'. On the other hand, in the professional panel, there was consensus that dementia and schizophrenia should always be included in multimorbidity measurement.

Please rate the following mental health conditions as to whether they should be included or excluded, which allows us to see if your responses have changed between rounds. If you are not sure, please tick 'Don't know'.

Condition	Exclude (not important)	Usually exclude (unless a good reason to include in a particular context)	Could include or exclude	Usually include (unless a good reason to exclude in particular context)	Always include (extremely important)	Don't know
Anxiety						
Depression						

Dementia			
Schizophrenia			
Substance use disorder(includin g alcohol and drug dependence)			
Nicotine dependence			
Post-traumatic stress disorder			
(PTSD)			
Bipolar disorder			
Chronic insomnia			
Dissociative or personality disorders (altering to be a person with different personality and the alters are			
imaginary)			
Eating disorders (including bulimia or anorexia)			
Learning disability			
Autism			
Obsessive compulsive disorder (OCD)			
Somatoform disorders (psychological disorder where a person experiences physical symptoms that cannot be explained by medical doctors)			
Attention deficit hyperactivity disorder (ADHD)			
Other (please specify either one or more conditions):			

Q16: In cancer domain, none of the individual cancer conditions was rated by more than 70% of panellists as 'always include'. On the other hand, in the professional panel, there was consensus that solid organ cancers, haematological cancers and metastatic cancers should always be included in multimorbidity measurement. Please rate the following cancer

conditions as to whether they should be included or excluded. If you are not sure, p	olease	tick
'Don't know'.		

Condition	Exclude (not important)	Usually exclude (unless a good reason to include in a particular context)	Could include or exclude	Usually include (unless a good reason to exclude in particular context)	Always include (extremely important)	Don't know
Solid organ cancers (e.g. lung, colon, prostate, breast etc)						
Haematological cancers (cancer that affects blood, e.g. leukaemia, lymphoma, myeloma)						
Melanoma (serious form of malignant skin cancer)						
Non-melanoma skin cancer						
Benign cancers (excluding benign skin lumps and bumps)						
Metastatic cancers (which have spread to other parts of the body)						
Other (please specify either one or more conditions):						

Q17: Based on the panellists' responses, we have you to choose the method you would recommend	*
(e.g. count gastric cancer and liver cancer	irrespective of which systems they affect separately). if they affect different systems (e.g. breast
□ Other	

Q18: Note inserted: In cardiovascular domain, none of the individual cardiovascular conditions was rated by more than 70% of panellists as 'always include'. On the other hand, in the professional panel, there was consensus that stroke, coronary heart disease, heart failure and peripheral artery disease should always be included in multimorbidity measurement. Please rate the following cardiovascular conditions as to whether they should be included or excluded. If you are not sure, please tick 'Don't know'.

Condition	Exclude (not important)	Usually exclude (unless a good reason to include in a particular context)	Could include or exclude	Usually include (unless a good reason to exclude in particular context)	Always include (extremely important)	Don't know
Hypertension (treated)						
High blood pressure (untreated)						
Stroke						
Transient Ischaemic Attack (mini stroke)						
High cholesterol (treated)						
High cholesterol (untreated)						
Coronary artery disease (heart attack or angina)						
Heart failure						
Peripheral artery disease (circulation problems in the legs)						
Heart rhythm problem (irregular or very fast heart beats)						
Heart valves problem (leaking or tight heart valves)						
Aneurysm (a weakening or bulging of an artery wall)						
Other (please specify either one or more conditions):						

Q19: Consistent with the results in Q14 that less than 70% of panellists would always include haematological system in multimorbidity measurement, none of the individual haematological conditions was rated by more than 70% of panellists as 'always include'.

Please rate the following haematological conditions as to whether they should be included or excluded. If you are not sure, please tick 'Don't know'.

Condition	Exclude (not important)	Usually exclude (unless a good reason to include in a particular context)	Could include or exclude	Usually include (unless a good reason to exclude in particular context)	Always include (extremely important)	Don't know
Anaemia						
Venous thrombotic disease (blood clots formed inside a blood vessel that can block circulation)						
Other (please specify either one or more conditions):						

Q20: Note inserted: In neurological domain, none of the individual neurological conditions was rated by more than 70% of panellists as 'always include'. On the other hand, in the professional panel, there was consensus that Parkinson's disease, epilepsy, multiple sclerosis, and paralysis/ hemiplegia/ paraplegia (not including those caused by stroke) should always be included in multimorbidity measurement.

Please rate the following neurological conditions as to whether they should be included or excluded. If you are not sure, please tick 'Don't know'.

Condition	Exclude (not important)	Usually exclude (unless a good reason to include in a particular context)	Could include or exclude	Usually include (unless a good reason to exclude in particular context)	Always include (extremel y important)	Don't know
Parkinson's disease						
Epilepsy (fits/ seizures)						
Migraine or other						
regular headache						
Multiple sclerosis						
Peripheral neuropathy						
(damage to the nerves						
going to legs or arms)						
Paralysis/ Hemiplegia/						
Paraplegia (not						
including those caused						
by stroke)						

Round	l two	pub	lic	pane	l
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Other (please specify			
either one or more			
conditions):			

Q21: Note inserted: In metabolic and endocrine domain, none of the individual conditions was rated by more than 70% of panellists as 'always include'. On the other hand, in the professional panel, there was consensus that diabetes (any types) should always be included in multimorbidity measurement.

Please rate the following metabolic and endocrine conditions as to whether they should be included or excluded. If you are not sure, please tick 'Don't know'.

Condition	Exclude (not important)	Usually exclude (unless a good reason to include in a particular context)	Could include or exclude	Usually include (unless a good reason to exclude in particular context)	Always include (extremely important)	Don't know
Diabetes (including type 1 and type 2)						
Thyroid problems (abnormal production of thyroid hormones)						
Malnutrition (including protein energy deficiency, commonly in low-income countries)						
Addison's disease (adrenal insufficiency)						
Other (please specify either one or more conditions):						

Q22: Note inserted: In musculoskeletal domain, none of the individual conditions was rated by more than 70% of panellists as 'always include'. On the other hand, in the professional panel, there was consensus that connective tissue disease (including rheumatoid arthritis or lupus) should always be included in multimorbidity measurement.

Please rate the following musculoskeletal conditions as to whether they should be included or excluded. If you are not sure, please tick 'Don't know'.

Condition	Exclude (not important)	Usually exclude (unless a good reason to include in a particular context)	Could include or exclude	Usually include (unless a good reason to exclude in particular context)	Always include (extremely important)	Don't know
Osteoporosis (thinning of the bones)						

Round two	dua c	lic pa	ane
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Osteoarthritis			
(wearing and tearing			
of joints that leads to			
inflammation and			
degeneration			
Connective tissue			
disease (autoimmune			
disease that can affect			
our bones, cartilages			
or joints, such as			
rheumatoid arthritis or			
lupus)			
Gout (crystals that			
formed inside or			
around joints, causing			
pain)			
Long-term			
musculoskeletal			
problems due to			
injury (including low			
back pain)			
Other (please specify			
either one or more			
conditions):			

Q23: Note inserted: In respiratory domain, none of the individual conditions was rated by more than 70% of panellists as 'always include'. On the other hand, in the professional panel, there was consensus that COPD, asthma and cystic fibrosis should always be included in multimorbidity measurement.

Please rate the following respiratory condition as to whether it should be included or excluded. If you are not sure, please tick 'Don't know'.

Condition	Exclude (not important)	Usually exclude (unless a good reason to include in a particular context)	Could include or exclude	Usually include (unless a good reason to exclude in particular context)	Always include (extremely important)	Don't' know
Chronic obstructive pulmonary disease (COPD – smoking related lung damage)						
Asthma						
Sleep apnoea (breathing problems when asleep)						
Cystic fibrosis (inherited condition that damages lungs)						

Round tv	dua ov	lic	panel
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Chronic/Allergic			
rhinitis (long-term			
nasal			
inflammation)			
Bronchiectasis			
(consistent			
infection in the			
lungs)			
Post-acute COVID-			
19			
Other (please			
specify either one			
or more			
conditions):			
·			

Q24: Note inserted: In digestive domain, none of the individual conditions was rated by more than 70% of panellists as 'always include'. On the other hand, in the professional panel, there was consensus that chronic liver disease and inflammatory bowel disease should always be included in multimorbidity measurement.

Please rate the following digestive conditions as to whether they should be included or excluded. If you are not sure, please tick 'Don't know'.

Condition	Exclude (not important)	Usually exclude (unless a good reason to include in a particular context)	Could include or exclude	Usually include (unless a good reason to exclude in particular context)	Always include (extremely important)	Don't know
Chronic liver disease (including liver cirrhosis, liver failure and chronic hepatitis)						
Inflammatory bowel disease (including ulcerative colitis and Crohn's disease)						
Irritable bowel syndrome						
Diverticulosis (small pockets in the lining of the intestine)						
Pancreatic disease						
Upper digestive disease—Peptic ulcer (including gastric/ stomach ulcers and duodenal ulcers) and Gastroesophageal reflux (acid reflux and heartburn)						
Gall bladder problems (including gallstones)						

Round t	two	bubl	ic	panel
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Other (please specify			
either one or more			
conditions):			

Q25: Note inserted: In urogenital domain, none of the individual conditions was rated by more than 70% of panellists as 'always include'. On the other hand, in the professional panel, there was consensus that chronic kidney disease and end-stage kidney disease should always be included in multimorbidity measurement.

Please rate the following urogenital conditions as to whether they should be included or excluded. If you are not sure, please tick 'Don't know'.

Condition	Exclude (not important)	Usually exclude (unless a good reason to include in a particular context)	Could include or exclude	Usually include (unless a good reason to exclude in particular context)	Always include (extremely important)	Don't know
Chronic kidney disease						
End-stage kidney disease (including kidney dialysis and transplant)						
Kidney or bladder stones						
Chronic urinary tract infections (including chronic bladder infections)						
Urinary incontinence (loss of control over passing urine)						
Uterus (womb) problems (including prolapse and fibroid)						
Polycystic ovary syndrome						
Prostatic hypertrophy (large prostate glands)						
Endometriosis						
Infertility						
Sexual dysfunction						
Other (please specify either one or more conditions):						

Round two public panel

Q26: Note inserted: In chronic infections domain, none of the individual conditions was rated by more than 70% of panellists as 'always include'. On the other hand, in the professional panel, there was consensus that HIV/AIDS should always be included in multimorbidity measurement.

Please rate the following chronic infections conditions as to whether they should be included or excluded. If you are not sure, please tick 'Don't know'.

Condition	Exclude (not important)	Expect to usually exclude (unless a good reason to include in a particular context)	Could include or exclude	Expect to usually include (unless a good reason to exclude in particular context)	Always include (extremely important)
HIV/AIDS					
Tuberculosis					
Lyme disease (a bacterial infection that can be spread to humans by infected ticks)					
Other (please specify either one or more conditions):					

Q27: Note inserted: In skin domain, none of the individual conditions was rated by more than 70% of panellists as 'always include', in both members of the public panel and professional panel.

Please rate the following skin conditions as to whether they should be included or excluded again. If you are not sure, please tick 'Don't know'.

Condition	Exclude (not important)	Usually exclude (unless a good reason to include in a particular context)	Could include or exclude	Usually include (unless a good reason to exclude in particular context)	Always include (extremely important)	Don't know
Psoriasis						
Eczema						
Chronic urticarial (chronic hives)						
Other (please specify either one or more conditions):						

Round two public panel

Q28: Note inserted: In ENT, eye and oral domains, none of the individual conditions was rated by more than 70% of panellists as 'always include', in both members of the public panel and professional panel.

Please rate the following ENT, eye and oral conditions as to whether they should be included or excluded again. If you are not sure, please tick 'Don't know'.

Condition	Exclude (not important)	Usually exclude (unless a good reason to include in a particular context)	Could include or exclude	Usually include (unless a good reason to exclude in particular context)	Always include (extremely important)	Don't know
Hearing impairment or Deafness (that cannot be easily corrected with hearing aids)						
Meniere's disease (an ear condition that causes sudden attacks of vertigo)						
Chronic sinusitis (sinus infection)						
Vision impairment or Blindness (that cannot be easily corrected with glasses)						
Cataract						
Glaucoma						
Edentulism (having no teeth)						
Chronic gum disease						
Other (please specify either one or more conditions):						

Q29: Note inserted: In congenital disease domain, none of the individual conditions was rated by more than 70% of panellists as 'always include', in both members of the public panel and professional panel.

Please rate the following statement as to whether congenital conditions should be included or excluded again. If you are not sure, please tick 'Don't know'.

Condition	Exclude (not	Usually	Could	Usually	Always	Don't
	important)	exclude	include or	include	include	know
		(unless a	exclude	(unless a		

Round two public panel

	good reason to include in a particular context)	good reason to exclude in particular context)	(extremely important)	
Congenital disease (conditions that babies are born with, including congenial heart disease, chromosomal aberrations)				
Other (please specify either one or more conditions):				

Q30: Note inserted: The following risk factors/symptoms have been included in multimorbidity measures by existing studies, but none of them was rated by more than 70% of panellists as 'always include'. Some suggested that those should be measured separately rather than as part of a multimorbdiity measure.

Having read a summary of the results from round one, please rate the following statement as to whether these conditions should be included or excluded again. If you are not sure, please tick 'Don't know'.

Condition	Exclude (not important)	Usually exclude (unless a good reason to include in a particular context)	Could include or exclude	Usually include (unless a good reason to exclude in particular context)	Always include (extremely important)	Don't know
Obesity (body mass index ≥30)						
Smoking						
Sedentary lifestyle (spending most of the day sitting)						
Physical disability						
Dizziness (without a specific diagnosis)						
Chronic cough (without a specific diagnosis)						
Post-sepsis syndrome						
Side effects of medications						
Treatment burden (the sum of all the hassles of taking medicines or attending appointments)						

Round two public

Social deprivation			
and poverty			
Other (please specify either one			
or more			
conditions):			

Weighting

Q31:

Reflecting on a summary of the results from round one and your responses so far, please rate the degree to which these outcomes, you think, are important to be considered when investigating the impact of multimorbidity.

Outcome	Not at all	Slightly	Important	Sufficiently	Very	No
	important	important		important	important	opinion
Death						
Healthcare use						
(e.g. number of						
emergency						
admissions to						
hospital;						
outpatient						
appointments)						
Quality of life						
Physical disability						
Frailty (general						
physical and/or						
mental weakness						
and vulnerability)						
Treatment burden						
(the sum of all the						
hassles of taking						
medicines or						
attending						
appointments)						
Healthcare costs						
(how much						
treatment and care						
for each individual						
costs)						
How people						
perceive their						
general health overall (e.g.						
whether it is						
excellent, very						
good, fair, poor,						
very poor)						
Other (please						
specify)						
Other (please						
specify)						
	1					

Round three public panel

Round-three survey in the public panel

Socio-demographic information
Q1: Please describe the country where you are currently living.
Q2: What is your age?
Q2. What is your age:
□ 18-34
□ 35-54
□ 55-64
$\square \geq 65$
☐ Prefer not to answer
Q3: What gender do you identify as?
☐ Female
☐ Male
Other
☐ Prefer not to answer
Q4: We are seeking your views as a member of the public, but we know that some of you will have worked in healthcare or academia. Do you currently or did you previously work in academia, health and social care practice or healthcare policy?
□ Yes
□ No
Q5: Do you have multiple chronic conditions?
□ Yes
□ No
Q6: Do you have family or friends who have multiple chronic conditions?
□ Yes
□ No
What is multimorbidity?
Q7: Please rate the degree to which you agree or disagree with the following statement again so we can see if there is a change between round 2 and round 3.

Neither

disagree

nor agree

Agree

Strongly

disagree

Disagree

Strongly

agree

Don't

know

Round th	ree pu	blic pa	nel
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Do you agree that defining complex multimorbidity in addition to a core definition of simple multimorbidity is useful?		П	П			
Q8: Irrespective of wheth would like to invite you to (please choose one).						
☐ 3+ conditions ☐ 3+ conditions ☐ 3+ conditions ☐ 4+ conditions ☐ No opinion ☐ Other	from 2+ bo from 3+ b	ody systems ody systems	S			
Note: In the public and pr patterns of conditions sho statements based on pane	uld be incl	luded in the				
Sub-question: Please choo the definition of "complex" No other patter Any combinate mental health condition Any combinate limitation Difficulty in refere.g. poverty). Any combinate Other	x multimorerns that I vition of two ditions ion of two managing i	rbidity". would like to or more cor or more cor llnesses due	o include nditions whit nditions with to social factors	ich includes b a significant pa	oth physical	and

Q9. Recurrence or remission

Please rate the degree to which you agree or disagree with the following statements.

Statement	Strongly disagree	Disagree	Neither disagree nor agree	Agree	Strongly agree	Don't know

Include remitting-relapsing conditions that require ongoing			

conditions that require ongoing treatment/ therapy/ care (e.g. depression, epilepsy)—newly added			
Include remitting-relapsing conditions which have relapsed during the last five years			

Q10: Treatment, care or surveillance

Round three public panel

Note inserted: More than 70% of panellists strongly agreed to include conditions that require current treatment, care, or therapy, and this question will not be asked again in round three. Please rate the degree to which you agree or disagree with the following statements.

Statement	Strongly disagree	Disagree	Neither disagree nor agree	Agree	Strongly agree	Don't know
Include conditions which usually require treatment, care or therapy 'at some point in the future' even if not currently treated (one panellist suggested to conceptualise this as 'current risk to future health outcomes')						
Include conditions which usually require surveillance (one panellist suggested to conceptualise this as 'current health needs or complexity of providing care')						

Q11: Categorisation of conditions/counting methods

Please choose one option that you would prefer to use.

☐ Broad disease category based on body and mind system (e.g. cardiovascular disease, mental health problems, skin conditions).
☐ Individual conditions (e.g. TIA (mini stroke) and stroke are counted separately).
☐ Grouping together similar conditions that are in the same category and treated similarly (e.g. grouping together 'Angina and Myocardial Infarction')
☐ Other. Please explain:

Round	three	public	panel
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□ Don't know

Q12: Data source

Please rate the following statements again.

Statement	Strongly disagree	Disagree	Neither disagree nor agree	Agree	Strongly agree	Don't know
Conditions included for multimorbidity measurement should be the same/similar in both self-report and databases (e.g. identifying a core set of conditions or condition framework).						
Other comments						

What conditions should be included?

Note inserted: In the previous section, more than 70% of panellists have agreed to include formal medical diagnoses, clinical risk factors, conditions that are permanent in their effects, conditions that last 12 months or longer, and require current treatment, care, or therapy.

Q13: Note inserted: In cancer conditions, we added three new statements based on panellists' comments. Please rate the following cancer conditions as to whether they should be included or excluded. If you are not sure, please tick 'Don't know'.

Condition	Exclude (not important)	Usually exclude (unless a good reason to include in a particular context)	Could include or exclude	Usually include (unless a good reason to exclude in particular context)	Always include (extremely important)	Don't know
Treated cancer that requires surveillance						
Treated cancer that did not recur over the past 5 years and does not require surveillance						
Benign cerebral tumours (brain tumours that can cause functional limitations)						

Round three public panel	
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Q14: Please choose th	e categorisation n	nethod you w	ould prefer	again.				
☐ Count ind affect (e.g. cou ☐ Count ind	Please rate the fo	ancers separa and liver cancancers separa er). n and consequence and conseque	ver separatel tely if they a uences of in itions as to v	y). Iffect differe jury were rev whether they	nt systems	(e.g.		
Condition	Exclude (not important)	Usually exclude (unless a	Could include or exclude	Usually include (unless a	Always include (extremel	Don't know		
		good reason to include in a particular context)	CACILLIC	good reason to exclude in particular context)	y important)			
Chronic or recurrent low back pain								
Chronic primary pain (defined as pain with no clear underlying condition but significant impact on the person)								
Long-term musculoskeletal problems due to injury (e.g. consequences of accidental injuries)								
Q16: Conditions newly added in round two Please rate again the following conditions so we can see if your responses have changed between round two and round three								
Condition	Exclude (not important)	Usually exclude (unless a good reason to include in a particular context)	Could include or exclude	Usually include (unless a good reason to exclude in particular context)	Always include (extremel y important)	Don't know		
Addison's disease (adrenal insufficiency, which is an uncommon								

Round three public panel

disorder which is fatal			
without lifelong			
treatment)			
Bronchiectasis (a lung			
condition that causes			
cough, sputum			
production, and			
recurrent respiratory			
infections)			
Post-acute covid 19			
("long COVID")			
Chronic Lyme disease			
(a bacterial infection			
that can be spread to			
humans by infected			
ticks)			

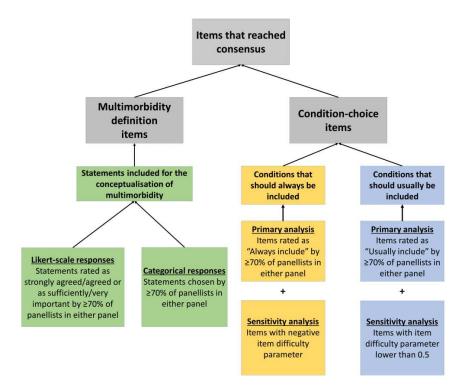
Measuring multimorbidity in research: a Delphi consensus study

Appendix 3

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Figure S1: Criteria for defining consensus of items



Box 1: Rasch Model used for sensitivity analysis

The Rasch dichotomous model is an item response theory model with one item parameter [1]. We used Rasch modelling to examine the degree to which condition-relevant items were endorsed by panelists. The common feature in the Rasch model is conditional maximum likelihood estimation which leads to the separability of item and person parameters [2]. Item discrimination (the level of endorsement varies with person ability) in the Rasch analysis is fixed for all items [3].

The mathematical form of the Rasch dichotomous model is given as follows[1, 3]:

 $P(X_{is} = 1 | \theta s, \beta i) = e(\theta s - \beta i)/1 + e(\theta s - \beta i)$

 X_{is} refers to response (X) made by subject s to item i.

 $X_{is} = 1$ refers to an endorsement of the item

 βi refers to the difficulty of item i

 θ refers to person's trait/ability

To estimate the trait level, we can take the natural log of a ratio of proportion endorsed responses to proportion unendorsed responses:

 $\theta = LN(Ps/1 - Ps)$

To estimate item difficulty, we can take the natural log of a ratio of proportion unendorsed responses to proportion endorsed responses.

 $\beta = LN(1 - Ps/Ps)$

Table S1: Characteristics of non-respondents and unidentified panelists‡

Table S1: Characteristics of Characteristics	Professional round	Professional round	Public round one	Public round two
Characteristics	one participant	two participant	participant who	participant who
	who did not	who did not	did not respond or	did not respond or
	respond or could	respond in round	could not be	could not be
	not be tracked in	three Professional	tracked in round	tracked in round
	round two (n=42)	round two (n=15)b	two (n=10) ^c	three (n=7)d
	No (%)	No (%)	No (%)	No (%)
	NO (%)	NO (%)	NO (76)	NO (%)
Continent				
Europe	24 (57.1%)	9 (60.0%)	7 (70.0%)	7 (100.0%)
North America	7 (16.7%)	2 (13.3%)	1 (10.0%)	0
Australasia	1 (2.4%)	2 (13.3%)	0	0
Asia	9 (21.4%)	1 (6.7%)	1 (10.0%)	0
South America	1 (2.4%)	0	0	0
Africa	0	1 (6.7%)	0	0
Not stated	0	0	1 (10.0%)	0
Country income			, ,	
High-income	39 (92.9%)	14 (93.3%)	8 (80.0%)	7 (100.0%)
Low and middle income	3 (7.1%)	1 (6.7%)	1 (10.0%)	0
Not stated	0	0	1 (10.0%)	0
Participant has multimorbidity			2 (20.070)	
Yes	2 (4.8%)	3 (20.0%)	13 (52.0%)	19 (61.3%)
No	40 (95.2%)	12 (80.0%)	12 (48.0%)	12 (38.7%)
Not stated	0	0	0	0
	0			
Family or friends have multimorbidity				
Yes	20 (66 70/)	10 (66 70/)	C (CO 00()	F (74 40/)
No	28 (66.7%)	10 (66.7%)	6 (60.0%)	5 (71.4%)
Type of work*	14 (33.3%)	5 (33.3%)	4 (40.0%)	2 (28.6%)
••	21 /72 00/\	12 (96 70/)		
Research	31 (73.8%)	13 (86.7%)	-	-
Public policy	4 (9.5%)	2 (13.3%)	-	-
Clinical practice	16 (38.1%)	4 (26.7%)	-	-
Teaching	2 (4.8%)	0	-	-
Main work setting	4 (0 50()			
Government	4 (9.5%)	0	-	-
Academia	26 (61.9%)	13 (86.7%)	-	-
Hospital	2 (4.8%)	0	-	-
Primary care	10 (23.8%)	2 (13.3%)	-	-
Other	0	0	-	-
Populations of interest*	27 (64 22()	44 (70 000)		
General population	27 (64.3%)	11 (73.3%)	-	-
Older people	29 (69.0%)	7 (46.7%)	-	-
Middle-aged and older	26 (61.9%)	6 (40.0%)	-	-
Socially-deprived population	23 (54.8%)	6 (40.0%)	-	-
Women	10 (23.8%)	2 (13.3%)	-	-
Men	10 (23.8%)	1 (6.7%)	-	-
Children	5 (11.9%)	1 (6.7%)	-	-
Ethnic minority/indigenous	1 (2.4%)	3 (20.0%)	-	-
People with disability	0	2 (13.3%)	-	-
Age group				
18-34	-	-	1 (10.0%)	0
35-54	-	-	2 (20.0%)	2 (28.6%)
55-64	-	-	3 (30.0%)	1 (14.3%)
≥65	-	-	4 (40.0%)	4 (57.1%)

Characteristics	Professional round one participant who did not respond or could not be tracked in round two (n=42) ^a No (%)	Professional round two participant who did not respond in round three Professional round two (n=15) ^b No (%)	Public round one participant who did not respond or could not be tracked in round two (n=10) ^c No (%)	Public round two participant who did not respond or could not be tracked in round three (n=7) ^d No (%)
Gender				
Female	-	-	4 (40.0%)	6 (85.7%)
Male	-	-	6 (60.0%)	1 (14.3%)

^{‡&}quot;Unidentified panelists" is defined as "panelists who did not provide their email or preferred name in the previous and following rounds". Some who dropped out in the following round did not provide their email and preferred name either. Thus, they were in the pool of those who either dropped out or cannot be identified". As we couldn't distinguish these two groups in the pool, we reported the characteristics of the two groups together.

*multiple select question

a. The 42 panelists consist of 38 who dropped out in round 2, and 4 are unidentified.

b. The 15 non-respondents in round 2

c. The 10 panelists consist of 6 who dropped out in round 2 and 4 are unidentified

d. The 7 panelists consist of 6 who dropped out in round 3 and 1 are unidentified

Table S2: Responses to questions on conceptualising multimorbidity

Question	Choice	Profession	al panel		Public pan	el	
		Round 1	Round 2	Round 3	Round 1	Round 2	Round 3
		(n=150)	(n=112)	(n=97)	(n=25)	(n=31)	(n=25)
Number of conditions to	2 or more	68.3%	84.8%	-	88.0%	-	-
define multimorbidity	3 or more	29.3%	12.5%		8.0%		
	4 or more	1.3%	0.9%		0		
	5 or more	0.7%	0		4.0%		
	Other	0.7%	1.8%		0		
How long-term a condition	3 months or more	16.7%	3.6%	-	8.0%	-	-
should be	6 months or more	42.7%	70.5%		8.0%		
	12 months or more	33.3%	23.2%		76.0%		
	Other	7.3%	2.7%		8.0%		
Do you agree that defining	Strongly agree	-	36.4%	60.4%	-	33.3%	36.0%
complex multimorbidity in	Agree		32.7%	27.1%		53.3%	48.0%
addition to a core definition of	Neither agree nor disagree		9.1%	4.2%		3.3%	4.0%
simple multimorbidity is	Disagree		15.5%	5.2%		6.7%	4.0%
useful?	Strongly disagree		6.4%	3.1%		3.3%	8.0%
How would you define	3 or more conditions irrespective of how many body	-	17.9%	25.8%	-	41.9%	76.0%
complex multimorbidity based	systems		29.5%	33.0%		16.1%	12.0%
on number of conditions?	3 or more conditions from two or more body systems		14.3%	26.8%		19.4%	12.0%
	3 or more conditions from three or more body systems		6.3%	3%		9.7%	0
	4 or more conditions irrespective of how many body		1.8%	0		0	0
	systems		3.6%	0		0	0
	4 or more conditions four or more body systems		1.8%	0		0	0
	5 or more conditions irrespective of how many body		25.0%	11.3%		12.9%	0
	systems						
	5 or more conditions five or more body systems						
	Other/No opinion						
Other patterns I would like to	No other patterns that I would like to include	-	29.5%	22.7%	-	16.1%	20.0%
include in the definition of	Any combination of two or more conditions which includes		36.6%	33.0%		64.5%	44.0%
complex multimorbidity*	both physical and mental health conditions						
	Any combination of two or more conditions with		0	30.9%		0	32.0%
	significantly physical functional limitation						
	Difficulty in managing illnesses due to social factors/social		0	26.8%		0	28.0%
	determinants of health						
	Any combination of two or more conditions and frailty		0	25.8%		0	12.0%
What kind of conditions to	Formal medical diagnosis	98.7%	99.1%	-	96.0%	96.8%	-
include*	Clinical risk factor	54.0%	49.1%		60.0%	74.2%	
	Symptom	22.7%	20.5%		28.0%	58.1%	

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	Health behaviour (e.g. smoking)	11.3%	8.9%		16.0%	51.6%	
	Health impact (e.g. disability)	39.3%	36.6%		40.0%	25.8%	
	Social deprivation and poverty	15.3%	10.7%		16.0%	38.7%	
	Consequences of treatment	22.7%	21.4%		40.0%	29.0%	
	Environmental factors	-	4.5%		-	22.6%	
How to count conditions	Count broad disease category based on body and mind	23.1%	-	-	12.0%	-	-
(round 1)	system	72.0%			88.0%		
	Count individual conditions	4.9%			0		
	Other						
How to count conditions	Count all of the conditions that are currently active	84.0%	-	-	80.0%	-	-
(round 2)	Count the primary health condition	4.0%			16.0%		
	Only count the complications	2.0%			0		
How to count cancers (round	Count all cancers as one	25.3%	-	-	24%	-	-
1)	Count individual cancers separately	62.0%			76%		
	Other	12.7%			0		
How to count cancers (round	Count all cancers as one	-	14.3%	-	-	16.1%	-
2)	Count individual cancers separately irrespective of which		38.4%			38.7%	
	systems they affect						
	Count individual cancers separately if they affect different		45.5%			45.2%	
	systems						
	Other		1.8%			0	
How to count cancers (round	Count all primary cancers as one	-	-	11.3%	-	-	12.0%
3)	Count individual primary cancers separately irrespective of			55.7%			48.0%
	which systems they affect						
	Count individual primary cancers separately only if they			33.0%			36.0%
	affect different systems			0			4.0%
	Other						
Conditions included for	Strongly agree	-	32.1%	29.1%	-	36.7%	44.0%
multimorbidity measurement	Agree		41.3%	42.7%		60.0%	52.0%
should be the same/similar in	Neither agree nor disagree		11.9%	5.5%		3.3%	4.0%
both self-report and databases	Disagree		11.9%	7.3%		0	0
	Strongly disagree		2.8%	1.8%		0	0

^{*} multiple select question

Supplemental material

Figure S2: Type of conditions to include when defining multimorbidity

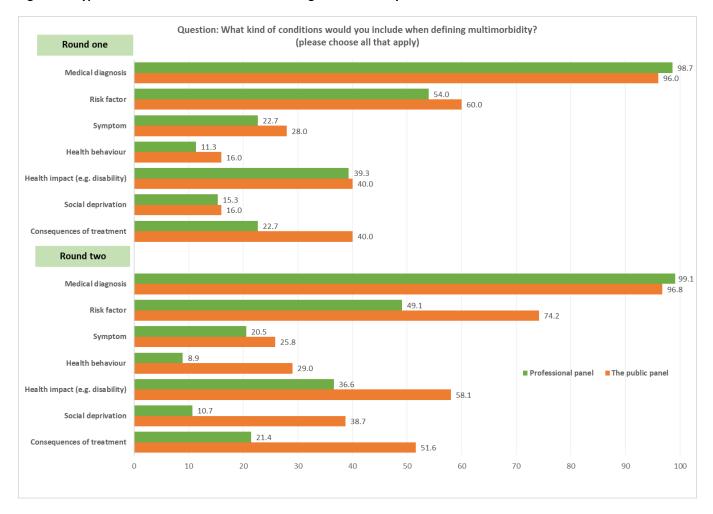


Table S3: Defining "condition" in the concept of multimorbidity (questions changed from round to round based on participant free-text feedback)

Statement	Rating	Pro	fessional pa	nel		Public pane	
		Round 1 (n=150)	Round 2 (n=112)	Round 3 (n=97)	Round 1 (n=25)	Round 2 (n=31)	Round 3 (n=25)
Include conditions	Strongly agree	89.9%			80.0%		
which are permanent in	Agree	8.7%			16.0%		
their effects	Neither agree/disagree	0			0		
	Disagree	0			0		
	Strongly disagree	1.3%			4.0%		
Include conditions	Strongly agree	73.2%			60.0%	64.5%	
which are currently	Agree	25.5%			28.0%	29.0%	
active	Neither agree/disagree	0			8.0%	3.2%	
	Disagree	0.7%			0	0	
	Strongly disagree	0.7%			4.0%	3.2%	
Include conditions	Strongly agree	13.6%			28.0%		
which may recur but	Agree	25.9%			16.0%		
happen rarely	Neither agree/disagree	19.7%			36.0%		
	Disagree	35.4%			20.0%		
	Strongly disagree	5.4%			0		
Include conditions	Strongly agree	3.170	27.3%			41.9%	
which may recur or	Agree		48.2%			45.2%	
remit	Neither agree/disagree		14.5%			9.7%	
Territ	Disagree		7.3%			0	
	Strongly disagree		2.7%			3.2%	
Include remitting-	Strongly agree		23.4%	12.9%		43.3%	21.0%
relapsing conditions			56.1%	48.4%		40.0%	37.5%
which have relapsed	Agree Neither agree/disagree		10.3%	22.6%		10.0%	37.5%
			10.3%	15.1%		3.3%	0
during the last five years	Disagree Strongly disagree		0	15.1%		3.3%	4.2%
Include remitting	Strongly agree		U	49.0%		3.3%	52.0%
Include remitting-							
relapsing conditions	Agree			44.8%			40.0%
that require ongoing	Neither agree/disagree			2.1%			0
treatment/ therapy/	Disagree			3.1%			0
care	Strongly disagree	42.40/		1.0%	20.00/		8.0%
Only include conditions	Strongly agree	13.4%			28.0%		
which usually require	Agree	31.5%			20.0%		
formal treatment or	Neither agree/disagree	14.1%			8.0%		
care	Disagree	32.9%			28.0%		
	Strongly disagree	8.1%			16.0%		
Include conditions	Strongly agree		77.5%			71.0%	
which usually require	Agree		22.5%			25.8%	
'current' treatment,	Neither agree/disagree		0			0	
care or therapy	Disagree		0			0	
	Strongly disagree		0			3.2%	
Include conditions	Strongly agree		36.7%	13.7%		27.6%	24.0%
which usually require	Agree		38.5%	43.2%		51.7%	44.0%
treatment, care or	Neither agree/disagree		11.9%	23.2%		10.3%	20.0%
therapy 'at some point	Disagree		12.8%	17.9%		3.4%	8.0%
in the future' even if not	Strongly disagree		0	2.1%		6.9%	4.0%
currently treated							
Include conditions	Strongly agree		38.0%	20.0%		41.9%	32.0%
which usually require	Agree		37.0%	54.7%		45.2%	56.0%
surveillance	Neither agree/disagree		11.1%	12.6%		9.7%	8.0%
	Disagree		13.0%	11.6%		0	0
	Strongly disagree		0.9%	1.1%	ĺ	3.2%	4.0%

Percentages were rounded.

Number of panelists voting "No opinion/Don't know" was not added in the denominator when calculating percentages

Table S4: Consideration of conditions to include based on impact (based on participant free-text comments, questions changed from round to round)

Statement	Rating	Professio	nal panel	Public panel		
		Round one (n=150)	Round two (n=112)	Round one (n=25)	Round two (n=31)	
Include conditions	Strongly agree	68.7%	69.4%	80.0%		
which significantly	Agree	20.4%	25.2%	20.0%		
increase risk of death	Neither agree nor disagree	5.4%	2.7%	0	-	
	Disagree	4.8%	0.9%	0		
	Strongly disagree	0.7%	1.8%	0		
	Strongly agree	76.5%		60.0%	67.7%	
Include conditions	Agree	20.1%		28.0%	25.8%	
which significantly	Neither agree nor disagree	2.7%	-	8.0%	3.2%	
reduce quality of life	Disagree	0.7%		4.0%	0	
	Strongly disagree	0		0	3.2%	
	Strongly agree	69.8%		68.0%	64.5%	
Include conditions	Agree	23.5%		24.0%	32.3%	
which cause significant	Neither agree nor disagree	5.4%	-	8.0%	0	
physical disability	Disagree	1.3%		0	0	
	Strongly disagree	0		0	3.2%	
	Strongly agree	65.8%	61.5%	44.0%	51.6%	
Include conditions	Agree	24.8%	28.4%	40.0%	38.7%	
Include conditions	Neither agree nor disagree	7.4%	5.5%	16.0%	6.5%	
which cause frailty	Disagree	2.0%	1.8%	0	0	
	Strongly disagree	0	2.8%	0	3.2%	
	Strongly agree	71.8%		62.5%	64.5%	
Include conditions	Agree	20.8%		16.7%	22.6%	
which significantly	Neither agree nor disagree	6.0%	-	20.8%	9.7%	
worsen mental health	Disagree	1.3%		0	0	
	Strongly disagree	0		0	3.2%	
Include conditions	Strongly agree	49.7%	48.6%	17.4%	16.7%	
which significantly	Agree	28.9%	28.8%	43.5%	46.7%	
worsen self-perceived	Neither agree nor disagree	17.4%	16.2%	26.1%	30.0%	
health status	Disagree	4.0%	4.5%	13.0%	0	
nealth status	Strongly disagree	0	1.8%	0	6.7%	
Include conditions	Strongly agree		55.0%		51.6%	
which significantly	Agree		32.4%		35.5%	
increase treatment	Neither agree nor disagree	-	7.2%	-	9.7%	
burden	Disagree		3.6%		0	
Durucii	Strongly disagree		1.8%		3.2%	
Include conditions	Strongly agree		25.5%		38.7%	
which could be	Agree		38.7%		35.5%	
impacted by or impact	Neither agree nor disagree	-	21.7%	-	16.1%	
social deprivation and	Disagree		11.3%		3.2%	
poverty	Strongly disagree		2.8%		6.5%	

Percentages were rounded.

Table S5: Responses to questions relevant to weighting (based on participant free-text comments, questions changed from round to round)

Question	Responses	Professional round 1 No. (%)	Professional round 2 No. (%)	Professional round 3 No. (%)	Public round 1 No. (%)	Public round 2 No. (%)	Public round 3 No. (%)
Please describe what type of	Weighted morbidity measure	91 (53.2%)	-	-	-	-	-
measures you would use to	A simple count of conditions	69 (40.4%)					
measure multimorbidity*	Other	11 (6.4%)					
Please choose what type of	Weighted morbidity measure	92 (61.3%)	-	-	-	-	-
measures you would use to	A simple count of conditions	38 (25.3%)					
understand the impact of	Both	7 (4.7%)					
multimorbidity on outcomes	Other	13 (8.7%)					
What weighting methods	Use existing weighted indices	47 (27.3%)	20 (21.7%)	-	-	-	-
would you use in a multimorbidity measure?a*	Empirically derive weights based on the individual impact of diseases on an outcome (e.g. use regression models to calculate weights)	60 (34.9%)	43 (46.7%)				
	Set rules based on level of severity to grade each condition/disease category (e.g. if having "presence of a condition": 1 point, "treatment": additional 1 point, "functional limitation": additional 1 point)	47 (27.3%)	24 (26.1%)				
	Other	18 (10.5%)	5 (5.4%)				
Please rate the degree to which the outcomes listed are important to weight against ^a							
Death	Not at all important	2 (1.4%)	-	-	0	0	-
	Slightly important	2 (1.4%)			2 (8.3%)	0	
	Important	6 (4.3%)			2 (8.3%)	1 (3.2%)	
	Sufficiently important	12 (8.6%)			4 (16.7%)	3 (9.7%)	
	Very important	117 (84.2%)			16 66.7%)	27 (87.1%)	
Healthcare use	Not at all important	1 (0.7%)	-	-	0	0	-
	Slightly important	5 (3.5%)			0	0	
	Important	17 (12.1%)			7 (29.2%)	3 (9.7%)	
	Sufficiently important	31 (22.0%)			4 (16.7%)	11 (35.5%)	
	Very important	87 (61.7%)			13 54.2%)	17 (54.8%)	
Health-related quality of life	Not at all important	1 (0.7%)	-	-	0	1 (3.2%)	-
, ,	Slightly important	3 (2.1%)			3 (12.5%)	0	
	Important	7 (4.9%)			3 (12.5%)	2 (6.5%)	
	Sufficiently important	17 (11.9%)			7 (29.2%)	8 (25.8%)	
	Very important	115 (80.4%)			11 45.8%)	20 (64.5%)	

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Question	Responses	Professional round 1 No. (%)	Professional round 2 No. (%)	Professional round 3 No. (%)	Public round 1 No. (%)	Public round 2 No. (%)	Public round 3 No. (%)
Physical disability	Not at all important	1 (0.7%)	-	-	0	0	-
	Slightly important	3 (2.2%)			1 (4.2%)	1 (3.3%)	
	Important	13 (9.4%)			10 41.7%)	3 (10.0%)	
	Sufficiently important	30 (21.6%)			6 (25.0%)	12 (40.0%)	
	Very important	92 (66.2%)			7 (29.2%)	14 (46.7%)	
Frailty	Not at all important	2 (1.4%)	-	-	0	0	-
	Slightly important	3 (2.2%)			5 (20.0%)	1 (3.3%)	
	Important	14 (10.1%)			10 40.0%)	6 (20.0%)	
	Sufficiently important	42 (30.2%)			4 (16.0%)	11 (36.7%)	
	Very important	78 (56.1%)			6 (24.0%)	12 (40.0%)	
Mental health	Not at all important	2 (1.4%)	-	-	0	_b	-
	Slightly important	6 (4.3%)			1 (4.0%)		
	Important	14 (10.1%)			8 (32.0%)		
	Sufficiently important	36 (26.1%)			5 (20.0%)		
	Very important	80 (58.0%)			11(44.0%)		
Treatment burden	Not at all important	3 (2.2%)	-	-	1 (4.2%)	1 (3.3%)	-
	Slightly important	9 (6.5%)			10 40.0%)	6 (20%)	
	Important	24 (17.3%)			2 (8.0%)	3 (10%)	
	Sufficiently important	32 (23.0%)			7 (28.0%)	12 (40.0%)	
	Very important	71 (51.1%)			5 (20.0%)	8 (26.7%)	
Healthcare costs	Not at all important	5 (3.6%)	-	-	3 (12.5%)	2 (6.7%)	-
	Slightly important	8 (5.8%)			6 (24.0%)	10 (33.3%)	
	Important	26 (18.7%)			4 (16.0%)	8 (26.7%)	
	Sufficiently important	46 (33.1%)			6 (24.0%)	5 (16.7%)	
	Very important	54 (38.8%)			6 (24.0%)	5 (16.7%)	
Self-perceived health status	Not at all important	3 (2.1%)	-	-	2 (8.0%)	2 (6.5%)	-
	Slightly important	11 (7.9%)			5 (20.0%)	6 (19.4%)	
	Important	22 (15.7%)			6 (24.0%)	7 (22.6%)	
	Sufficiently important	40 (28.6%)			6 (24.0%)	9 (29.0%)	
	Very important	64 (45.7%)			6 (24.0%)	7 (22.6%)	

Number of panelists voting "no opinion" was not added in the denominator when calculating percentages. *multiple select question; a: Non-mandatory question; b: we did not ask this question in round two because some panelists commented that conditions cannot be weighted against based on the general term "mental health"

Table S6: Professional ratings for choice of simple counts vs weighted measures depending on the purpose of a study

Question	Response	Round 2	Round 3	Round 3 summary
		No. of panelists (%)	No. of panelists (%)	
For the purpose of estimating the prevalence of multimorbidity	Prefer simple count Prefer weighted measure Either is acceptable	66 (59.5%) 18 (16.2%) 27 (24.3%)	58 (63.0%) 15 (16.3%) 19 (20.7%)	77 (83.7%) prefer a simple count or think it equally acceptable 34 (37.0%) prefer a weighted measure or think it equally acceptable
For the purpose of identifying and counting disease clusters	Prefer simple count Prefer weighted measure Either is acceptable	66 (68.8%) 15 (15.6%) 15 (15.6%)	59 (64.8%) 18 (19.8%) 14 (15.4%)	73 (80.2%) prefer a simple count or think it equally acceptable 32 (35.2%) prefer a weighted measure or think it equally acceptable
For the purpose of exploring trajectories of multimorbidity (e.g. trends of multimorbidity prevalence and the number of conditions an individual has had)	Prefer simple count Prefer weighted measure Either is acceptable	35 (32.4%) 27 (25.0%) 46 (42.6%)	34 (38.6%) 24 (27.3%) 30 (34.1%)	64 (72.7%) prefer a simple count or think it equally acceptable 54 (61.4%) prefer a weighted measure or think it equally acceptable
For the purpose of exploring/identifying predictors of multimorbidity (e.g. sociodemographic information)	Prefer simple count Prefer weighted measure Either is acceptable	30 (27.5%) 35 (32.1%) 44 (40.4%)	28 (30.8%) 32 (35.2%) 31 (34.1%)	59 (64.9%) prefer a simple count or think it equally acceptable 63 (69.3%) prefer a weighted measure or think it equally acceptable
For the purpose of assessing the severity of disease burden	Prefer simple count Prefer weighted measure Either is acceptable	12 (10.8%) 56 (50.5%) 43 (38.7%)	5 (5.4%) 66 (71.7%) 21 (22.8%)	26 (28.2%) prefer a simple count or think it equally acceptable 87 (94.5%) prefer a weighted measure or think it equally acceptable
For the purpose of risk adjustment or outcome prediction	Prefer simple count Prefer weighted measure Either is acceptable	8 (7.3%) 57(52.3%) 44 (40.4%)	8 (8.8%) 62 (68.1%) 21 (23.1%)	29 (31.9%) prefer a weighted measure or think it equally acceptable 83 (91.2%) prefer a weighted measure or think it equally acceptable

Table S7: Professional ratings for choice of simple counts vs weighted measures for outcome prediction depending on the outcome being measured

For the following outcomes of interest, please choose which type of multimorbidity measures you would prefer to use for outcome prediction ^a		Round 2 No. of panelists (%)	Round 3 No. of panelists (%)	Round 3 summary
Death	Prefer simple count Prefer weighted measure Either is acceptable	25 (24.5%) 41 (40.2%) 36 (35.3%)	10 (10.6%) 54 (57.4%) 30 (31.9%)	40 (42.5%) prefer a simple count or think it equally acceptable 94 (89.3%) prefer a weighted measure or think it equally acceptable
Healthcare use	Prefer simple count Prefer weighted measure Either is acceptable	28 (26.2%) 39 (36.4%) 40 (37.4%)	15 (16.1%) 41 (44.1%) 37 (39.8%)	52 (55.1%) prefer a simple count or think it equally acceptable 78 (83.9%) prefer a weighted measure or think it equally acceptable
Health-related quality of life	Prefer simple count Prefer weighted measure Either is acceptable	15 (14.2%) 49 (46.2%) 42 (39.6%)	5 (5.3%) 54 (57.4%) 35 (37.2%)	40 (42.5%) prefer a simple count or think it equally acceptable 89 (94.6%) prefer a weighted measure or think it equally acceptable
Physical disability	Prefer simple count Prefer weighted measure Either is acceptable	18 (16.8%) 49 (45.8%) 40 (37.4%)	8 (8.5%) 54 (57.4%) 32 (34.0%)	40 (42.5%) prefer a simple count or think it equally acceptable 86 (91.4%) prefer a weighted measure or think it equally acceptable
Frailty	Prefer simple count Prefer weighted measure Either is acceptable	17 (15.9%) 47 (43.9%) 43 (23.4%)	9 (9.6%) 52 (55.3%) 33 (35.1%)	42 (44.7%) prefer a simple count or think it equally acceptable 85 (90.4%) prefer a weighted measure or think it equally acceptable
Mental health	Prefer simple count Prefer weighted measure Either is acceptable	25 (40.2%) 43 (56.2%) 39 (36.4%)	14 (15.2%) 42 (45.7%) 36 (39.1%)	50 (54.3%) prefer a simple count or think it equally acceptable 78 (84.8%) prefer a weighted measure or think it equally acceptable
Treatment burden	Prefer simple count Prefer weighted measure Either is acceptable	19 (17.8%) 47 (43.9%) 41 (38.3%)	12 (12.8%) 53 (56.4%) 29 (30.9%)	41 (43.7%) prefer a simple count or think it equally acceptable 82 (87.3%) prefer a weighted measure or think it equally acceptable
Healthcare costs	Prefer simple count Prefer weighted measure Either is acceptable	20 (18.9%) 40 (37.7%) 46 (43.4%)	14 (15.2%) 41 (44.6%) 37 (40.2%)	51 (55.4%) prefer a simple count or think it equally acceptable 78 (84.8%) prefer a weighted measure or think it equally acceptable
Self-perceived health status	Prefer simple count Prefer weighted measure Either is acceptable	24 (23.1%) 41 (39.4%) 39 (37.5%)	15 (16.1%) 40 (43.0%) 38 (40.9%)	53 (57.0%) prefer a simple count or think it equally acceptable 78 (83.9%) prefer a weighted measure or think it equally acceptable

Table S8: Professional scores for 'always include'

Condition	Round 1	Round 2	Round 3	Professional
	N=150	N=112	N=97	panel
	%	%	%	consensus
Heart failure ^a	90.0	-	-	Always include
Chronic liver disease ^a	88.5	_	-	Always include
Diabetes ^a	87.3	-	-	Always include
Parkinson's disease ^a	86.6	-	-	Always include
End-stage kidney disease ^a	86.4	-	-	Always include
Chronic obstructive pulmonary disease ^a	85.9	-	-	Always include
Coronary artery disease ^a	82.7	-	-	Always include
Dementia ^a	82.6	-	-	Always include
Inflammatory bowel disease ^a	82.6	-	-	Always include
Multiple sclerosis ^a	80.7	-	-	Always include
Stroke ^a	80.0	-	-	Always include
Connective tissue disease ^a	79.7	-	-	Always include
Chronic kidney disease ^a	79.3	-	-	Always include
HIV/AIDS ^a	78.5	-	-	Always include
Metastatic cancers ^a	77.4	-	-	Always include
Haematological cancers ^a	77.2	-	-	Always include
Solid organ cancers ^a	76.5	-	-	Always include
Paralysis ^a	76.0	-	-	Always include
Cystic fibrosis ^a	75.8	-	-	Always include
Schizophrenia ^a	75.2	-	-	Always include
Epilepsy ^a	73.0	-	-	Always include
Peripheral artery disease ^a	71.1	-	-	Always include
Asthma ^a	70.7	-	-	Always include
Depression	69.6	67.0	-	
Heart valve disorders	57.7	53.6	-	
Bipolar disorder	61.9	61.8	-	
Melanoma	65.8	54.5	-	
Addison's disease ^b	-	48.1	34.5	
Bronchiectasis ^b	-	36.9	31.1	
Osteoarthritis	59.7	51.4	-	
Pancreatic disease	66.0	38.5	-	
Arrhythmia	58.4	46.4	-	
Thyroid problems Venous thrombotic disease	59.7	40.0	-	
	45.9	43.5	-	
Drug/alcohol misuse Anaemia	51.4 43.9	51.8 44.0	-	
Chronic Lyme disease ^b	45.9	29.8	25.3	
Transient ischaemic attack	49.3	48.2	25.5	
Hypertension	56.7	43.8	_	
Anxiety	52.0	47.3	_	
Treated cancer that requires surveillance ^c	-		35.9	
Eating disorders	45.0	44.5	-	
Vision impairment or blindness that	13.0	1 1.3		
cannot be corrected with glasses	52.7	46.4	_	
Musculoskeletal problems due to injury	44.5	41.1	36.1	
Tuberculosis	61.5	46.8	-	
Gout	45.3	38.5	-	
Endometriosis	42.4	35.5	-	
Chronic primary pain ^d	48.3	-	33.0	
Peptic ulcer	48.6	36.9	-	
Hearing impairment or deafness that				
cannot be corrected with hearing aids	48.0	39.6	-	
Post-acute COVID-19 ^b	-	27.7	24.5	
Post-traumatic stress disorder	38.0	42.2		

Benign cerebral tumours* Peripheral neuropathy \$2.4 36.1 - Peripheral neuropathy \$5.4 36.1 - Sleep apnoea \$4.5 29.9 - Congenital disease and chromosomal abnormalities \$4.2 35.8 - Chronic low back pain* Chronic low back pain* Chronic urinary tract infection 38.6 30.0 Aneurysm* Chronic urinary tract infection 38.6 30.0 Aneurysm* Aneurysm* Aneurysm* See and Chronic low back pain* Chronic urinary tract infection 38.6 30.0 Aneurysm* Aneurysm* Aneurysm* Solate and Sac 20.0 Migraine or other regular headache 37.8 28.7 - Osteoporosis Osteoporosis Osteoporosis Autism 36.2 29.0 - Obsessive compulsive disorder 38.6 32.7 - Dissociative or personality disorders 38.8 31.5 - Sidaucoma 44.2 36.0 - Urinary incontinence 38.8 31.5 - Sirviable bowel syndrome 38.8 29.4 - Disyociative ory syndrome 38.8 29.4 - Disyociative ory syndrome 38.8 29.4 - Disyolidaemial (treated) 35.0 30.8 - Obestity 48.3 35.1 - Obestity 48.3 35.1 - Oyslipidaemial (treated) 39.6 33.6 - Utreine (womb) problems 31.0 21.6 - Cataract 31.8 26.1 - Gall bladder problems 35.9 26.4 - Prostatic hypertrophy 29.5 27.2 - Attention deflict hyperactivity disorder 30.7 25.5 - Gastroesophageal reflux 32.0 27.3 - Chronic uriticaria 32.1 29.1 19.2 - Mainutrition 36.2 28.3 - High cholesterol (untreated) 44.9 25.5 - Learning disability 44.9 19.9 - Ferenan Social deprivation and poverty 14.4 19.9 - Post-sepsis syndrome 16.1 14.5 - Districtions 16.1 14.5 - Districtions 17.4 19.9 - Freated and one of the problems 18.7 - Freated and one of the problems 18.8 19.8 - Nicotine dependence 20.0 22.2 - Sexual dysfunction 14.0 16.2 - Sexual dysfunction 14.0 16.9 - Sexual dysfunction 14.0 16.9 - Sexual dysfunction 14.0 16.9 - Sexual dysfunction	Psoriasis	43.2	31.2		
Peripheral neuropathy		43.2	51.2	- 28 a	
Hypertension (untreated) 54.7 39.6 29.9 3.5 29.9 3.5 29.9 3.5 3.	_	52.4	36.1		
Siepe apnoea 45.5 29.9 -				_	
Congenital disease and chromosomal abnormalities				-	
abnormalities		45.5	23.3	-	
Chronic low back paind - - 30.9	_	46.3	25.0		
Chronic urinary tract infection				-	
Aneurym"	· ·				
Meniere's disease 34.9 26.9 - Migraine or other regular headache 37.8 28.7 - Osteoporosis 52.0 41.4 - Autism 36.2 29.0 - Obsessive compulsive disorders 41.0 40.7 - Glaucoma 44.2 36.0 - Urinary incontinence 38.8 31.5 - Irritable bowel syndrome 38.8 29.4 - Polycystic ovary syndrome 36.6 29.8 - Chronic insomnia 35.0 30.8 - Obesity 48.3 35.1 - Dyslipidaemia (treated) 39.6 33.6 - Uterine (womb) problems 31.0 21.6 - Cataract 31.8 26.1 - Galt bladder problems 35.9 26.4 - Prostatic hypertrophy 29.5 27.2 - Attention deficit hyperactivity disorder 30.7 25.5 -					
Migraine or other regular headache 37.8 28.7 - Osteoporosis 52.0 41.4 - Autism 36.2 29.0 - Obsessive compulsive disorder 38.6 32.7 - Dissociative or personality disorders 41.0 40.7 - Glaucoma 44.2 36.0 - Urinary incontinence 38.8 31.5 - Irritable bowle syndrome 36.6 29.8 - Polycystic ovary syndrome 36.6 29.8 - Chronic insomnia 35.0 30.8 - Obesity 48.3 35.1 - Dyslipidaemia (treated) 39.6 33.6 - Uterine (womb) problems 31.0 21.6 - Cataract 31.8 26.1 - Gall bladder problems 35.9 26.4 - Prostatic hypertrophy 29.5 27.2 - Attention deficit hyperactivity disorder 30.7 25.5 -	1			22.7	
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Autism				-	
Obsessive compulsive disorder 38.6 32.7 - Dissociative or personality disorders 41.0 40.7 - Glaucoma 44.2 36.0 - Urinary incontinence 38.8 31.5 - Irritable bowel syndrome 36.6 29.8 - Chronic insomnia 35.0 30.8 - Obesity 48.3 35.1 - Dyslipidaemia (treated) 39.6 33.6 - Uterine (womb) problems 31.0 21.6 - Cataract 31.8 26.1 - Gall bladder problems 35.9 26.4 - Prostatic hypertrophy 29.5 27.2 - Attention deficit hyperactivity disorder 27.0 25.9 - Somatoform disorders 30.7 25.5 - Gastroesophageal reflux 30.0 27.3 - Chronic urticaria 29.1 19.2 - Malanutrition 36.2 28.3 -	I			-	
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Irritable bowel syndrome 38.8 29.4 -	Glaucoma	44.2	36.0	-	
Polycystic ovary syndrome	Urinary incontinence	38.8	31.5	-	
Chronic insomnia 35.0 30.8 -	Irritable bowel syndrome	38.8	29.4	-	
Obesity 48.3 35.1 - Dyslipidaemia (treated) 39.6 33.6 - Uterine (womb) problems 31.0 21.6 - Cataract 31.8 26.1 - Gall bladder problems 35.9 26.4 - Prostatic hypertrophy 29.5 27.2 - Attention deficit hyperactivity disorder 27.0 25.9 - Somatoform disorders 30.7 25.5 - Somatoform disorders 30.7 25.5 - Gastroesophageal reflux 32.0 27.3 - Chronic urticaria 29.1 19.2 - Malnutrition 36.2 28.3 - High cholesterol (untreated) 44.9 25.5 - Learning disability 28.1 25.2 - Kidney or bladder stones 28.6 22.4 - Physical disability 38.9 27.9 - Eczema 25.9 19.3 -	Polycystic ovary syndrome	36.6	29.8	-	
Dyslipidaemia (treated) 39.6 33.6 -	Chronic insomnia	35.0	30.8	-	
Uterine (womb) problems 31.0 21.6 - Cataract 31.8 26.1 - Gall bladder problems 35.9 26.4 - Prostatic hypertrophy 29.5 27.2 - Attention deficit hyperactivity disorder 27.0 25.9 - Somatoform disorders 30.7 25.5 - Gastroesophageal reflux 32.0 27.3 - Chronic urticaria 29.1 19.2 - Malnutrition 36.2 28.3 - High cholesterol (untreated) 44.9 25.5 - Learning disability 28.1 25.2 - Kidney or bladder stones 28.6 22.4 - Physical disability 38.9 27.9 - Eczema 25.9 19.3 - Non-melanoma skin cancer 30.3 23.1 - Chronic sinusitis 27.4 17.3 - Chronic gum disease 16.8 19.8 - Nicotine dependence 20.0 22.2 - <td< td=""><td>Obesity</td><td>48.3</td><td>35.1</td><td>-</td><td></td></td<>	Obesity	48.3	35.1	-	
Cataract 31.8 26.1 - Gall bladder problems 35.9 26.4 - Prostatic hypertrophy 29.5 27.2 - Attention deficit hyperactivity disorder 27.0 25.9 - Somatoform disorders 30.7 25.5 - Gastroesophageal reflux 32.0 27.3 - Chronic urticaria 29.1 19.2 - Malnutrition 36.2 28.3 - High cholesterol (untreated) 44.9 25.5 - Learning disability 28.1 25.2 - Kidney or bladder stones 28.6 22.4 - Physical disability 38.9 27.9 - Eczema 25.9 19.3 - Non-melanoma skin cancer 30.3 23.1 - Chronic sinusitis 27.4 17.3 - Chronic/allergic rhinitis 30.1 17.4 - Diverticulosis 38.7 22.2 - Chronic gum disease 16.8 19.8 - Ni	Dyslipidaemia (treated)	39.6	33.6	-	
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Edentulism 20.1 18.7 - Smoking 29.3 27.9 - Social deprivation and poverty 24.3 17.9 - Infertility 10.6 14.0 - Post-sepsis syndrome 14.3 15.7 - Treatment burden 16.1 14.5 - Dizziness 10.3 10.8 - Side effects of medications 12.0 11.9 - Chronic cough 8.3 10.7 - Benign cancers 18.6 9.3 - Sedentary lifestyle 14.4 9.9 -				-	
Smoking 29.3 27.9 - Social deprivation and poverty 24.3 17.9 - Infertility 10.6 14.0 - Post-sepsis syndrome 14.3 15.7 - Treatment burden 16.1 14.5 - Dizziness 10.3 10.8 - Side effects of medications 12.0 11.9 - Chronic cough 8.3 10.7 - Benign cancers 18.6 9.3 - Sedentary lifestyle 14.4 9.9 -				-	
Social deprivation and poverty 24.3 17.9 - Infertility 10.6 14.0 - Post-sepsis syndrome 14.3 15.7 - Treatment burden 16.1 14.5 - Dizziness 10.3 10.8 - Side effects of medications 12.0 11.9 - Chronic cough 8.3 10.7 - Benign cancers 18.6 9.3 - Sedentary lifestyle 14.4 9.9 -				-	
Infertility				-	
Post-sepsis syndrome 14.3 15.7 - Treatment burden 16.1 14.5 - Dizziness 10.3 10.8 - Side effects of medications 12.0 11.9 - Chronic cough 8.3 10.7 - Benign cancers 18.6 9.3 - Sedentary lifestyle 14.4 9.9 -				-	
Treatment burden 16.1 14.5 - Dizziness 10.3 10.8 - Side effects of medications 12.0 11.9 - Chronic cough 8.3 10.7 - Benign cancers 18.6 9.3 - Sedentary lifestyle 14.4 9.9 -	·			-	
Dizziness 10.3 10.8 - Side effects of medications 12.0 11.9 - Chronic cough 8.3 10.7 - Benign cancers 18.6 9.3 - Sedentary lifestyle 14.4 9.9 -				-	
Side effects of medications 12.0 11.9 - Chronic cough 8.3 10.7 - Benign cancers 18.6 9.3 - Sedentary lifestyle 14.4 9.9 -				-	
Chronic cough 8.3 10.7 - Benign cancers 18.6 9.3 - Sedentary lifestyle 14.4 9.9 -				-	
Benign cancers 18.6 9.3 - Sedentary lifestyle 14.4 9.9 -				-	
Sedentary lifestyle 14.4 9.9 -	_			-	
	=	18.6	9.3	-	
Treated cancer not requiring surveillance ^c - 5.4	Sedentary lifestyle	14.4	9.9	-	
a Reached (always include) consensus after round 1		-	-	5.4	

a. Reached 'always include' consensus after round 1

b. Proposed as additional conditions by participants after round $\ensuremath{\mathbf{1}}$

c. Proposed as additional conditions by participants after round 2

d. Chronic pain asked about in rounds 1, split into chronic primary pain and chronic low back pain by participants after round 2

Table S9: Public scores for 'always include'

Condition	Round 1	Round 2	Round 3	Public panel
	N=150	N=112	N=97	consensus
	%	%	%	
End-stage kidney disease	68.0	90.3	-	Always include
Heart failure	68.0	83.9	-	Always include
Dementia	60.0	83.3	-	Always include
Chronic liver disease	64.0	80.6	-	Always include
Stroke	60.0	80.6	-	Always include
Chronic kidney disease	64.0	80.6	-	Always include
Parkinson's disease	56.0	77.4	-	Always include
Multiple sclerosis	60.0	77.4	_	Always include
Coronary artery disease	64.0	74.2	_	Always include
Cystic fibrosis	62.5	74.2	_	Always include
Diabetes	64.0	71.0	_	Always include
HIV/AIDS	37.5	71.0	_	Always include
Epilepsy	44.0	71.0	_	Always include
Addison's disease	-	40.0	70.8	Always include
Metastatic cancers	70.8	80.0	-	Always include
Haematological cancers	70.8	77.4	_	Always include
Solid organ cancers	70.8	80.0		Always include
Inflammatory bowel disease	60.0	58.1		Aiways illulude
Heart valve disorders	36.0	61.3	_	
Melanoma	58.3	67.7	_	
	60.0	67.7	_	
Chronic obstructive pulmonary disease	24.0	35.5	-	
Peripheral artery disease Venous thrombotic disease	48.0		-	
		64.5	-	
Transient ischaemic attack	32.0	64.5	-	
Aneurysm	40.0	67.7	-	
Pancreatic disease	48.0	53.3	-	
Anaemia	24.0	43.3	-	
Peripheral neuropathy	44.0	26.7	-	
Schizophrenia	56.5	67.7	-	
Bipolar disorder	52.2	51.6	-	
Connective tissue disease	54.2	66.7	-	
Paralysis	56.0	63.3	-	
Post-acute COVID-19	-	44.8	60.0	
Tuberculosis	54.2	54.8	-	
Congenital disease	44.0	53.3	-	
Prostatic hypertrophy	25.0	23.3	-	
Endometriosis	36.4	50.0	-	
Bronchiectasis	-	45.2	48.0	
Osteoarthritis	41.7	41.9	-	
Thyroid problems	37.5	35.5	-	
Autism	22.7	35.5	-	
Chronic urinary tract infection	20.0	40.0	-	
Polycystic ovary syndrome	40.9	37.9	-	
Arrhythmia	28.0	39.3	-	
Peptic ulcer	29.2	35.5	-	
Learning disability	25.0	38.7	-	
Asthma	32.0	58.1	-	
Osteoporosis	33.3	45.2	-	
Uterine (womb) problems	22.7	35.5	-	
Treated cancer that requires surveillance	-	-	36.0	
Chronic primary pain	24.0	-	20.0	
Malnutrition	14.3	36.7	-	
Non-melanoma skin cancer	30.4	36.7	-	
Chronic Lyme disease	-	38.7	45.8	

Depression	37.5	25.8	-	
Irritable bowel syndrome	20.0	26.7	-	
Kidney or bladder stones	20.0	26.7	-	
Benign cerebral tumours	-		52.0	
Diverticulosis	25.0	24.1	-	
Drug/alcohol misuse	20.8	35.5	-	
Eating disorders	34.8	45.2	-	
Vision impairment or blindness that				
cannot be corrected with glasses	36.0	41.9	-	
Hearing impairment or deafness that				
cannot be corrected with hearing aids	25.0	38.7	-	
Post-traumatic stress disorder	30.4	25.8	-	
Dyslipidaemia (treated)	24.0	29.0	-	
Physical disability	29.2	41.9	-	
Hypertension (untreated)	29.2	35.5	-	
Meniere's disease	29.2	25.8	-	
Dissociative or personality disorders	42.9	41.9	-	
Musculoskeletal problems due to injury	25.0	16.1	45.8	
Glaucoma	39.1	33.3	-	
Gall bladder problems	29.2	36.7	-	
Sleep apnoea	37.5	19.4	-	
Obsessive compulsive disorder	14.3	22.6	-	
Urinary incontinence	20.8	22.6	-	
Chronic/allergic rhinitis	16.7	16.1	-	
Gout	25.0	19.4	-	
Psoriasis	16.7	16.1	-	
Obesity	20.8	16.1	-	
Attention deficit hyperactivity disorder	19.0	12.9	-	
Somatoform disorders	5.0	17.2	-	
Hypertension (treated)	29.2	16.7	-	
Anxiety	25.0	16.7	-	
Benign cancers	17.4	17.2	-	
Migraine or other regular headache	20.8	12.9	-	
Gastroesophageal reflux	24.0	12.9	-	
Eczema	20.8	6.5	-	
Post-sepsis syndrome	23.8	28.6	-	
Chronic low back pain	-	-	24.0	
Cataract	20.8	16.1	-	
Treatment burden	12.5	20.0	-	
Side effects of medications	16.7	23.3	-	
Social deprivation and poverty	17.4	23.3	-	
Chronic sinusitis	16.0	9.7	-	
Nicotine dependence	13.6	22.6	-	
High cholesterol (untreated)	29.2	12.9	-	
Smoking	17.4	16.1	-	
Chronic insomnia	18.2	13.3	-	
Chronic urticaria	13.0	6.9	-	
Chronic gum disease	21.7	9.7	-	
Chronic cough	12.5	3.2	-	
Sedentary lifestyle	0.0	9.7	-	
Sexual dysfunction	13.6	13.8	-	
Edentulism	8.3	6.5	-	
Dizziness	17.4	3.2	-	
Infertility	22.7	17.2	-	
Treated cancer not requiring surveillance	-	-	8.0	
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Table S10: Professional scores for 'usually include' (the sum of 'always' and 'usually')

Condition	Round 1	Round 2	Round 3	Professional
	N=150	N=112	N=97	panel consensus
	%	%	%	
Heart failure	98.7	-	-	Always include
Chronic liver disease	98.6	-	-	Always include
Diabetes	98.7	-	-	Always include
Parkinson's disease	98.0	-	-	Always include
End-stage kidney disease	94.6	-	-	Always include
Chronic obstructive pulmonary disease	98.7	-	-	Always include
Coronary artery disease	97.3	-	-	Always include
Dementia	96.6	-	-	Always include
Inflammatory bowel disease	94.0	-	-	Always include
Multiple sclerosis	94.0	-	-	Always include
Stroke	97.3	-	-	Always include
Connective tissue disease	97.3	-	-	Always include
Chronic kidney disease	97.3	-	-	Always include
HIV/AIDS	94.0	-	-	Always include
Metastatic cancers	91.1	-	-	Always include
Haematological cancers	94.0	-	-	Always include
Solid organ cancers	94.6	-	-	Always include
Paralysis	88.4	-	-	Always include
Cystic fibrosis	90.6	-	-	Always include
Schizophrenia	94.0	-	-	Always include
Epilepsy	95.3	-	-	Always include
Peripheral artery disease	92.6	-	-	Always include
Asthma	92.7	-	-	Always include
Depression	93.9	92.9	-	Usually include
Heart valve disorder	85.2	92.0	-	Usually include
Bipolar disorder	90.5	90.0	-	Usually include
Melanoma	87.7	88.2	-	Usually include
Addison's disease	-	83.7	86.9	Usually include
Bronchiectasis	-	78.6	86.7	Usually include
Osteoarthritis	87.9	84.7	-	Usually include
Pancreatic disease	88.9	84.4	-	Usually include
Arrhythmia	85.9	83.9	-	Usually include
Thyroid problems	83.9	82.7	-	Usually include
Venous thrombotic disease	77.4	82.4	-	Usually include
Drug/alcohol misuse	82.4	81.8	-	Usually include
Anaemia	75.0	81.7	-	Usually include
Chronic Lyme disease	-	61.5	81.3	Usually include
Transient ischaemic attack	76.7	80.4	-	Usually include
Hypertension (treated)	78.7	80.4	-	Usually include
Anxiety	86.5	80.0	-	Usually include
Treated cancer that requires surveillance	-	-	79.3	Usually include
Eating disorders	74.3	79.1	-	Usually include
Vision impairment or blindness that				
cannot be corrected with glasses	80.1	78.6	-	Usually include
Musculoskeletal problems due to injury	80.1	78.6	78.4	Usually include
Tuberculosis	82.4	76.6	-	Usually include
Gout	79.1	76.1	-	Usually include
Endometriosis	73.6	75.7	-	Usually include
Chronic primary pain	78.5	-	75.3	Usually include
Peptic ulcer	79.7	73.9	-	Usually include
Hearing impairment or deafness that				
cannot be corrected with hearing aids	70.9	73.9	-	Usually include
Post-acute COVID-19	-	70.3	73.4	Usually include
Post-traumatic stress disorder	77.5	73.4	-	Usually include

-				
Psoriasis	77.0	73.4	-	Usually include
Benign cerebral tumours	-	-	73.3	Usually include
Peripheral neuropathy	83.4	73.1	-	Usually include
Hypertension (untreated)	73.6	73.0	-	Usually include
Sleep apnoea	77.2	72.9	-	Usually include
Congenital disease	79.7	72.6	-	Usually include
Chronic low back pain	-	-	72.2	Usually include
Chronic urinary tract infection	73.8	71.8	-	Usually include
Aneurysm	-	-	71.6	Usually include
Meniere's disease	63.0	71.3	-	Usually include
Migraine or other regular headache	68.9	71.3	-	Usually include
Osteoporosis	77.3	70.3	-	Usually include
Autism	63.0	70.1	-	Usually include
Obsessive compulsive disorder	63.6	70.1	-	Usually include
Dissociative or personality disorders	69.8	69.4	-	,
Glaucoma	72.8	69.4	-	
Urinary incontinence	67.3	66.7	-	
Irritable bowel syndrome	63.9	66.1	_	
Polycystic ovary syndrome	66.9	65.4	-	
Chronic insomnia	69.3	64.5	_	
Obesity	69.1	64.0	_	
Dyslipidaemia (treated)	63.1	63.6	_	
Uterine (womb) problems	71.0	63.1	_	
Cataract	62.2	63.1	_	
Gall bladder problems	67.6	62.7	_	
Prostatic hypertrophy	55.0	62.1	_	
Attention deficit hyperactivity disorder	56.9	60.2	-	
Somatoform disorders	55.7	59.4	_	
Gastroesophageal reflux	60.5	59.1	_	
Chronic urticaria	59.6	58.7	-	
Malnutrition	64.5	57.5	-	
High cholesterol (untreated)	63.9	57.3	-	
Learning disability	53.3	57.3 55.1	-	
Kidney or bladder stones	55.5 57.1	55.1 55.1	-	
	60.4	55.0	-	
Physical disability Eczema	60.4		-	
Non-melanoma skin cancer	54.5	54.1	-	
		53.7	-	
Chronic sinusitis	54.1	53.6	-	
Chronic/allergic rhinitis Diverticulosis	58.2	53.2	-	
	69.7	50.0	-	
Chronic gum disease	46.2	49.1	-	
Nicotine dependence	42.9	48.1	-	
Sexual dysfunction	42.0	46.7	-	
Edentulism	38.8	41.1	-	
Smoking	45.6	37.8	-	
Social deprivation and poverty	41.0	36.6	-	
Infertility	33.3	33.6	-	
Post-sepsis syndrome	42.1	32.4	-	
Treatment burden	27.3	31.8	-	
Dizziness	33.6	30.6	-	
Side effects of medications	28.9	29.4	-	
Chronic cough	29.0	27.7	-	
Benign cancers	35.9	26.2	-	
Sedentary lifestyle	28.8	18.0	-	
Treated cancer not requiring surveillance	-	-	21.5	

Table S11: Public scores for 'usually include' (the sum of 'always' and 'usually')

Condition	Round 1	Round 2	Round 3	Public panel
	N=25	N=31	N=25	consensus
	%	%	%	
End-stage kidney disease	96.0	100.0	-	Always include
Heart failure	100.0	96.8	-	Always include
Dementia	92.0	96.7	-	Always include
Chronic liver disease	100.0	100.0	-	Always include
Stroke	92.0	96.8	-	Always include
Chronic kidney disease	100.0	100.0	-	Always include
Parkinson's disease	92.0	96.8	-	Always include
Multiple sclerosis	100.0	100.0	-	Always include
Coronary artery disease	96.0	96.8	-	Always include
Cystic fibrosis	100.0	96.8	-	Always include
Diabetes	92.0	100.0	-	Always include
HIV/AIDS	91.7	90.3	-	Always include
Epilepsy	92.0	100.0	-	Always include
Addison's disease	-	88.0	95.8	Always include
Metastatic cancers	91.7	96.7	-	Always include
Haematological cancers	91.7	100.0	-	Always include
Solid organ cancers	91.7	100.0	-	Always include
Inflammatory bowel disease	100.0	100.0	-	Usually include
Heart valve disorders	76.0	100.0	-	Usually include
Melanoma	91.7	100.0	-	Usually include
Chronic obstructive pulmonary disease	100.0	96.8	-	Usually include
Peripheral artery disease	88.0	96.8	-	Usually include
Venous thrombotic disease	80.0	96.8	-	Usually include
Transient ischaemic attack	88.0	96.8	-	Usually include
Aneurysm	-	96.8	-	Usually include
Pancreatic disease	96.0	96.7	-	Usually include
Anaemia	60.0	96.7	-	Usually include
Peripheral neuropathy	76.0	96.7	-	Usually include
Schizophrenia	91.3	93.5	-	Usually include
Bipolar disorder	95.7	93.5	-	Usually include
Connective tissue disease	83.3	93.3	-	Usually include
Paralysis	84.0	93.3	-	Usually include
Post-acute COVID-19	-	93.1	92.0	Usually include
Tuberculosis	91.7	90.3	-	Usually include
Congenital disease	88.0	90.0	-	Usually include
Prostatic hypertrophy	58.3	90.0	-	Usually include
Endometriosis	72.7	89.3	-	Usually include
Bronchiectasis	-	87.1	88.0	Usually include
Osteoarthritis	79.2	87.1	-	Usually include
Thyroid problems	75.0	87.1	-	Usually include
Autism	63.6	87.1	-	Usually include
Chronic urinary tract infection	60.0	86.7	-	Usually include
Polycystic ovary syndrome	72.7	86.2	-	Usually include
Arrhythmia	68.0	85.7	-	Usually include
Peptic ulcer	66.7	83.9	-	Usually include
Learning disability	50.0	83.9	-	Usually include
Asthma	96.0	80.6	-	Usually include
Osteoporosis	70.8	80.6	-	Usually include
Uterine (womb) problems	63.6	80.6	-	Usually include
Treated cancer that requires surveillance	-	-	80.0	Usually include
Chronic primary pain	76.0	-	80.0	Usually include
Malnutrition	57.1	80.0	-	Usually include
Non-melanoma skin cancer	69.6	80.0	-	Usually include
Chronic Lyme disease	-	83.9	79.2	Usually include

Depression	87.5	77.4	-	Usually include
Irritable bowel syndrome	68.0	76.7	-	Usually include
Kidney or bladder stones	52.0	76.7	-	Usually include
Benign cerebral tumours	58.3	-	76.0	Usually include
Diverticulosis	62.5	75.9	-	Usually include
Drug/alcohol misuse	78.3	74.2	-	Usually include
Eating disorders	72.0	74.2	-	Usually include
Vision impairment or blindness that				
cannot be corrected with glasses	58.3	74.2	-	Usually include
Hearing impairment or deafness that				
cannot be corrected with hearing aids	65.2	74.2	-	Usually include
Post-traumatic stress disorder	64.0	74.2	-	Usually include
Dyslipidaemia (treated)	70.8	74.2	-	Usually include
Physical disability	79.2	74.2	-	Usually include
Hypertension (untreated)	75.0	71.0	-	Usually include
Meniere's disease	81.0	71.0	-	Usually include
Dissociative or personality disorders	70.8	71.0	-	Usually include
Musculoskeletal problems due to injury	69.6	67.7	70.8	Usually include
Glaucoma	58.3	70.0	-	Usually include
Gall bladder problems	62.5	70.0	-	Usually include
Sleep apnoea	47.6	67.7	-	
Obsessive compulsive disorder	41.7	67.7	-	
Urinary incontinence	50.0	67.7	-	
Chronic/allergic rhinitis	41.7	67.7	-	
Gout	54.2	64.5	-	
Psoriasis	58.3	64.5	-	
Obesity	47.6	61.3	-	
Attention deficit hyperactivity disorder	45.0	61.3	-	
Somatoform disorders	83.3	58.6	-	
Hypertension (treated)	62.5	56.7	-	
Anxiety	52.2	56.7	-	
Benign cancers	58.3	55.2	-	
Migraine or other regular headache	56.0	54.8	-	
Gastroesophageal reflux	45.8	54.8	-	
Eczema	66.7	54.8	-	
Post-sepsis syndrome	-	53.6	-	
Chronic low back pain	41.7	-	52.0	
Cataract	29.2	51.6	-	
Treatment burden	41.7	50.0	-	
Side effects of medications	39.1	50.0	-	
Social deprivation and poverty	44.0	46.7	-	
Chronic sinusitis	40.9	45.2	-	
Nicotine dependence	66.7	45.2	-	
High cholesterol (untreated)	34.8	41.9	-	
Smoking	54.5	41.9	-	
Chronic insomnia	43.5	40.0	-	
Chronic urticaria	34.8	37.9	-	
Chronic gum disease	54.2	35.5	-	
Chronic cough	13.0	35.5	-	
Sedentary lifestyle	45.5	35.5	-	
Sexual dysfunction	25.0	34.5	-	
Edentulism	47.8	32.3	-	
Dizziness	45.5	32.3	-	
Infertility	-	27.6	-	
Treated cancer not requiring surveillance		-	24.0	

Table S12: Conditions not recommended to 'always include' or 'usually include' by both panels

Condition	Professional	Public	Both panels	Final consensus
	panel	panel (%)	(difficulty	
	(%)	` ` `	parameter,	
	` `		logit)	
Consensus to usually include in one panel (n=22))			
Hypertension (treated)	80.4 (R2)	56.7 (R2)	0.3 (R2)	Usually include
Anxiety	80.0 (R2)	56.7 (R2)	0.4 (R2)	Usually include
Gout	76.1 (R2)	64.5 (R2)	0.4 (R2)	Usually include
Psoriasis	73.4 (R2)	64.5 (R2)	0.6 (R2)	No consensus
Sleep apnoea	72.9 (R2)	67.7 (R2)	0.6 (R2)	No consensus
Obsessive compulsive disorder	70.1 (R2)	67.7 (R2)	0.7 (R2)	No consensus
Prostatic hypertrophy	62.1 (R2)	90.0 (R2)	0.7 (R2)	No consensus
Polycystic ovary syndrome	65.4 (R2)	86.2 (R2)	0.7 (R2)	No consensus
Dissociative or personality disorders	69.4 (R2)	71.0 (R2)	0.7 (R2)	No consensus
Glaucoma	69.4 (R2)	70.0 (R2)	0.7 (R2)	No consensus
Chronic low back pain	72.2 (R3)	52.0 (R3)	0.8 (R3)	No consensus
Migraine	71.3 (R2)	54.8 (R2)	0.8 (R2)	No consensus
Uterine (womb) problems	63.1 (R2)	80.0 (R2)	0.8 (R2)	No consensus
Irritable bowel syndrome	66.1 (R2)	76.7 (R2)	0.8 (R2)	No consensus
Gall bladder problems	62.7 (R2)	70.0 (R2)	1.0 (R2)	No consensus
Malnutrition	57.5 (R2)	80.0 (R2)	1.1 (R2)	No consensus
Learning disability	55.1 (R2)	83.9 (R2)	1.2 (R2)	No consensus
Kidney or bladder stones	55.1 (R2)	76.7 (R2)	1.2 (R2)	No consensus
High cholesterol (untreated)	57.3 (R2)	74.2 (R2)	1.2 (R2)	No consensus
Non-melanoma skin cancer	53.7 (R2)	80.6 (R2)	1.3 (R2)	No consensus
Physical disability	55.0 (R2)	74.2 (R2)	1.3 (R2)	No consensus
Diverticulosis	50.0 (R2)	75.9 (R2)	1.5 (R2)	No consensus
No consensus in either panel (n=27)				
Urinary incontinence	66.7 (R2)	67.7 (R2)	0.9 (R2)	No consensus
Obesity	64.0 (R2)	61.3 (R2)	1.0 (R2)	No consensus
Cataract	63.1 (R2)	51.6 (R2)	1.2 (R2)	No consensus
Attention deficit hyperactivity disorder	60.2 (R2)	61.3 (R2)	1.2 (R2)	No consensus
Gastroesophageal reflux	59.1 (R2)	54.8 (R2)	1.3 (R2)	No consensus
Chronic insomnia	64.5 (R2)	40.0 (R2)	1.3 (R2)	No consensus
Dyslipidaemia (treated)	63.6 (R2)	41.9 (R2)	1.3 (R2)	No consensus
Somatoform disorders	59.4 (R2)	58.6 (R2)	1.3 (R2)	No consensus
Chronic/Allergic rhinitis	53.2 (R2)	67.7 (R2)	1.4 (R2)	No consensus
Eczema	54.1 (R2)	54.8 (R2)	1.5 (R2)	No consensus
Chronic urticarial (chronic hives)	58.7 (R2)	37.9 (R2)	1.6 (R2)	No consensus
Chronic sinusitis	53.6 (R2)	45.2 (R2)	1.7 (R2)	No consensus
Chronic gum disease	49.1 (R2)	35.5 (R2)	1.9 (R2)	No consensus
Nicotine dependence	48.1 (R2)	45.2 (R2)	2.0 (R2)	No consensus
Sexual dysfunction	46.7 (R2)	34.5 (R2)	2.1 (R2)	No consensus
Edentulism	41.1 (R2)	32.3 (R2)	2.3 (R2)	No consensus
Smoking	37.8 (R2)	41.9 (R2)	2.4 (R2)	No consensus
Social deprivation and poverty	36.6 (R2)	46.7 (R2)	2.4 (R2)	No consensus
Post-sepsis syndrome	32.4 (R2)	53.6 (R2)	2.5 (R2)	No consensus
Treatment burden	31.8 (R2)	50.0 (R2)	2.5 (R2)	No consensus
Benign cancers	26.2 (R2)	55.2 (R2)	2.7 (R2)	No consensus
Infertility	33.6 (R2)	27.6 (R2)	2.7 (R2)	No consensus
Side effects of medications	29.4 (R2)	50.0 (R2)	2.8 (R2)	No consensus
Chronic cough	27.7 (R2)	35.5 (R2)	2.9 (R2)	No consensus
Dizziness	30.6 (R2)	32.3 (R2)	2.9 (R2)	No consensus
Treated cancer not requiring surveillance	21.5 (R3)	24.0 (R3)	3.4 (R3)	No consensus
Sedentary lifestyle	18.0 (R2)	35.5 (R2)	3.4 (R2)	No consensus

^{*} Consensus to 'usually include' in one panel and difficulty parameter across both panels ≤0.5

Table S13: Differential item functioning (used to examine which condition-choice items were more likely to be endorsed by panelists with varying interest in different populations [4])

Subgroup	Condition	Group 1 (b	Group 0 (b	Z-statistic	p-value
		parameter, logit)	parameter, logit)		
Panelists specifically	Anaemia	-0.6	-0.2	-0.7	0.51
interested in	Depression	-1.0	-1.8	0.9	0.35
multimorbidity in women	Anxiety	0.04	-0.1	0.3	0.78
(group1) versus those not	Dementia	-1.9	-2.8	0.9	0.35
specifically interested	Eating disorder	-0.3	0.06	-0.5	0.62
(group 0)	Osteoarthritis	-1.5	-0.3	-1.5	0.14
	Osteoporosis	0.04	0.7	-1.2	0.25
	Connective tissue disease	-2.6	-2.8	0.2	0.86
	Endometriosis	0.08	0.3	-0.3	0.78
	Chronic urinary tract infection	0.3	0.5	-0.4	0.71
	Polycystic ovary syndrome	0.9	0.9	-0.004	1.0
	Urinary incontinence	0.3	1.0	-1.2	0.25
	Uterus (womb) problems	1.0	1.0	0.004	1.0
Panelists specifically	Anxiety	-0.8	0.01	-0.9	0.37
interested in	Depression	-0.8	-1.7	1.0	0.32
multimorbidity in	Drug/Alcohol misuse	-0.8	-0.2	-0.7	0.50
children (group1) versus	Eating disorder	-0.2	0.02	-0.2	0.83
those not specifically	Autism	-0.8	0.8	-1.8	0.07
interested (group 0)	Learning disability	1.4	1.5	-0.08	0.94
	Attention deficit hyperactivity disorder	-0.2	1.4	-2.0	0.04
	Haematological cancers	-1.8	-1.9	0.01	1.0
	Benign brain tumour	2.9	3.2	-0.4	0.65
	Epilepsy	-1.0	-2.4	1.5	0.12
	Cystic fibrosis	-1.8	-1.3	-0.5	0.61
	Congenital diseases	-0.08	0.5	-0.7	0.46
	Asthma	-1.8	-1.6	-0.2	0.84
Panelists specifically	Anxiety	-0.7	0.4	-2.1	0.04
interested in	Depression	-1.5	-2.0	0.6	0.56
multimorbity in socially-	Schizophrenia	-1.5	-2.6	1.4	0.15
deprived populations	Drug/Alcohol misuse	-0.4	-0.2	-0.4	0.71
(group1) versus those not	Nicotine dependence	1.6	1.9	-0.6	0.52
specifically interested	Post-traumatic stress disorder	0.3	0.4	-0.2	0.85
(group 0)	Bipolar disorder	-1.2	-1.1	-0.1	0.91
	Chronic liver disease	-3.4	-3.0	0.4	0.69
	HIV/AIDS	-2.0	-2.1	0.2	0.88

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Subgroup	Condition	Group 1 (b parameter, logit)	Group 0 (b parameter, logit)	Z-statistic	p-value
	Tuberculosis	0.07	0.01	0.1	0.91

p-value larger than 0.05 indicates that there is no statistical difference between group 1 and group 0. A positive value for the z-statistic indicates that group 0 is more likely to endorse the condition than group 1.

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Table S14: Level of endorsement of the condition-related items

Condition	Difficulty parameter (log(odds), 95% Confidence Interval)			
	Professional panel	Public panel	Both panels combined	
System domain				
Cardiovascular diseases	-4.2 (-6.1 to -2.2)	-1.0 (-2.2 to 0.2)	-2.8 (-3.8 to -1.8)	
Respiratory diseases	-3.8 (-5.8 to -1.8)	endorsed by all panelists	-3.9 (-5.9 to -2.0)	
Neurological diseases	-3.4 (-4.9 to -2.0)	-1.5 (-2.9 to -0.03)	-2.8 (-3.8 to -1.8)	
Metabolic and endocrine	-2.7 (-3.7 to -1.7)	-1.0 (-2.2 to 0.2)	-2.2 (-3.0 to -1.4)	
Mental health	-2.2 (-3.1 to -1.4)	-0.2 (-1.1 to 0.8)	-1.6 (-2.2 to -0.9)	
Haematological disorders	-2.0 (-3.0 to -1.1)	-1.0 (-2.2 to 0.2)	-1.7 (-2.5 to -1.0)	
Cancers	-1.9 (-2.7 to 1.2)	-1.5 (-2.9 to -0.03)	-1.8 (-2.4 to -1.1)	
Musculoskeletal diseases	-1.6 (-2.3 to 1.0)	-0.7 (-1.8 to 0.4)	-1.4 (-2.0 to -0.8)	
Digestive diseases	-1.1 (-1.8 to -0.4)	-1.0 (-2.2 to 0.2)	-1.0 (-1.6 to -0.4)	
Chronic infections	-1.2 (-1.9 to -0.5)	0.3 (-0.6 to 1.1)*	-0.7 (-1.2 to -0.2)	
Urogenital disorders	-0.8 (-1.4 to -0.1)	-0.7 (-1.8 to 0.4)	-0.7 (-1.2 to -0.2)	
Eye diseases	0.05 (-0.5 to 0.6)	0.3 (-0.6 to 1.1)*	0.1 (-0.3 to 0.6)*	
Ear, Nose and Throat	0.3 (-0.2 to 0.8)*	-0.2 (-1.1 to 0.8)	0.2 (-0.2 to 0.7)*	
disorders				
Oral disorders	0.4 (-0.05 to 0.9)*	0.1 (-0.8 to 1.0)*	0.4 (-0.02 to 0.8)*	
Skin disorders	0.5 (0.02 to 1.0)*	0.1 (-0.8 to 1.0)*	0.4 (-0.02 to 0.9)*	
Mental health and				
behaviour disorders				
Dementia	-2.4 (-3.4 to -1.5)	-2.2 (-4.2 to -0.2)	-2.3 (-3.2 to -1.5)	
Schizophrenia	-1.7 (-2.4 to -1.0)	-1.5 (-2.9 to -0.03)	-1.6 (-2.2 to -1.0)	
Depression	-1.5 (-2.3 to -0.7)	0.1 (-0.8 to 1.0)	-0.9 (-1.5 to -0.4)	
Bipolar disorder	-1.0 (-1.7 to -0.3)	-1.5 (-2.9 to -0.03)	-1.1 (-1.7 to -0.5)	
Anxiety	-0.02 (-0.6 to 0.5)	1.2 (0.4 to 1.9)	0.4 (-0.08 to 0.8)*	
Drug/Alcohol misuse	-0.2 (-0.7 to 0.4)	0.3 (-0.6 to 1.1)*	-0.02 (-0.5 to 0.4)	
Eating disorder	0.1 (-0.5 to 0.6)*	0.3 (-0.6 to 1.1)*	0.2 (-0.3 to 0.6)*	
Post-traumatic stress	0.5 (-0.002 to 1.0)*	0.3 (-0.6 to 1.1)*	0.5 (0.05 to 0.9)*	
disorder				
Dissociative or personality disorder	0.7 (0.3 to 1.2)	0.4 (-0.4 to 1.3)*	0.7 (0.3 to 1.1)	
Chronic insomnia	1.0 (0.6 to 1.5)	1.9 (1.1 to 2.8)	1.3 (0.9 to 1.7)	
Autism	0.7 (0.2 to 1.2)	-0.7 (-1.8 to 0.4)	0.5 (0.03 to 0.9)*	
Obsessive compulsive disorder	0.5 (0.2 to 1.2)*	0.6 (0.2 to 1.4)	0.7 (0.3 to 1.1)	
Attention deficit hyperactivity disorder	1.3 (0.8 to 1.7)	1.4 (0.7 to 2.2)	1.2 (0.8 to 1.6)	
Somatoform disorders	1.3 (0.9 to 1.8)	1.0 (0.2 to 1.7)	1.3 (0.9 to 1.7)	
Learning disability	1.5 (1.1 to 2.0)	-0.4 (-1.4 to 0.6)	1.2 (0.8 to 1.5)	
Nicotine dependence	1.9 (1.5 to 2.3)	1.8 (1.0 to 2.5)	1.9 (1.5 to 2.3)	
Cardiovascular diseases	,	, ,		
Heart failure	-3.4 (-4.9 to -2.0)	-2.2 (-4.2 to -0.2)	-3.1 (-4.3 to -2.0)	
Coronary heart diseases	-2.7 (-3.7 to -1.7)	-2.2 (-4.2 to -0.2)	-2.6 (-3.5 to -1.6)	
Stroke	-2.7 (-3.7 to -1.7)	-2.2 (-4.2 to -0.2)	-2.6 (-3.5 to -1.6)	
Peripheral artery diseases	-1.5 (-2.2 to -0.9)	-2.2 (-4.2 to -0.2)	-1.6 (-2.2 to -0.9)	
Arrhythmia	-0.4 (-1.0 to 0.2)	-0.6 (1.7 to 0.5)	-0.4 (-0.9 to 0.1)	
Heart valve disorders	-0.3 (-0.8 to 0.2)	endorsed by all panelists	-1.6 (-2.3 to -0.9)	
Venous thrombotic disease	-0.3 (-0.8 to -0.3)	-2.2 (-4.2 to -0.2)	-0.5 (-1.0 to -0.003)	
Hypertension (treated)	-0.1 (-0.6 to 0.4)	1.2 (0.4 to 2.0)	0.3 (-0.1 to 0.7)*	
rispertension (treated)	5.1 (5.5 to 6.4)	1.2 (0.7 to 2.0)	3.3 (0.1 to 0.7)	

Hypertension (untreated)	0.4 (-0.05 to 0.9)*	0.4 (-0.4 to 1.3)*	0.4 (-0.1 to 0.9)*
Aneurysm	0.6 (0.05 to 1.1)	-2.2 (-4.2 to -0.2)	0.5 (0.1 to 0.9)*
Dyslipidaemia (treated)	1.0 (0.6 to 1.5)	1.9 (1.1 to 2.7)	1.3 (0.9 to 1.7)
High cholesterol	1.4 (0.9 to 1.8)	0.4 (-0.4 to 1.3)*	1.2 (0.8 to 1.5)
(untreated)		,	, ,
Disorders of the blood			
Anaemia	-0.2 (-0.7 to 0.4)	-2.2 (-4.2 to -0.2)	-0.4 (-0.9 to 0.08)
Cancers			
Solid organ cancers	-1.9 (-2.6 to -1.1)	endorsed by all panelists	-2.0 (-2.7 to -1.3)
Haematological cancers	-1.7 (-2.5 to -1.0)	endorsed by all panelists	-1.9 (-2.6 to -1.2)
Metastatic cancers	-1.3 (-1.9 to -0.6)	-2.2 (-4.2 to -0.2)	-1.3 (-1.9 to -0.7)
Melanoma	-0.8 (-1.5 to -0.2)	endorsed by all panelists	-1.1 (-1.7 to -0.5)
Treated cancers requiring	-0.04 (-0.6 to 0.5)	0.03 (-1.0 to 1.1)*	0.01 (-0.5 to 0.5)*
surveillance	,	,	
Benign cerebral tumours	0.3 (-0.2 to 0.9)*	0.3 (-0.7 to 1.3)*	0.4 (-0.1 to 0.8)*
Non-melanoma skin	1.6 (1.2 to 2.0)	-0.1 (-1.0 to 0.9)	1.3 (0.9 to 1.7)
cancers			
Benign cancers	3.2 (2.7 to 3.7)	1.3 (0.5 to 2.1)	2.7 (2.3 to 3.1)
Treated cancer not	3.4 (2.9 to 4.0)	3.1 (2.1 to 4.1)	3.4 (2.9 to 3.9)
requiring surveillance			
Neurological diseases			
Parkinson's disease	-3.0 (-4.2 to -1.8)	-2.2 (-4.2 to -0.2)	-2.8 (-3.8 to -1.8)
Epilepsy	-2.1 (-2.9 to -1.3)	endorsed by all panelists	-2.2 (-3.0 to -1.4)
Multiple sclerosis	-1.8 (-2.5 to -1.1)	endorsed by all panelists	-1.9 (-2.6 to -1.2)
Paralysis	-0.9 (-1.5 to -0.3)	-1.4 (-2.9 to 0.01)	-0.9 (-1.4 to -0.4)
Transient ischaemic attack	-0.1 (-0.6 to 0.4)	-2.2 (-4.2 to -0.2)	-0.4 (-0.8 to 0.1)
Chronic primary pain	0.2 (-0.3 to 0.8)*	0.003 (-1.0 to 1.1)*	0.2 (-0.2 to 0.7)*
Peripheral neuropathy	0.5 (-0.02 to 1.0)*	-2.2 (-4.2 to -0.2)	0.1 (-0.3 to 0.6)*
Sleep apnoea	0.5 (-0.3 to 1.0)*	0.6 (-0.2 to 1.4)	0.6 (0.2 to 1.0)
Migraine	0.5 (0.1 to 1.1)*	1.3 (0.5 to 2.1)	0.8 (0.4 to 1.2)
Metabolic and endocrine			
disorders			
Diabetes	-3.4 (-4.9 to -2.0)	endorsed by all panelists	-3.5 (-4.9 to -2.1)
Cystic fibrosis	-1.2 (-1.8 to -0.6)	-2.2 (-4.2 to -0.2)	-1.3 (-1.9 to -0.7)
Thyroid disorders	-0.3 (-0.9 to 0.3)	-0.7 (-1.8 to 0.4)	-0.3 (-0.8 to 0.2)
Addison's disease	-0.6 (-1.3 to 0.1)	-1.7 (-3.8 to 0.3)	-0.8 (-1.4 to 0.1)
Malnutrition	1.4 (0.9 to 1.8)	-0.1 (-1.0 to 0.9)	1.1 (0.7 to 1.5)
Musculoskeletal diseases			
Connective tissue disease	-3.4 (-4.8 to -2.0)	-1.4 (-2.9 to 0.01)	-2.3 (-3.1 to -1.5)
Osteoarthritis	-0.5 (-1.0 to 0.1)	-0.7 (-1.8 to 0.4)	-0.5 (-1.0 to 0.04)
Long-term musculoskeletal	0.01 (-0.5 to 0.6)*	0.6 (-0.3 to 1.6)	0.2 (-0.3 to 0.7)*
impairment due to injuries			
Gout	0.2 (-0.3 to 0.7)*	0.8 (-0.009 to 1.6)	0.4 (0.01 to 0.9)*
Chronic low back pain	0.5 (-0.1 to 1.0)*	1.6 (0.7 to 2.4)	0.8 (0.3 to 1.2)
Osteoporosis	0.6 (0.2 to 1.1)	-0.2 (-1.1 to 0.8)	0.5 (0.08 to 0.9)*
Respiratory diseases			
Chronic obstructive	-3.4 (-4.8 to -2.0)	-2.2 (-4.2 to -0.2)	-3.1 (-4.2 to -1.9)
pulmonary disease			
Asthma	-1.5 (-2.2 to -0.9)	0.2 (-1.1 to 0.8)	-1.1 (-1.7 to -0.6)
Bronchiectasis	-0.7 (-1.3 to 0.001)	-1.1 (-2.6 to 0.4)	-0.7 (-1.3 to -0.1)
Post-acute COVID-19	0.4 (-0.2 to 0.9)*	-0.6 (-1.9 to 0.6)	0.2 (-0.3 to 0.7)*
Chronic/Allergic rhinitis	1.6 (1.2 to 2.0)	0.6 (-0.2 to 1.4)	1.4 (1.0 to 1.8)

Infectious diseases			
HIV/AIDS	-1.7 (-2.5 to -1.0)	-1.0 (-2.2 to 0.2)	-1.5 (-2.2 to -0.9)
Chronic Lyme disease	-0.1 (0.7 to 0.5)	0.2 (-0.9 to 1.2)*	-0.03 (-0.5 to 0.5)
Tuberculosis	0.2 (-0.3 to 0.7)*	-1.0 (-2.2 to 0.2)	0.07 (-0.4 to 0.5)*
Digestive diseases			·
Chronic liver diseases	-3.4 (-4.8 to -2.0)	endorsed by all panelists	-3.5 (-4.9 to -2.1)
Inflammatory bowel	-1.8 (-2.5 to -1.0)	endorsed by all panelists	-1.9 (-2.6 to -1.2)
disease			
Pancreatic disease	-0.4 (-1.0 to 0.1)	-2.2 (-4.2 to -0.2)	-0.7 (-1.2 to -0.1)
Peptic ulcer	0.4 (-1.0 to 0.9)*	-0.4 (-1.4 to 0.6)	0.3 (-0.2 to 0.7)*
Irritable bowel syndrome	0.9 (0.4 to 1.4)	0.1 (-0.8 to 1.0)*	0.8 (0.4 to 1.2)
Gall bladder disorders	1.1 (0.6 to 1.5)	0.5 (-0.3 to 1.4)*	1.0 (0.6 to 1.4)
Gastroesophageal reflux	1.3 (0.9 to 1.7)	1.3 (0.5 to 2.1)	1.3 (0.9 to 1.7)
Diverticulosis	1.8 (1.4 to 2.3)	0.1 (-0.8 to 1.0)*	1.5 (1.1 to 1.9)
Urogenital diseases			
Chronic kidney disease	-2.7 (-3.7 to -1.7)	endorsed by all panelists	-2.8 (-3.8 to -1.8)
End-stage kidney disease	-1.9 (-2.6 to -1.1)	endorsed by all panelists	-2.0 (-2.7 to -1.2)
Endometriosis	0.3 (-0.2 to 0.8)*	-0.7 (-2.0 to 0.5)	0.1 (-0.3 to 0.6)*
Chronic urinary tract	0.5 (-0.05 to 1.0)*	-0.6 (-1.7 to 0.4)	0.3 (-0.1 to 0.8)*
infection			
Polycystic ovary syndrome	0.9 (0.5 to 1.4)	-0.6 (-1.7 to 0.5)	0.7 (0.2 to 1.1)
Urinary incontinence	0.9 (0.4 to 1.3)	0.6 (-0.2 to 1.4)	0.9 (0.5 to 1.3)
Uterine (womb) problems	1.1 (0.6 to 1.5)	-0.2 (-1.1 to 0.8)	0.8 (0.4 to 1.2)
Prostatic hypertrophy	1.1 (0.7 to 1.6)	-1.0 (-2.2 to 0.2)	0.7 (0.3 to 1.2)
Kidney or bladder stones	1.5 (1.1 to 2.0)	0.1 (-0.8 to 1.0)*	1.2 (0.8 to 1.6)
Sexual dysfunction	2.0 (1.5 to 2.4)	2.3 (1.5 to 3.2)	2.1 (1.7 to 2.5)
Infertility	2.7 (2.2 to 3.2)	2.8 (1.9 to 3.7)	2.7 (2.3 to 3.1)
Skin disorders			
Psoriasis	0.4 (-0.05 to 0.9)*	0.8 (-0.009 to 1.6)	0.6 (0.2 to 1.0)
Eczema	1.6 (1.1 to 2.0)	1.3 (0.5 to 2.1)	1.5 (1.2 to 1.9)
Chronic urticarial	1.4 (0.9 to 1.8)	2.1 (1.3 to 2.9)	1.6 (1.2 to 2.0)
Eye diseases			
Vision impairment that	0.05 (-0.5 to -0.6)*	0.3 (-0.6 to 1.1)*	0.1 (-0.3 to 0.6)*
cannot be corrected with			
glasses			<u> </u>
Glaucoma	0.7 (0.2 to 1.1)	0.5 (-0.3 to 1.3)*	0.7 (0.3 to 1.1)
Cataract	1.0 (0.6 to 1.5)	1.4 (0.7 to 2.2)	1.2 (0.8 to 1.6)
ENT disorders			
Hearing impairment that	0.4 (-0.1 to -0.9)*	0.3 (-0.6 to 1.1)*	0.4 (-0.02 to 0.8)*
cannot be corrected with			
hearing aids	0.6 (0.1 to 1.1)	0.4 (0.4 +0.1.2)*	0 5 (0 2 to 1 0)*
Meniere's disease Chronic sinusitis	0.6 (0.1 to 1.1)	0.4 (-0.4 to 1.3)* 1.8 (1.0 to 2.5)	0.5 (0.2 to 1.0)* 1.7 (1.3 to 2.0)
Oral disorders	1.6 (1.1 to 2.0)	1.0 (1.0 (0 2.3)	1.7 (1.3 (0 2.0)
	1 2 (0 0 to 1 7)	2 2 (1 4 +0 2 1)	2 0 /1 6 to 2 /\
Chronic gum disease Edentulism	1.3 (0.9 to 1.7) 1.9 (1.5 to 2.3)	2.3 (1.4 to 3.1) 2.4 (1.6 to 3.3)	2.0 (1.6 to 2.4) 2.3 (1.9 to 2.7)
	0.5 (-0.003 to 1.0)*		0.2 (-0.2 to 0.7)*
Congenital and chromosomal abnormalities	0.5 (-0.003 to 1.0)*	-1.0 (-2.2 to 0.2)	0.2 (-0.2 (0 0.7)*
Risk factors/Symptoms/	<u> </u>		
Behaviour			
Obesity	1.0 (0.5 to 1.4)	1.0 (0.2 to 1.7)	1.0 (0.6 to 1.4)
		· · · · · · · · · · · · · · · · · · ·	, ,

Smoking	2.5 (2.0 to 2.9)	1.9 (1.1 to 2.7)	2.4 (2.0 to 2.8)
Social deprivation or	2.5 (2.1 to 3.0)	1.7 (0.9 to 2.5)	2.4 (2.0 to 2.8)
poverty			
Post-sepsis syndrome	2.8 (2.3 to 3.3)	1.4 (0.5 to 2.2)	2.5 (2.1 to 2.9)
Treatment burden	2.8 (2.4 to 3.3)	1.5 (0.4 to 2.3)	2.5 (2.2 to 2.9)
Dizziness	2.9 (2.4 to 3.4)	2.4 (1.6 to 3.3)	2.9 (2.4 to 3.4)
Side effects of medicines	3.0 (2.5 to 3.4)	1.5 (0.4 to 2.3)	2.8 (2.4 to 3.2)
Chronic cough	3.1 (2.6 to 3.6)	2.3 (1.4 to 3.1)	2.9 (2.5 to 3.3)
Sedentary lifestyle	3.8 (3.3 to 4.4)	2.3 (1.4 to 3.1)	3.4 (3.0 to 3.9)

Lower difficulty parameter estimates indicate more frequent endorsement of items. Here we classified the numbers to three levels— ≤0 (in bold) indicates strongly endorsed, 0-0.5 indicates endorsed (*), and >0.5 indicates infrequently endorsed [5, 6]

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