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Policies to prevent the spillover of zoonotic diseases: protocol for a systematic scoping review of evaluative evidence

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1 Abstract

2 Introduction

The increasing incidence of disease transmission from vertebrate animals to humans (zoonotic spillover events) has been attributed to ecological, behavioural and socioeconomic change. As these events sometimes involve diseases with epidemic or pandemic potential, they pose a serious threat to population health. Public policies may play a key role in preventing these events. The aim of this review is to identify evaluations of public policies that target the determinants of zoonotic spillover, examining approaches taken to evaluation, choice of outcomes measures and evidence of effectiveness. Our approach to identifying and analysing this literature will be informed by a population health perspective and a One Health lens, acknowledging the inter-connectedness of human, animal and environmental health.

12 Methods and analysis

A systematic scoping review methodology will be used. To identify articles, we will search Medline, SCOPUS, Web of Science and Global Health using search terms combining public policy, prevention, zoonoses and spillover events. We will screen titles and abstracts and extract data according to published guidelines for scoping reviews. All evaluations of public policies aiming to prevent zoonotic spillover events will be eligible for inclusion. We will summarise key data from each study, mapping policies along the disease transmission pathway and outlining the range of policies, approaches to evaluation and outcome measures. Review findings will provide a useful reference for researchers and practitioners, outlining the state of the evaluative evidence around policies to prevent zoonotic spillover.

³⁹ 22 Ethics and dissemination40

Formal ethical approval is not required, as the study does not involve primary data collection.
The findings of this study will be disseminated through a peer-reviewed publication,
presentations, and summaries for key stakeholders.

26 Strengths and limitations

- This scoping review protocol outlines the first piece of work to systematically identify
 and review evaluations of public policies designed to prevent the spillover of zoonotic
 diseases, and will be undertaken in line with published guidelines for best practice in
 scoping reviews.
- The review will be informed by a One Health lens, encompassing distal determinants and risk factors for spillover events and acknowledging the interconnectedness of human, animal and environmental health.

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2 3	1 •	Due to the complex drivers of spillover events, some potentially relevant policy
4 5	2	evaluations may not be identified in searches where their outcome measures are too
6	3	far removed from the spillover of zoonotic diseases.
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1 Introduction

The increasing incidence of zoonotic emerging infectious diseases (EIDs) has been attributed to ecological, behavioural and socioeconomic change, and is predicted to continue in the coming years (1). Higher levels of anthropogenic activity, including agricultural intensification, urbanisation and other forms of land use change, have led to increased interactions between wildlife, humans and livestock, increasing the risk of cross-species transmission (2). In response, a call has been issued by leading organisations and experts, including the United Nations Environment Programme, the International Livestock Research Institute and the Intergovernmental Science-Policy Platform on Biodiversity and Ecosystem Services, to complement reactive policy responses with policies that prevent zoonotic EIDs (1,3-6).

20 11 A preventive medicine approach to tackling the 'causes of the causes' of spillover events

Zoonotic spillover, defined as the transmission of a pathogen from a vertebrate animal to a human, depends on the alignment of ecological, epidemiological and behavioural factors (7). Zoonotic pathogens must meet a series of conditions in order to induce spillover infections in humans, including appropriate density and distribution of reservoir hosts, pathogen prevalence, infection intensity and human exposure (7). Across this transmission pathway, a number of drivers of zoonotic spillover have been identified, including changes in wildlife and livestock populations (8); deforestation, urbanisation and other forms of land use change (9); and a variety of necessary human practices including hunting, farming, animal husbandry, and trade (5,6,10,11).

These large-scale changes have on multiple occasions given rise to spillover events, sometimes involving diseases with epidemic or pandemic potential. In this context, the concept of preventive medicine can be useful in determining how to intervene to reduce the risk of spillover events (12), where spillover events may be understood as 'deviant' cases situated within a much broader 'population' of non-events or possible events. A broad-based approach to tackling the 'causes of the causes' of these events could be a key part of the solution.

Considering the causes of the causes of spillover events, there are a number of modifiable determinants that may be targeted with preventive interventions. These interventions include surveillance of pools of viruses in wildlife and management of wildlife populations (13); enhanced food safety measures in both the wildlife and livestock value chain, pre- and post-farm gate (11,14–16); replacement of traditional 'wet' markets with supermarkets (17); controls on wildlife hunting, trade and consumption (10,18,19); and phasing out of unsustainable agriculture practices (5,20).

⁵⁸ 34 Preventive policies in a real-world context: multi-sectoral efforts, One Health governance and
 35 complex systems

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While some evaluative evidence exists around the effectiveness of these interventions (21-24), they have often been implemented as short- to medium-term programmes or academic investigations (5). In some cases, zoonoses have re-emerged after successful programmes have ended (24). As a result, experts have argued for the incorporation of successful interventions into policy frameworks, providing interventions with the sustainability required for long-term disease control (5). By policy, we refer here to public policy: 'a set of interrelated decisions taken by a political actor or group of actors concerning the selection of goals and the means of achieving them' (25). While policy implementation requires the involvement of public and private actors, we understand policy decisions as being ultimately in the hands of government, which for this review we extend to include supranational governing bodies, and having greater longevity compared to many programmes, which are often implemented for a fixed term.

Although the longevity and scope of government actions may make policy an effective vehicle
 for disease prevention, implementing policy is a complex process involving numerous
 stakeholders with competing views and interests (26). The responsibility for addressing
 zoonotic disease frequently spans multiple sectors of governance due to its relevance for both
 animals and humans. Where relevant policies are designed and implemented in isolation,
 opportunities for synergy may be missed and efforts may even be counter-productive.

Successful policy measures require not only a sound evidence base, but also governance structures that enable action to be taken. Given the range of possible risk factors that might contribute to emerging zoonoses, and the possible impacts of preventive policies, a One Health response has been advocated, requiring coordination between institutions and government departments involved in human and animal health, trade, agriculture and the environment (27). At the international level, the World Health Organization, the Food and Agriculture Organization and the World Organisation for Animal Health have endorsed a One Health policy framework to respond to zoonotic infectious diseases, emphasising collaboration between agencies (28). Within countries, national and local governments have also emphasised the need for multi-sectoral efforts, although many report that further integration is still required (29).

Further, given the complex social-ecological systems within which policies to prevent zoonotic spillover are implemented, the risk of unintended consequences is high. For example, bans on economic activities associated with higher risks of disease transmission may lead to the emergence or growth of illegal marketplaces where regulation is impossible (30,31). Region-specific closures of live animal markets have been shown to spread diseases further afield as vendors seek new venues to sell their animals (32). Meanwhile, attempts to manage

populations of wild animals may alter disease dynamics, unintentionally increasing the risk of
 spillover into livestock or people (33).

Given these particular characteristics of policy development and implementation, they may be usefully considered as a particular case of intervention, and the evidence around them assessed accordingly. Different types of interventions might be more or less feasibly implemented by governments (or their partners), and their impacts might be different given potentially more complex implementation contexts, longer timespans and broader geographic ranges. Evaluations of these policies should also include consideration and monitoring of potential unintended consequences. In order to facilitate this, multi-sectoral involvement in both policy development and evaluation may be required.

20 11 Aims and scope

Preventive approaches to managing epidemic and pandemic infectious diseases once they have entered human populations have been systematically catalogued in the medical literature (34–40). These measures include hand washing, face masks, school closures, and contract tracing and case isolation. Further upstream, systematic reviews of interventions targeting the spillover pathway have predominantly focused on programmes rather than policies, and have been restricted by various characteristics such as geographic region (23) or disease type (24), or focused on programmes with an explicit endorsement of a One Health approach (22). In consequence, a comprehensive understanding of how policies have been evaluated, and what evidence there is of their effectiveness, is lacking. To address these research gaps, our objectives are to:

- Identify evaluations of population health policies that target the determinants of zoonotic spillover;
 - 2. Synthesise the nature of how the interventions were evaluated; and
- 3. Examine the effectiveness of the interventions and identify gaps in the literature.

Our approach to identifying and analysing this literature will be informed by a One Health lens,
 acknowledging the inter-connectedness of human, animal and environmental health.

1 2			
3 4	1	Methods and analysis	
5 6	2	We will conduct a systematic scoping review	of evaluations of policies aimed at preventing
7	3	zoonotic spillover events. The scoping revie	ew will be conducted in line with guidelines
8 9	4	published by Arksey and O'Malley and refin	ed by Levac and colleagues (41-43), which
10	5	emphasise an iterative approach suited to an e	exploratory research question.
11 12 13	6	Stage 1: Identifying the research question	
14 15	7	The aim of this review is to use a One Health le	ns to identify and describe the range of policies
15 16	8	that have been evaluated, the approaches	to evaluation, and the evaluative evidence.
17 18	9	Informed by this aim, our research questions a	ire:
19 20	10	1. What population health policies aimed	at preventing the spillover of emerging
21 22	11	infectious diseases of zoonotic origin h	ave been evaluated?
22	12	a. What types of policies?	
24 25	13	b. Which policy actors (single depa	artment, multi-sectoral, whole of government)?
26	14	2. How are these interventions evaluated	?
27 28	15	a. What methods/study designs?	
29	16	b. What outcomes?	
30 31	17	3. What is the evidence around the relativ	e effectiveness of these interventions?
32 33	18	Stage 2: Identifying relevant studies	
34 35	19	We will systematically search four electronic of	databases (Medline, Scopus, Web of Science,
36 37	20	Global Health). The search strategy will be in	formed by the main concepts in our research
38	21	question using the PICO framework designed t	to frame research questions in evidence-based
39 40	22	medicine (44) (see Table 1). See Supplementa	ary File 1 for details of search strategy.
41 42	23	Table 1: Concepts from the research question	used in developing the search strategy
43 44	24	according to the PICO Framework.	
45		Population	All actors within systems (animal or human
40 47			populations: institution including
48 40			and industry)
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51 52		Intervention	Preventive public policy
53 54 55		Comparator	N/A
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Outcome	Spillover events, or any upstream
	determinants or risk factors for spillover
	events

Stage 3: Study selection

Records identified through the searches will be collated and double screened using the online
 platform Covidence (45). Studies will be included where they meet all of the following criteria:

- 1. Primary empirical study from any country or region;
- 2. Report empirical findings from an evaluation of any sort; and
- 3. Focus on a policy implemented by government that targets the determinants of zoonotic spillover.

Titles and abstracts will initially be screened, followed by full-text screening. Title and abstract screening of an initial set of 100 papers will be undertaken by two independent researchers. Results will be compared in order to ensure consistency in decisions around study eligibility, and discrepancies resolved through discussion of the inclusion criteria. This process will be repeated until an acceptable level of agreement (>90%) is reached. The remaining papers will then be screened by one of the two reviewers. Full-text screening will be undertaken by two independent researchers and discrepancies will be resolved by discussing reasons for inclusion or exclusions among the screeners.

- In line with published guidelines, the approach to study selection may be refined iteratively
 when reviewing articles for inclusion (41–43).
- 40 19 Stage 4: Charting the data

Data charting will be conducted using a data charting form designed to identify the information required to answer the research question and sub-research questions (see Table 2). As recommended, the data charting form will be piloted with ten records to ensure that it is consistent with the research question, and the data charting form will be revised iteratively in order to ensure the purpose of the research is being met (41–43).

25 Table 2: Data charting form

Record	Author(s)
	Year
Policy	Country

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	World region (World Bank grouping) (46)
	Country income (World Bank grouping) (46)
	Disease
	Stakeholders or sector responsible for implementing policy (retail,
	agriculture, conservation, etc.)
	Implementation date (start date, or range if the policy has been changed)
	Intervention type
	Location along spillover pathway adapted from Plowright et al. (7,21)
	Policy level (local, national, regional, global)
	Multi-sectoral initiative (Y/N)
	Sector(s) responsible for policy
Evaluation	Aim
	Type (Process/outcome)
	Study design
	Period of observation
	Outcome measure(s) and change in measure(s)
	Consideration of unintended consequences (Y/N)
	If yes, which unintended consequences? (e.g., economic outcomes, food
	security)

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Stage 5: Collating, summarising and reporting the results

We will undertake quality assessment of the included studies using the Quality Assessment
Tool for Quantitative Studies developed by the Effective Public Health Practice Project (47),
which has previously been used to assess the quality of natural experiments including public
policy evaluations (48).

We will analyse the extracted data, presenting a numerical summary of the included studies in table form, allowing us to describe the range of policy interventions that have been evaluated, approaches to evaluation, and evidence of effectiveness. We will also conduct a thematic analysis of the contents of the included articles in order to identify, if possible, the challenges encountered in evaluating these policies, as well as insights into why policies
 succeeded or failed in achieving their aims.

3 Patient and public involvement

Patients or the public were not involved in the design, conduct, reporting or disseminationplans of our research.

6 Strengths and weaknesses of the study

To our knowledge, this is the first attempt to systematically identify and document evaluations of policies aiming to prevent the spillover of zoonoses into human populations. However, because of the complex drivers of spillover events, some potentially relevant policy evaluations may be excluded where their outcome measures are too far removed from the spillover of zoonotic diseases. For example, it has been hypothesised that declines in vulture populations may increase the risk of disease transmission by increasing the number of uneaten carcasses, as well as, potentially, the population of feral dogs (49). In 2006, India, Pakistan and Nepal implemented a ban on the veterinary drug diclofenac, which had been identified as a driver of declining vulture populations. While policy evaluations suggest that this ban has resulted in a resurgence of vultures (50-53), the knock-on effects of this on zoonotic disease transmission risk have not been included in these evaluations. While relevant, such evaluations will be difficult to systematically identify as they make no reference to zoonotic disease.

In addition, this review will focus on policy evaluations that have been reported in the peer-reviewed literature. Policies that have been implemented but not evaluated, or evaluated but not published in the academic literature, will therefore be excluded from this review. As a result, potentially effective and important policies in the prevention of zoonotic spillover events may not be identified. However, we hope that the findings from this review will highlight these gaps in the evaluative evidence. We also hope that this review, by extracting practical dimensions such as study design, outcome measures and the challenges encountered in the evaluation process, will support policymakers and researchers in carrying out policy evaluations in this space.

51 29 Ethics and dissemination

Formal ethical approval is not required, as the study does not involve primary data collection.
 The findings of this study will be disseminated through a peer-reviewed publication,
 presentations, and summaries for key stakeholders.

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3 4	1	List of abbreviations
5 6	2	EID: Emerging infectious disease
7 8 9	3	Declarations
10 11 12	4	Ethics approval and consent to participate
13 14 15	5	Not applicable.
16 17 18	6	Patient consent for publication
19 20	7	Not applicable.
21 22 23 24	8	Availability of data and material
25 26	9	Data sharing not applicable to this article as no datasets were generated or analysed
20 27 28	10	during the current study.
29 30 31	11	Competing interests
32 33 34	12	The authors declare that they have no competing interests.
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43 44 45	17	developing the protocol.
46 47 48	18	Contributors
49 50	19	CCA, KML and TLP conceived and designed the study. CCA prepared the
51 52	20	manuscript. All authors provided critical input on the manuscript and methods and
53 54	21	have read and approved the final manuscript.
55 56 57	22	Acknowledgements
58 59 60	23	Not applicable.

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Supplementary file 1

Search strings for academic databases

Scopus search, all terms in TITLE-ABS-KEY; Block 1 AND 2 AND 3 AND 4

P – Population: All actors within a system (no specific search terms)

I – Intervention: Preventive public policy

Block 1 – Policy interventions

policy OR law OR legal OR legislat* OR regulat* OR tariff OR subsidy OR tax OR ban OR "voluntary agreement" OR incentive OR fiscal OR guidelines OR govern* OR federal* OR closure OR closing OR state* OR "rest day*" OR "border control*" OR "habitat protection" OR "wetland protection" OR "supplement* fed" OR "supplement* feed*" OR "market size"

Block 2 – Prevention

Prevent* OR "ecological intervention*" OR "non-pharmaceutical intervention*" OR "public health" OR "risk management" OR "risk minimisation" OR "control strateg*" OR "outbreak risk" OR "reduc* W/5 transmission" OR "reduc* W/5 infection"

C – Comparator: N/A

O - Outcome: Zoonotic spillover events or their upstream determinants and risk factors

Block 3 - Zoonoses

Zika OR ebola OR covid-19 OR sars-cov-2 OR coronavirus OR sars OR mers OR h1n1 OR h7n9 OR h5n1 OR "one health" OR dengue OR "nipah virus" OR influenza OR zoonoses OR zoonosis OR zoonotic OR "West Nile" OR "HIV/AIDS" OR "avian flu" OR "hendra virus" OR "marburg virus" OR "yellow fever" OR "tick-borne encephalitis" OR "emerging infectious diseases" OR "emergent infectious diseases" OR brucellosis OR rabies OR chikungunya OR "bovine spongiform encephalopathy" OR "rift valley fever"

Block 4 – Spillover events

Spillover OR "spill over" OR "cross-species transmission" OR poultry OR wildlife OR bushmeat OR "bush meat" OR livestock OR "animal market*" OR "wet market*" OR "bird market*" OR horse* OR waterfowl OR fowl OR bat OR bats OR mammal* OR swine OR pig* OR poaching OR "pet trade" OR pork OR "trade W/5 animal"

PRISMA-P 2015 Checklist

This checklist has been adapted for use with systematic review protocol submissions to BioMed Central journals from Table 3 in Moher D et al: Preferred reporting items for systematic review and meta-analysis protocols (PRISMA-P) 2015 statement. *Systematic Reviews* 2015 **4**:1

An Editorial from the Editors-in-Chief of *Systematic Reviews* details why this checklist was adapted - **Moher D, Stewart L & Shekelle P:** Implementing PRISMA-P: recommendations for prospective authors. *Systematic Reviews* 2016 **5**:15

O a ati a m lt a m i a	ш		Information	reported	Line	
Section/topic	#		Yes	No	number(s)	
ADMINISTRATIVE INFO	RMAT	ION				
Title						
Identification	1a	Identify the report as a protocol of a systematic review			p.1 l.1-2	
Update	1b	If the protocol is for an update of a previous systematic review, identify as such		\boxtimes	N/A	
Registration	2	If registered, provide the name of the registry (e.g., PROSPERO) and registration number in the Abstract		\square	N/A	
Authors						
Contact	3а	Provide name, institutional affiliation, and e-mail address of all protocol authors; provide physical mailing address of corresponding author			p.1 l.3-25	
Contributions	3b	Describe contributions of protocol authors and identify the guarantor of the review			p.12 l.16-18	
Amendments	4	If the protocol represents an amendment of a previously completed or published protocol, identify as such and list changes; otherwise, state plan for documenting important protocol amendments		\square	N/A	
Support						
Sources	5a	Indicate sources of financial or other support for the review	\square		p.12 l.14-16	
Sponsor	5b	Provide name for the review funder and/or sponsor			p.12 l.14-16	
Role of sponsor/funder	5c	Describe roles of funder(s), sponsor(s), and/or institution(s), if any, in developing the protocol			p.12 l.16	
INTRODUCTION						
Rationale	6	Describe the rationale for the review in the context of what is already known			p.4-6	
Objectives	7	Provide an explicit statement of the question(s) the review will address with reference to			p.6 l.12-27	



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Saction/topia	#	Chacklist item	Information reported		Line	
Section/topic	#		Yes	No	number(s)	
		participants, interventions, comparators, and outcomes (PICO)				
METHODS			11			
Eligibility criteria	8	Specify the study characteristics (e.g., PICO, study design, setting, time frame) and report characteristics (e.g., years considered, language, publication status) to be used as criteria for eligibility for the review			p.8 l.2-8	
Information sources	9	Describe all intended information sources (e.g., electronic databases, contact with study authors, trial registers, or other grey literature sources) with planned dates of coverage			p.7 l.19-20	
Search strategy	10	Present draft of search strategy to be used for at least one electronic database, including planned limits, such that it could be repeated	\square		p.11	
STUDY RECORDS						
Data management	11a	Describe the mechanism(s) that will be used to manage records and data throughout the review	\square		p.8 I.3-4; 20-24	
Selection process	11b	State the process that will be used for selecting studies (e.g., two independent reviewers) through each phase of the review (i.e., screening, eligibility, and inclusion in meta-analysis)	\square		p.8 l.9-16	
Data collection process	11c	Describe planned method of extracting data from reports (e.g., piloting forms, done independently, in duplicate), any processes for obtaining and confirming data from investigators	\square		p.8 l.19-25	
Data items	12	List and define all variables for which data will be sought (e.g., PICO items, funding sources), any pre-planned data assumptions and simplifications	\square		Table 2, p.8-9	
Outcomes and prioritization	13	List and define all outcomes for which data will be sought, including prioritization of main and additional outcomes, with rationale		\square	N/A – scoping review	
Risk of bias in individual studies	14	Describe anticipated methods for assessing risk of bias of individual studies, including whether this will be done at the outcome or study level, or both; state how this information will be used in data synthesis			p.9 l.3-6	
DATA						
	15a	Describe criteria under which study data will be quantitatively synthesized			Table 2, p.8-9	
Synthesis	15b	If data are appropriate for quantitative synthesis, describe planned summary measures, methods of handling data, and methods of combining data from studies, including any planned exploration of consistency (e.g., <i>I</i> ² , Kendall's tau)		\square	N/A	
	15c	Describe any proposed additional analyses (e.g., sensitivity or subgroup analyses, meta-regression)		\square	N/A	
	15d	If quantitative synthesis is not appropriate, describe the type of summary planned			p.9 l.7-10, p.10 l.1-2	
Meta-bias(es)	16	Specify any planned assessment of meta-bias(es) (e.g., publication bias across studies, selective		\square	N/A – scoping	



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Saction/tonic	#	Checklist item	Information	Line	
Section/topic	π		Yes	No	number(s)
		reporting within studies)			review
Confidence in cumulative evidence	17	Describe how the strength of the body of evidence will be assessed (e.g., GRADE)			N/A – scoping review

The body of evidence will be assessed (e.g., GK.



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Policies to prevent zoonotic spillover: protocol for a systematic scoping review of evaluative evidence

Journal:	BMJ Open
Manuscript ID	bmjopen-2021-058437.R1
Article Type:	Protocol
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Primary Subject Heading :	Health policy
Secondary Subject Heading:	Global health, Health policy, Infectious diseases
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1 2		
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1 Abstract

2 Introduction

The increasing incidence of pathogen transmission from animals to humans (zoonotic spillover events) has been attributed to behavioural practices and ecological and socioeconomic change. As these events sometimes involve pathogens with epidemic or pandemic potential, they pose a serious threat to population health. Public policies may play a key role in preventing these events. The aim of this review is to identify evaluations of public policies that target the determinants of zoonotic spillover, examining approaches taken to evaluation, choice of outcomes measures and evidence of effectiveness. Our approach to identifying and analysing this literature will be informed by a One Health lens, acknowledging the inter-connectedness of human, animal and environmental health.

12 Methods and analysis

A systematic scoping review methodology will be used. To identify articles, we will search Medline, SCOPUS, Web of Science and Global Health in March 2022 using search terms combining animal health and the animal-human interface, public policy, prevention and zoonoses. We will screen titles and abstracts and extract data according to published guidelines for scoping reviews. All evaluations of public policies aiming to prevent zoonotic spillover events will be eligible for inclusion. We will summarise key data from each study, mapping policies along the spillover pathway and outlining the range of policies, approaches to evaluation and outcome measures. Review findings will provide a useful reference for researchers and practitioners, outlining the state of the evaluative evidence around policies to prevent zoonotic spillover.

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4123Ethics and dissemination

Formal ethical approval is not required, because the study does not involve primary data
collection. The findings of this study will be disseminated through a peer-reviewed publication,
presentations, and summaries for key stakeholders.

48 27 Strengths and limitations

- This scoping review protocol outlines the first piece of work to systematically identify and review evaluations of public policies designed to prevent zoonotic spillover, and will be undertaken in line with published guidelines for best practice in scoping reviews.
- The review will be informed by a One Health lens, encompassing distal determinants and risk factors for spillover events and acknowledging the interconnectedness of human, animal and environmental health.

1 2 3 4 5 6 7 8	1 2 3	 Due to the complex drivers of spillover events, some potentially relevant policy evaluations may not be identified where outcome measures are too far removed from zoonotic spillover.
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1 Introduction

The increasing incidence of zoonotic emerging infectious diseases (EIDs) has been attributed to behavioural practices and ecological and socioeconomic change, and is predicted to continue in the coming years (1). Higher levels of anthropogenic activity, including agricultural intensification, urbanisation and other forms of land use change, have led to increased interactions between wildlife, humans and livestock, increasing the risk of cross-species transmission (2,3). In response, a call has been issued by leading organisations and experts, including the United Nations Environment Programme, the International Livestock Research Institute and the Intergovernmental Science-Policy Platform on Biodiversity and Ecosystem Services, to complement reactive policy responses with policies that prevent zoonotic EIDs (1,4–7).

12 Preventing zoonotic spillover from a One Health perspective

Zoonotic spillover, defined as the transmission of a pathogen from an animal to a human, depends on the alignment of ecological, epidemiological and behavioural factors (8). Zoonotic pathogens must meet a series of conditions in order to induce spillover infections in humans, including appropriate density and distribution of reservoir hosts, pathogen prevalence, infection intensity and human exposure (8). Across this transmission pathway, a number of drivers of zoonotic spillover have been identified, including changes in wildlife and livestock populations (9); deforestation, urbanisation and other forms of land use change (10); and a variety of human practices including hunting, farming, animal husbandry, keeping of exotic pets and trade (6,7,11,12). These large-scale changes have on multiple occasions given rise to spillover events, sometimes involving pathogens with epidemic or pandemic potential.

A One Health perspective, which recognises the health of humans, animals and ecosystems as being closely linked and inter-dependent (13), can be useful in conceptualizing a range of potential determinants of spillover events. From this perspective, interventions could include surveillance of pools of viruses in wildlife and management of wildlife populations (14); enhanced food safety measures in both the wildlife and livestock value chain, pre- and post-farm gate (12,15–17); replacement of traditional 'wet' markets with supermarkets (18); controls on wildlife hunting, trade and consumption (11,19,20); and phasing out of unsustainable agriculture practices (6,21).

While some evaluative evidence exists around the effectiveness of interventions (22–25), they
 have often been implemented as short- to medium-term programmes or academic
 investigations (6). In some cases, zoonoses have re-emerged after successful programmes
 have ended (25). As a result, experts have argued for the incorporation of successful

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interventions into policy frameworks, providing interventions with the sustainability required for
 long-term disease control (6).

3 Governance, systems and the role of multi-sectoral actors

Public policy is 'a set of interrelated decisions taken by a political actor or group of actors concerning the selection of goals and the means of achieving them' (26). Public policy decisions are ultimately in the hands of government and supranational governing bodies, and have greater longevity compared to many programmes, which are often implemented for a fixed term. Non-government actors, including vested interest stakeholders, can also play a powerful role in shaping government decisions (27,28).

Although the longevity and scope of government actions may make policy an effective vehicle for prevention of emergent diseases, implementing policy is a complex process involving numerous stakeholders with competing views and interests (29). The responsibility for addressing zoonotic disease frequently spans multiple sectors of governance due to its relevance for both animals and humans. Where relevant policies are designed and implemented in isolation, opportunities for synergy may be missed and efforts may even be counter-productive.

Successful policy measures require not only a sound evidence base, but also governance structures that enable action to be taken. Given the range of possible risk factors that might contribute to emerging zoonoses, and the possible impacts of policies to prevent zoonotic spillover, a One Health response has been advocated, requiring coordination between institutions and government departments involved in human and animal health, trade, agriculture and the environment (30). At the international level, the World Health Organization, the Food and Agriculture Organization and the World Organisation for Animal Health have endorsed a One Health policy framework to respond to zoonotic infectious diseases, emphasising collaboration between agencies (31). Within countries, national and local governments have also emphasised the need for multi-sectoral efforts, although many report that further integration is still required (32).

Further, given the complex social-ecological systems within which policies to prevent zoonotic spillover are implemented, the risk of unintended consequences is high. For example, region-specific closures of live animal markets have been shown to spread pathogens further afield as vendors seek new venues to sell their animals (33). Meanwhile, attempts to manage populations of wild animals may alter pathogen dynamics, unintentionally increasing the risk of spillover into livestock or people (34).

Given these particular characteristics of policy development and implementation, they may be
 usefully considered as a particular case of intervention, and the evidence around them

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1 assessed accordingly. Different types of interventions might be more or less feasibly 2 implemented by governments (or their partners), and their impacts might be different given 3 potentially more complex implementation contexts, longer timespans and broader geographic 4 ranges. Evaluations of these policies should also include consideration and monitoring of 5 potential unintended consequences. In order to facilitate this, multi-sectoral involvement in 6 both policy development and evaluation may be required.

7 Aims and scope

Approaches to managing epidemic and pandemic infectious pathogens once they have entered human populations have been systematically catalogued in the medical literature (35-41). These measures include hand washing, face masks, school closures, and contact tracing and case isolation. Further upstream, systematic reviews of interventions targeting the spillover pathway have predominantly focused on programmes rather than policies, and have been restricted by various characteristics such as geographic region (24) or pathogen type (25), or focused on programmes with an explicit endorsement of a One Health approach (23). In consequence, a comprehensive understanding of how policies to prevent zoonotic spillover have been evaluated, and what evidence there is of their effectiveness, is lacking. To address these research gaps, our objectives are to:

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1. Identify evaluations of policies that target the determinants of zoonotic spillover

- included in the spillover pathway (8) (i.e. human and animal health and interactions);
- 20 2. Synthesise the nature of how the interventions were evaluated; and
 - 3. Examine the effectiveness of the interventions and identify gaps in the literature.

Our approach to identifying and analysing this literature will be informed by a One Health lens,
 acknowledging the inter-connectedness of human, animal and environmental health.

2 3	1	
4	I	Methods and analysis
5 6	2	We will conduct a systematic scoping review of evaluations of policies aimed at preventing
7	3	zoonotic spillover events. The scoping review will be conducted in line with guidelines
8 9	4	published by Arksey and O'Malley and refined by Levac and colleagues (42-44), which
10	5	emphasise an iterative approach suited to an exploratory research question.
11 12 13 14	6	Stage 1: Identifying the research question
	7	The aim of this review is to use a One Health lens to identify and describe the range of policies
15 16	8	that have been evaluated, the approaches to evaluation, and the evaluative evidence.
17	9	Informed by this aim, our research questions are:
19	10	1 What policies aimed at preventing zoonotic spillover have been evaluated?
20 21	11	a What types of policies?
22	12	b. Which policy actors (single department, multi-sectoral, whole of government)?
23 24	12	2. How are these interventions evaluated?
25	13	2. How are these interventions evaluated?
26 27	14	a. vvnat methods/study designs?
28	15	b. vvnat outcomes?
29 30	16	3. What is the evidence around the relative effectiveness of these interventions?
31 32	17	Stage 2: Identifying relevant studies
33	18	We will systematically search four electronic databases (Medline, Scopus, Web of Science,
34 35	19	Global Health) in March 2022. The search strategy is organized by the main concepts in our
36	20	research question: the spillover pathway; public policy; prevention; and zoonotic pathogens (
 37 38 39 40 41 42 43 44 	21	see Supplementary File 1 for details of search strategy). The search strategy was developed
	22	iteratively, informed by existing systematic reviews focused on related concepts (24,45–49)
	23	and known indicator papers meeting inclusion criteria.
	24	Stage 3: Study selection
45	25	Records identified through the searches will be collated and double screened using the online
46 47	26	platform Covidence (50). Studies will be included where they meet all of the following criteria:
48 49	27	1 Primary empirical study from any country or region with English-language abstracts:
50	28	 Report empirical findings from an evaluation of any sort: and
51 52	20	2. Report empirical minings from an evaluation of any sort, and
52 53	29	5. Focus on a policy implemented by government that targets the determinants of
54 57	30	zoonotic spillover.
55 56	31	Titles and abstracts will initially be screened, followed by full-text screening. Title and abstract
57	32	screening of an initial set of 100 papers will be undertaken by two independent researchers.
59 60	33	Results will be compared in order to ensure consistency in decisions around study eligibility,

and discrepancies resolved through discussion of the inclusion criteria. This process will be repeated until an acceptable level of agreement (>90%) is reached. The remaining papers will then be screened by one of the two reviewers. Full-text screening will be undertaken by two independent researchers and discrepancies will be resolved by discussing reasons for inclusion or exclusions among the screeners. Studies with full-texts in languages other than English will be eligible for inclusion if they include an English-language abstract. Full-text studies published in French, Spanish or Chinese will be single-screened by a member of the research team fluent in that language. Studies published in other languages will be translated as necessary.

In line with published guidelines, the approach to study selection may be refined iteratively
 when reviewing articles for inclusion (42–44). Reporting on the search and screening process
 will follow the guidelines provided in the Preferred Reporting Items for Systematic Reviews
 and Meta-Analyses Extension for Scoping Reviews (51).

2425 14 Stage 4: Charting the data

Data charting will be conducted using a data charting form designed to identify the information required to answer the research question and sub-research questions (see Supplementary File 2). As recommended, the data charting form will be piloted with ten records to ensure that it is consistent with the research question, and the data charting form will be revised iteratively in order to ensure the purpose of the research is being met (42-44).

³⁵ 20 Stage 5: Collating, summarising and reporting the results
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We will undertake quality assessment of the included studies using the Quality Assessment
Tool for Quantitative Studies developed by the Effective Public Health Practice Project (52),
which has previously been used to assess the quality of natural experiments including public
policy evaluations (53).

We will analyse the extracted data, presenting a numerical summary of the included studies in table form, allowing us to describe the range of policy interventions that have been evaluated, approaches to evaluation, and evidence of effectiveness. We will also conduct a thematic analysis of the contents of the included articles in order to identify, if possible, barriers and facilitators to implementing and evaluating these policies, as well as insights into why policies succeeded or failed in achieving their aims.

54 31 Patient and public involvement 55

This scoping review is being undertaken as part of a larger project involving policy actors at
 national and international levels as research team members, knowledge users and
 participants. Insights from the project have informed protocol development and stakeholders

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are able to provide input and perspectives on the results of the review. Project-level
 dissemination events involving policy stakeholders are also planned, where findings from the
 proposed review will be shared.

4 Strengths and weaknesses of the study

To our knowledge, this is the first attempt to systematically identify and document evaluations of policies aiming to prevent the spillover of zoonoses into human populations. However, because of the complex drivers of spillover events, some potentially relevant policy evaluations may be excluded where their outcome measures are too far removed from zoonotic spillover. For example, it has been hypothesised that declines in vulture populations may increase the risk of pathogen transmission by increasing the number of uneaten carcasses, as well as, potentially, the population of feral dogs (54). In 2006, India, Pakistan and Nepal implemented a ban on the veterinary drug diclofenac, which had been identified as a driver of declining vulture populations. While policy evaluations suggest that this ban has resulted in a resurgence of vultures (55–58), the knock-on effects of this on zoonotic pathogen transmission risk have not been included in these evaluations. While relevant, such evaluations will be difficult to systematically identify as they make no reference to zoonotic disease.

In addition, this review will focus on policy evaluations that have been reported in the peer-reviewed literature. Policies that have been implemented but not evaluated, or evaluated but not published in the academic literature, will therefore be excluded from this review. As a result, potentially effective and important policies in the prevention of zoonotic spillover events may not be identified. However, we hope that the findings from this review will highlight these gaps in the evaluative evidence. We also hope that this review, by extracting practical dimensions such as study design, outcome measures and the challenges encountered in the evaluation process, will support policymakers and researchers in carrying out policy evaluations in this space.

27 Ethics and dissemination

Formal ethical approval is not required, because the study does not involve primary data
 collection. The findings of this study will be disseminated through a peer-reviewed publication,
 presentations, and summaries for key stakeholders.

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1	List of abbreviations
2	EID: Emerging infectious disease
3	Declarations
4	Ethics approval and consent to participate
5	Not applicable.
6	Patient consent for publication
7	Not applicable.
8	Availability of data and material
9 10	Data sharing not applicable to this article as no datasets were generated or analysed during the current study.
11	Competing interests
12	The authors declare that they have no competing interests.
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17	developing the protocol.
18	Contributors
19	CCA, KML and TLP conceived and designed the study. CCA prepared the
20	manuscript. KML, TLP, RA, AA, MB, JC, RL, AR, KCT, AMV, MW and AY provided
21	critical input on the manuscript and methods and have read and approved the final
22	manuscript.

23 Acknowledgements

Not applicable.

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Supplementary file 1

Example search string framed by core concepts

Example from Scopus search, all terms in TITLE-ABS-KEY; Block 1 AND 2 AND 3 AND 4

Block 1 - Spillover pathway (Animal populations and human-animal interface)

Spillover OR "spill over" OR "cross-species transmission" OR poultry OR wildlife OR bushmeat OR "bush meat" OR livestock OR "animal market" OR "wet market" OR "bird market" OR horse* OR waterfowl OR fowl OR bat OR bats OR mammal* OR swine OR pig* OR poaching OR "pet trade" OR pork OR "trade W/5 animal"

Block 2 – Public policy

policy OR law OR legal OR legislat* OR regulat* OR tariff OR subsidy OR tax OR ban OR "voluntary agreement" OR incentive OR fiscal OR guidelines OR govern* OR federal* OR closure OR closing OR state* OR "rest day*" OR "border control*" OR "habitat protection" OR "wetland protection" OR "supplement* fed" OR "supplement* feed*" OR "market size"

Block 3 – Prevention

Prevent* OR "ecological intervention*" OR "non-pharmaceutical intervention*" OR "public health" OR "risk management" OR "risk minimisation" OR "control strateg*" OR "outbreak risk" OR "reduc* W/5 transmission" OR "reduc* W/5 infection"

Block 4 - Zoonotic pathogens

Zika OR ebola OR covid-19 OR sars-cov-2 OR coronavirus OR sars OR mers OR h1n1 OR h7n9 OR h5n1 OR "one health" OR dengue OR "nipah virus" OR influenza OR zoonoses OR zoonosis OR zoonotic OR "West Nile" OR "HIV/AIDS" OR "avian flu" OR "hendra virus" OR "marburg virus" OR "yellow fever" OR "tick-borne encephalitis" OR "emerging infectious diseases" OR "emergent infectious diseases" OR brucellosis OR rabies OR chikungunya OR "bovine spongiform encephalopathy" OR "rift valley fever"

Search strings for all included academic databases

Scopus search

TITLE-ABS-KEY(Zika OR ebola OR covid-19 OR sars-cov-2 OR coronavirus OR sars OR mers OR h1n1 OR h7n9 OR h5n1 OR "one health" OR dengue OR "nipah virus" OR influenza OR zoonoses OR zoonosis OR zoonotic OR "West Nile" OR "HIV/AIDS" OR "avian flu" OR "hendra virus" OR "marburg virus" OR "yellow fever" OR "tick-borne encephalitis" OR "emerging

infectious diseases" OR "emergent infectious diseases" OR brucellosis OR rabies OR chikungunya OR "bovine spongiform encephalopathy" OR "rift valley fever") AND TITLE-ABS-KEY(Spillover OR "spill over" OR "cross-species transmission" OR poultry OR wildlife OR bushmeat OR "bush meat" OR livestock OR "animal market*" OR "wet market*" OR "bird market*" OR horse* OR waterfowl OR fowl OR bat OR bats OR mammal* OR swine OR pig* OR poaching OR "pet trade" OR pork OR "trade W/5 animal") AND TITLE-ABS-KEY(policy OR law OR legal OR legislat* OR regulat* OR tariff OR subsidy OR tax OR ban OR "voluntary agreement" OR incentive OR fiscal OR guidelines OR govern* OR federal* OR closure OR closing OR state* OR "rest day*" OR "border control*" OR "habitat protection" OR "wetland protection" OR "supplement* fed" OR "supplement* feed*" OR "market size") AND TITLE-ABS-KEY(Prevent* OR "ecological intervention*" OR "non-pharmaceutical intervention*" OR "public health" OR "risk management" OR "risk minimisation" OR "control strateg*" OR "outbreak risk" OR "reduc* W/5 transmission" OR "reduc* W/5 infection")

Medline search

(((Zika[Title/Abstract] OR ebola[Title/Abstract] OR covid-19[Title/Abstract] OR sars-cov-2[Title/Abstract] OR coronavirus[Title/Abstract] OR sars[Title/Abstract] OR mers[Title/Abstract] OR h1n1[Title/Abstract] OR h7n9[Title/Abstract] OR h5n1[Title/Abstract] OR "one health"[Title/Abstract] OR dengue[Title/Abstract] OR "nipah virus"[Title/Abstract] OR influenza[Title/Abstract] OR zoonoses[Title/Abstract] OR zoonosis[Title/Abstract] OR zoonotic[Title/Abstract] OR "West Nile"[Title/Abstract] OR "HIV/AIDS"[Title/Abstract] OR "avian flu"[Title/Abstract] OR "hendra virus"[Title/Abstract] OR "marburg virus"[Title/Abstract] OR "yellow fever"[Title/Abstract] OR "tick-borne encephalitis"[Title/Abstract] OR "emerging infectious diseases"[Title/Abstract] OR "emergent infectious diseases"[Title/Abstract] OR brucellosis[Title/Abstract] OR rabies[Title/Abstract] OR chikungunya[Title/Abstract] OR "bovine spongiform encephalopathy"[Title/Abstract] OR "rift valley fever"[Title/Abstract] OR zoonoses [mesh]) AND (Spillover[Title/Abstract] OR "spill over"[Title/Abstract] OR "cross-species transmission"[Title/Abstract] OR poultry[Title/Abstract] OR wildlife[Title/Abstract] OR bushmeat[Title/Abstract] OR "bush meat"[Title/Abstract] OR livestock[Title/Abstract] OR "animal market"[Title/Abstract] OR "animal markets"[Title/Abstract] OR "wet market"[Title/Abstract] OR "wet markets"[Title/Abstract] OR "bird market"[Title/Abstract] OR "bird markets"[Title/Abstract] OR horse[Title/Abstract] OR horses[Title/Abstract] OR waterfowl[Title/Abstract] OR fowl[Title/Abstract] OR bat[Title/Abstract] OR bats[Title/Abstract] OR mammal[Title/Abstract] OR

mammals[Title/Abstract] OR mammalian[Title/Abstract] OR swine[Title/Abstract] OR pig[Title/Abstract] OR pigs[Title/Abstract] OR poaching[Title/Abstract] OR "pet trade"[Title/Abstract] OR pork[Title/Abstract] OR animal N5 trade[Title/Abstract] OR disease reservoir [mesh])) AND (policy[Title/Abstract] OR law[Title/Abstract] OR legal[Title/Abstract] OR legislation[Title/Abstract] OR legislative[Title/Abstract] OR legislating[Title/Abstract] OR regulation[Title/Abstract] OR regulations[Title/Abstract] OR regulatory[Title/Abstract] OR tariff[Title/Abstract] OR subsidy[Title/Abstract] OR tax[Title/Abstract] OR ban[Title/Abstract] OR "voluntary agreement"[Title/Abstract] OR incentive[Title/Abstract] OR fiscal[Title/Abstract] OR guidelines[Title/Abstract] OR government[Title/Abstract] OR governments[Title/Abstract] OR federal[Title/Abstract] OR federally[Title/Abstract] OR closure[Title/Abstract] OR closing[Title/Abstract] OR state[Title/Abstract] OR "rest day"[Title/Abstract] OR "rest days"[Title/Abstract] OR "border control"[Title/Abstract] OR "border controls"[Title/Abstract] OR "habitat protection"[Title/Abstract] OR "wetland protection"[Title/Abstract] OR "supplemental feeding"[Title/Abstract] OR "market size"[Title/Abstract])) AND (Prevent[Title/Abstract] OR prevention[Title/Abstract] OR "ecological intervention"[Title/Abstract] OR "ecological interventions"[Title/Abstract] OR "non-pharmaceutical intervention"[Title/Abstract] OR "nonpharmaceutical interventions"[Title/Abstract] OR "public health"[Title/Abstract] OR "risk management"[Title/Abstract] OR "risk minimisation"[Title/Abstract] OR "control strategy"[Title/Abstract] OR "control strategies"[Title/Abstract] OR "outbreak risk"[Title/Abstract] OR reducing N5 transmission[Title/Abstract] OR reducing N5 infection[Title/Abstract])

Web of knowledge search

AB=(Zika OR ebola OR covid-19 OR sars-cov-2 OR coronavirus OR sars OR mers OR h1n1 OR h7n9 OR h5n1 OR "one health" OR dengue OR "nipah virus" OR influenza OR zoonoses OR zoonosis OR zoonotic OR "West Nile" OR "HIV/AIDS" OR "avian flu" OR "hendra virus" OR "marburg virus" OR "yellow fever" OR "tick-borne encephalitis" OR "emerging infectious diseases" OR "emergent infectious diseases" OR brucellosis OR rabies OR chikungunya OR "bovine spongiform encephalopathy" OR "rift valley fever")

AND

AB=(Spillover OR "spill over" OR "cross-species transmission" OR poultry OR wildlife OR bushmeat OR "bush meat" OR livestock OR "animal market*" OR "wet market*" OR "bird

market*" OR horse* OR waterfowl OR fowl OR bat OR bats OR mammal* OR swine OR pig* OR poaching OR "pet trade" OR pork OR trade NEAR animal)

AND

AB=(policy OR law OR legal OR legislat* OR regulat* OR tariff OR subsidy OR tax OR ban OR "voluntary agreement" OR incentive OR fiscal OR guidelines OR govern* OR federal* OR closure OR closing OR state* OR "rest day*" OR "border control*" OR "habitat protection" OR "wetland protection" OR "supplement* fed" OR "supplement* feed*" OR "market size")

AND

AB=(Prevent* OR "ecological intervention*" OR "non-pharmaceutical intervention*" OR "public health" OR "risk management" OR "risk minimisation" OR "control strateg*" OR "outbreak risk" OR reduc* NEAR transmission OR reduc* NEAR infection)

Ovid Global Health database search (all in abstract)

Zika OR ebola OR covid-19 OR sars-cov-2 OR coronavirus OR sars OR mers OR h1n1 OR h7n9 OR h5n1 OR one health OR dengue OR nipah virus OR influenza OR zoonoses OR zoonosis OR zoonotic OR West Nile OR HIV/AIDS OR avian flu OR hendra virus OR marburg virus OR yellow fever OR tick-borne encephalitis OR emerging infectious diseases OR emergent infectious diseases OR brucellosis OR rabies OR chikungunya OR bovine spongiform encephalopathy OR rift valley fever

AND

Spillover OR spill over OR cross-species transmission OR poultry OR wildlife OR bushmeat OR bush meat OR livestock OR animal market* OR wet market* OR bird market* OR horse* OR waterfowl OR fowl OR bat OR bats OR mammal* OR swine OR pig* OR poaching OR pet trade OR pork OR (trade adj5 animal)

AND

policy OR law OR legal OR legislat* OR regulat* OR tariff OR subsidy OR tax OR ban OR voluntary agreement OR incentive OR fiscal OR guidelines OR govern* OR federal* OR closure OR closing OR state* OR rest day* OR border control* OR habitat protection OR wetland protection OR supplement* fed OR supplement* feed* OR market size

AND

Prevent* OR ecological intervention* OR non-pharmaceutical intervention* OR public health OR risk management OR risk minimisation OR control strateg* OR outbreak risk OR (reduc* adj5 transmission) OR (reduc* adj5 infection)

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Supplementary file 2

Data charting form

Record	Author(s)
	Year
Policy	Country
	World region (World Bank grouping) (44)
	Country income (World Bank grouping) (44)
	Disease
	Stakeholders or sector responsible for implementing policy (retail,
	agriculture, conservation, etc.)
	Implementation date (start date, or range if the policy has been changed)
	Intervention type
	Location along spillover pathway adapted from Plowright et al. (7,21)
	Policy level (local, national, regional, global)
	Multi-sectoral initiative (Y/N)
	Sector(s) responsible for policy
Evaluation	Aim
	Type (Process/outcome)
	Study design
	Theoretical framework and/or logic model underpinning evaluation (if
	described)
	Period of observation
	Outcome measure(s) and change in measure(s)
	Consideration of unintended consequences (Y/N)
	If yes, which unintended consequences? (e.g., economic outcomes, food security)

PRISMA-P 2015 Checklist

This checklist has been adapted for use with systematic review protocol submissions to BioMed Central journals from Table 3 in Moher D et al: Preferred reporting items for systematic review and meta-analysis protocols (PRISMA-P) 2015 statement. *Systematic Reviews* 2015 **4**:1

An Editorial from the Editors-in-Chief of *Systematic Reviews* details why this checklist was adapted - **Moher D, Stewart L & Shekelle P:** Implementing PRISMA-P: recommendations for prospective authors. *Systematic Reviews* 2016 **5**:15

Continu/tonio	щ	Chaptelint item	Information	n reported	Line
Section/topic	#		Yes	No	number(s)
ADMINISTRATIVE INFO	RMAT	ION			
Title					
Identification	1a	Identify the report as a protocol of a systematic review			p.1 l.1-2
Update	1b	If the protocol is for an update of a previous systematic review, identify as such		\boxtimes	N/A
Registration	2	If registered, provide the name of the registry (e.g., PROSPERO) and registration number in the Abstract		\square	N/A
Authors					
Contact	3a	Provide name, institutional affiliation, and e-mail address of all protocol authors; provide physical mailing address of corresponding author	\square		p.1 l.3-25
Contributions	3b	Describe contributions of protocol authors and identify the guarantor of the review			p.12 l.16-18
Amendments	4	If the protocol represents an amendment of a previously completed or published protocol, identify as such and list changes; otherwise, state plan for documenting important protocol amendments		\square	N/A
Support					
Sources	5a	Indicate sources of financial or other support for the review			p.12 l.14-16
Sponsor	5b	Provide name for the review funder and/or sponsor	\square		p.12 l.14-16
Role of sponsor/funder	5c	Describe roles of funder(s), sponsor(s), and/or institution(s), if any, in developing the protocol			p.12 l.16
INTRODUCTION					
Rationale	6	Describe the rationale for the review in the context of what is already known			p.4-6
Objectives	7	Provide an explicit statement of the question(s) the review will address with reference to			p.6 l.12-27



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Castion/tonio			Informatior	n report <u>ed</u>	Line
Section/topic	#	Checklist item	Yes	No	number(s)
		participants, interventions, comparators, and outcomes (PICO)			
METHODS					•
Eligibility criteria	8	Specify the study characteristics (e.g., PICO, study design, setting, time frame) and report characteristics (e.g., years considered, language, publication status) to be used as criteria for eligibility for the review	\square		p.8 l.2-8
Information sources	9	Describe all intended information sources (e.g., electronic databases, contact with study authors, trial registers, or other grey literature sources) with planned dates of coverage	\square		p.7 l.19-20
Search strategy	10	Present draft of search strategy to be used for at least one electronic database, including planned limits, such that it could be repeated	\square		p.11
STUDY RECORDS					
Data management	11a	Describe the mechanism(s) that will be used to manage records and data throughout the review	\square		p.8 l.3-4; 20-24
Selection process	11b	State the process that will be used for selecting studies (e.g., two independent reviewers) through each phase of the review (i.e., screening, eligibility, and inclusion in meta-analysis)			p.8 l.9-16
Data collection process	11c	Describe planned method of extracting data from reports (e.g., piloting forms, done independently, in duplicate), any processes for obtaining and confirming data from investigators			p.8 l.19-25
Data items	12	List and define all variables for which data will be sought (e.g., PICO items, funding sources), any pre-planned data assumptions and simplifications			Table 2, p.8-9
Outcomes and prioritization	13	List and define all outcomes for which data will be sought, including prioritization of main and additional outcomes, with rationale		\square	N/A – scoping review
Risk of bias in individual studies	14	Describe anticipated methods for assessing risk of bias of individual studies, including whether this will be done at the outcome or study level, or both; state how this information will be used in data synthesis			p.9 l.3-6
DATA					
	15a	Describe criteria under which study data will be quantitatively synthesized	\square		Table 2, p.8-9
Synthesis	15b	If data are appropriate for quantitative synthesis, describe planned summary measures, methods of handling data, and methods of combining data from studies, including any planned exploration of consistency (e.g., <i>I</i> ² , Kendall's tau)		\square	N/A
	15c	Describe any proposed additional analyses (e.g., sensitivity or subgroup analyses, meta-regression)		\square	N/A
	15d	If quantitative synthesis is not appropriate, describe the type of summary planned			p.9 l.7-10, p.10 l.1-2
Meta-bias(es)	16	Specify any planned assessment of meta-bias(es) (e.g., publication bias across studies, selective		\square	N/A – scoping



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Section/topic	#	Checklist item	Informatio	n reported	Line
	"		Yes	No	number(s)
		reporting within studies)			review
Confidence in cumulative evidence	17	Describe how the strength of the body of evidence will be assessed (e.g., GRADE)		\square	N/A – scoping review

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BMJ Open

Policies to prevent zoonotic spillover: protocol for a systematic scoping review of evaluative evidence

Journal:	BMJ Open
Manuscript ID	bmjopen-2021-058437.R2
Article Type:	Protocol
Date Submitted by the Author:	07-Sep-2022
Complete List of Authors:	Clifford Astbury, Chloe; York University Lee, Kirsten M.; York University Aguiar, Raphael; York University, Dahdaleh Institute for Global Health Research Atique, Asma; York University Balolong, Marilen; University of the Philippines Manila Clarke, Janielle; York University Labonte, Ronald; University of Ottawa Ruckert, Arne; University of Ottawa Togño, Kathleen Chelsea; University of the Philippines Manila Viens, A.M.; York University, School of Global Health Wiktorowicz, M; York University Yau, Amy; London School of Hygiene & Tropical Medicine Penney, Tarra; York University
Primary Subject Heading :	Health policy
Secondary Subject Heading:	Global health, Health policy, Infectious diseases
Keywords:	Public health < INFECTIOUS DISEASES, Health policy < HEALTH SERVICES ADMINISTRATION & MANAGEMENT, PUBLIC HEALTH



Page 1 of 27

1 2		
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60		

1 Abstract

2 Introduction

The increasing incidence of pathogen transmission from animals to humans (zoonotic spillover events) has been attributed to behavioural practices and ecological and socioeconomic change. As these events sometimes involve pathogens with epidemic or pandemic potential, they pose a serious threat to population health. Public policies may play a key role in preventing these events. The aim of this review is to identify evaluations of public policies that target the determinants of zoonotic spillover, examining approaches taken to evaluation, choice of outcomes measures and evidence of effectiveness. Our approach to identifying and analysing this literature will be informed by a One Health lens, acknowledging the inter-connectedness of human, animal and environmental health.

12 Methods and analysis

A systematic scoping review methodology will be used. To identify articles, we will search Medline, SCOPUS, Web of Science and Global Health in March 2022 using search terms combining animal health and the animal-human interface, public policy, prevention and zoonoses. We will screen titles and abstracts and extract data according to published guidelines for scoping reviews. All evaluations of public policies aiming to prevent zoonotic spillover events will be eligible for inclusion. We will summarise key data from each study, mapping policies along the spillover pathway and outlining the range of policies, approaches to evaluation and outcome measures. Review findings will provide a useful reference for researchers and practitioners, outlining the state of the evaluative evidence around policies to prevent zoonotic spillover.

40
4123Ethics and dissemination

Formal ethical approval is not required, because the study does not involve primary data
collection. The findings of this study will be disseminated through a peer-reviewed publication,
presentations, and summaries for key stakeholders.

48 27 Strengths and limitations

- This scoping review protocol outlines the first piece of work to systematically identify and review evaluations of public policies designed to prevent zoonotic spillover, and will be undertaken in line with published guidelines for best practice in scoping reviews.
- The review will be informed by a One Health lens, encompassing distal determinants and risk factors for spillover events and acknowledging the interconnectedness of human, animal and environmental health.

1 2 3 4 5 6 7 8 9 10 11	1 • 2 3 4	Due to the complex drivers of spillover events, some potentially relevant policy evaluations may not be identified where outcome measures are too far removed from zoonotic spillover.
13 14 15 16 17 18 19 20 21 22 23 24 25		
26 27 28 29 30 31 32 33 34 35 36 37 38		
39 40 41 42 43 44 45 46 47 48 49 50		
51 52 53 54 55 56 57 58 59 60		

1 Introduction

The increasing incidence of zoonotic emerging infectious diseases (EIDs) has been attributed to behavioural practices and ecological and socioeconomic change, and is predicted to continue in the coming years (1). Higher levels of anthropogenic activity, including agricultural intensification, urbanisation and other forms of land use change, have led to increased interactions between wildlife, humans and livestock, increasing the risk of cross-species transmission (2,3). In response, a call has been issued by leading organisations and experts, including the United Nations Environment Programme, the International Livestock Research Institute and the Intergovernmental Science-Policy Platform on Biodiversity and Ecosystem Services, to complement reactive policy responses with policies that prevent zoonotic EIDs (1,4–7).

12 Preventing zoonotic spillover from a One Health perspective

Zoonotic spillover, defined as the transmission of a pathogen from an animal to a human, depends on the alignment of ecological, epidemiological and behavioural factors (8). Zoonotic pathogens must meet a series of conditions in order to induce spillover infections in humans, including appropriate density and distribution of reservoir hosts, pathogen prevalence, infection intensity and human exposure (8). Across this transmission pathway, a number of drivers of zoonotic spillover have been identified, including changes in wildlife and livestock populations (9); deforestation, urbanisation and other forms of land use change (10); and a variety of human practices including hunting, farming, animal husbandry, keeping of exotic pets and trade (6,7,11,12). These large-scale changes have on multiple occasions given rise to spillover events, sometimes involving pathogens with epidemic or pandemic potential.

A One Health perspective, which recognises the health of humans, animals and ecosystems as being closely linked and inter-dependent (13), can be useful in conceptualizing a range of potential determinants of spillover events. From this perspective, interventions could include surveillance of pools of viruses in wildlife and management of wildlife populations (14); enhanced food safety measures in both the wildlife and livestock value chain, pre- and post-farm gate (12,15–17); replacement of traditional 'wet' markets with supermarkets (18); controls on wildlife hunting, trade and consumption (11,19,20); and phasing out of unsustainable agriculture practices (6,21).

While some evaluative evidence exists around the effectiveness of interventions (22–25), they
 have often been implemented as short- to medium-term programmes or academic
 investigations (6). In some cases, zoonoses have re-emerged after successful programmes
 have ended (25). As a result, experts have argued for the incorporation of successful

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interventions into policy frameworks, providing interventions with the sustainability required for
 long-term disease control (6).

3 Governance, systems and the role of multi-sectoral actors

Public policy is 'a set of interrelated decisions taken by a political actor or group of actors concerning the selection of goals and the means of achieving them' (26). Public policy decisions are ultimately in the hands of government and supranational governing bodies, and have greater longevity compared to many programmes, which are often implemented for a fixed term. Non-government actors, including vested interest stakeholders, can also play a powerful role in shaping government decisions (27,28).

Although the longevity and scope of government actions may make policy an effective vehicle for prevention of emergent diseases, implementing policy is a complex process involving numerous stakeholders with competing views and interests (29). The responsibility for addressing zoonotic disease frequently spans multiple sectors of governance due to its relevance for both animals and humans. Where relevant policies are designed and implemented in isolation, opportunities for synergy may be missed and efforts may even be counter-productive.

Successful policy measures require not only a sound evidence base, but also governance structures that enable action to be taken. Given the range of possible risk factors that might contribute to emerging zoonoses, and the possible impacts of policies to prevent zoonotic spillover, a One Health response has been advocated, requiring coordination between institutions and government departments involved in human and animal health, trade, agriculture and the environment (30). At the international level, the World Health Organization, the Food and Agriculture Organization and the World Organisation for Animal Health have endorsed a One Health policy framework to respond to zoonotic infectious diseases, emphasising collaboration between agencies (31). Within countries, national and local governments have also emphasised the need for multi-sectoral efforts, although many report that further integration is still required (32).

Further, given the complex social-ecological systems within which policies to prevent zoonotic spillover are implemented, the risk of unintended consequences is high. For example, region-specific closures of live animal markets have been shown to spread pathogens further afield as vendors seek new venues to sell their animals (33). Meanwhile, attempts to manage populations of wild animals may alter pathogen dynamics, unintentionally increasing the risk of spillover into livestock or people (34).

Given these particular characteristics of policy development and implementation, they may be
 usefully considered as a particular case of intervention, and the evidence around them

1 assessed accordingly. Different types of interventions might be more or less feasibly 2 implemented by governments (or their partners), and their impacts might be different given 3 potentially more complex implementation contexts, longer timespans and broader geographic 4 ranges. Evaluations of these policies should also include consideration and monitoring of 5 potential unintended consequences. In order to facilitate this, multi-sectoral involvement in 6 both policy development and evaluation may be required.

7 Aims and scope

Approaches to managing epidemic and pandemic infectious pathogens once they have entered human populations have been systematically catalogued in the medical literature (35-41). These measures include hand washing, face masks, school closures, and contact tracing and case isolation. Further upstream, systematic reviews of interventions targeting the spillover pathway have predominantly focused on programmes rather than policies, and have been restricted by various characteristics such as geographic region (24) or pathogen type (25), or focused on programmes with an explicit endorsement of a One Health approach (23). In consequence, a comprehensive understanding of how policies to prevent zoonotic spillover have been evaluated, and what evidence there is of their effectiveness, is lacking. To address these research gaps, our objectives are to:

- 18
 1. Identify evaluations of policies that target the determinants of zoonotic spillover
 included in the spillover pathway (8) (i.e. human and animal health and interactions);
 - Identify insights around policy success and failure, and unintended consequences of
 policy implementation; and
 - Describe approaches to evaluation and key barriers and facilitators to evaluating policies to reduce the risk of zoonotic spillover.
 - Our approach to identifying and analysing this literature will be informed by a One Health lens,
 acknowledging the inter-connectedness of human, animal and environmental health.

2 3	1	
4	1	Methods and analysis
5 6	2	We will conduct a systematic scoping review of evaluations of policies aimed at preventing
7	3	zoonotic spillover events. The scoping review will be conducted in line with guidelines
8 9	4	published by Arksey and O'Malley and refined by Levac and colleagues (42-44), which
10	5	emphasise an iterative approach suited to an exploratory research question.
11 12	6	Stage 1: Identifying the research question
13	0	Stage 1. Identifying the research question
14 15	7	The aim of this review is to use a One Health lens to identify and describe the range of policies
16	8	that have been evaluated, the approaches to evaluation, and the evaluative evidence.
17 18	9	Informed by this aim, our research questions are:
19	10	1 What policies aimed at preventing zoonotic spillover have been evaluated?
20 21	11	a What are the types of policies?
22	17	a. Which policy actors (single department multi sectoral whole of government)
23 24	12	b. Which policy actors (single department, multi-sectoral, whole of government)
25	13	are engaged?
26 27	14	2. What are the reasons for policy success and failure, and the unintended
28	15	consequences of implementing these policies?
29 30	16	3. How has evaluation of these policies been approached in the literature?
31	17	a. What are the methods or study designs used?
32 33	18	b. What are the outcomes?
34	19	c. What are the barriers and facilitators to evaluation?
35 36	20	Stage 2: Identifying relevant studies
37	01	
38 39	21	We searched four electronic databases (Mediline, Scopus, Web of Science, Global Health) in
40	22	May 2021. The search strategy is organized by the main concepts in our research question:
41 42	23	the spillover pathway; public policy; prevention; and zoonotic pathogens. The search strategy
43	24	was developed iteratively, informed by existing systematic reviews focused on related
44 45	25	concepts (24,45–49) and known indicator papers meeting inclusion criteria. We also searched
46	26	the websites of 18 organisations involved in the prevention of zoonotic spillover to identify
47 48	27	relevant grey literature. See Supplementary File 1 for details of search strategy and websites
49	28	searched.
50 51	29	Stage 3: Study selection
52 53	30	Records identified through the searches will be collated and double screened using the online
54	21	netform Covidence (50). Studies will be included where they meet all of the following criteria:
55 56	51	plation condence (50). Studies will be included where they meet all of the following chiefla.
57	32	1. Primary empirical study from any country or region with English-language abstracts;
58 59	33	2. Report empirical findings from an evaluation of any sort; and
60		

- Focus on a policy implemented by government that targets a determinant of zoonotic spillover located on the spillover pathway (see Figure 1).
- 3 [Insert Figure 1]

4 Figure 1 Spillover pathway adapted from Plowright et al. (8,22)

Titles and abstracts will initially be screened, followed by full-text screening. Title and abstract screening of an initial set of 100 papers will be undertaken by two independent researchers. Results will be compared in order to ensure consistency in decisions around study eligibility, and discrepancies resolved through discussion of the inclusion criteria. This process will be repeated until an acceptable level of agreement (>90%) is reached. The remaining papers will then be screened by one of the two reviewers. Full-text screening will be undertaken by two independent researchers and discrepancies will be resolved by discussing reasons for inclusion or exclusions among the screeners. Studies with full-texts in languages other than English will be eligible for inclusion if they include an English-language abstract. Full-text studies published in French, Spanish or Chinese will be single-screened by a member of the research team fluent in that language. Studies published in other languages will be translated as necessary.

- In line with published guidelines, the approach to study selection may be refined iteratively
 when reviewing articles for inclusion (42–44). Reporting on the search and screening process
 will follow the guidelines provided in the Preferred Reporting Items for Systematic Reviews
 and Meta-Analyses Extension for Scoping Reviews (51).
- ³⁶₃₇ 21 Stage 4: Charting the data

Data charting will be conducted using a data charting form designed to identify the information required to answer the research question and sub-research questions (see Supplementary File 2). Data charting focused on characteristics of the study, the policy and the evaluation. For each policy, this included identifying which determinant of zoonotic spillover situated along the spillover pathway was being targeted. For the purpose of this study, we used a model of the spillover pathway adapted from Plowright et al.'s work (8,22), in which we differentiated between wildlife and domesticated animals (Figure 1). This differentiation is important in the policy context, as the wildlife-domesticated animal interface is an important site for intervention, as well as the human-animal interface.

As recommended, the data charting form will be piloted with ten records to ensure that it is
 consistent with the research question, and the data charting form will be revised iteratively in
 order to ensure the purpose of the research is being met (42–44).

- ⁵⁸₅₉ 34 Stage 5: Collating, summarising and reporting the results

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1 We will undertake quality assessment of the included studies using the Quality Assessment

Tool for Quantitative Studies developed by the Effective Public Health Practice Project (52),
which has previously been used to assess the quality of natural experiments including public
policy evaluations (53).

We will analyse the extracted data, presenting a numerical summary of the included studies in table form, allowing us to describe the range of policy interventions that have been evaluated, approaches to evaluation, and evidence of effectiveness. We will also conduct a thematic analysis of the contents of the included articles in order to identify, if possible, barriers and facilitators to implementing and evaluating these policies, as well as insights into why policies succeeded or failed in achieving their aims.

20 11 Patient and public involvement 21

This scoping review is being undertaken as part of a larger project involving policy actors at national and international levels as research team members, knowledge users and participants. Insights from the project have informed protocol development and stakeholders are able to provide input and perspectives on the results of the review. Project-level dissemination events involving policy stakeholders are also planned, where findings from the proposed review will be shared.

32 18 Strengths and weaknesses of the study 33

To our knowledge, this is the first attempt to systematically identify and document evaluations of policies aiming to prevent the spillover of zoonoses into human populations. However, because of the complex drivers of spillover events, some potentially relevant policy evaluations may be excluded where their outcome measures are too far removed from zoonotic spillover. For example, it has been hypothesised that declines in vulture populations may increase the risk of pathogen transmission by increasing the number of uneaten carcasses, as well as, potentially, the population of feral dogs (54). In 2006, India, Pakistan and Nepal implemented a ban on the veterinary drug diclofenac, which had been identified as a driver of declining vulture populations. While policy evaluations suggest that this ban has resulted in a resurgence of vultures (55–58), the knock-on effects of this on zoonotic pathogen transmission risk have not been included in these evaluations. While relevant, such evaluations will be difficult to systematically identify as they make no reference to zoonotic disease.

In addition, this review will focus on policy evaluations that have been reported in the peer reviewed and grey literature. Policies that have been implemented but not evaluated, or
 evaluated but not reported in the literature, will therefore be excluded from this review. As a
 result, potentially effective and important policies in the prevention of zoonotic spillover events

may not be identified. However, we hope that the findings from this review will highlight these gaps in the evaluative evidence. We also hope that this review, by extracting practical dimensions such as study design, outcome measures and the challenges encountered in the evaluation process, will support policymakers and researchers in carrying out policy sevaluations in this space.

6 Ethics and dissemination

Formal ethical approval is not required, because the study does not involve primary data
collection. The findings of this study will be disseminated through a peer-reviewed publication,
presentations, and summaries for key stakeholders.

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List of abbreviations

Declarations

Not applicable.

Not applicable.

EID: Emerging infectious disease

Patient consent for publication

Availability of data and material

during the current study.

developing the protocol.

Competing interests

Funding

Contributors

manuscript.

Acknowledgements

Ethics approval and consent to participate

1 2

Data sharing not applicable to this article as no datasets were generated or analysed

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CCA, KML and TLP conceived and designed the study. CCA prepared the

manuscript. KML, TLP, RA, AA, MB, JC, RL, AR, KCT, AMV, MW and AY provided

critical input on the manuscript and methods and have read and approved the final

The authors declare that they have no competing interests.

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Figure 1 Spillover pathway adapted from Plowright et al. (8,22)

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Supplementary file 1

Example search string framed by core concepts

Example from Scopus search, all terms in TITLE-ABS-KEY; Block 1 AND 2 AND 3 AND 4

Block 1 - Spillover pathway (Animal populations and human-animal interface)

Spillover OR "spill over" OR "cross-species transmission" OR poultry OR wildlife OR bushmeat OR "bush meat" OR livestock OR "animal market" OR "wet market" OR "bird market" OR horse* OR waterfowl OR fowl OR bat OR bats OR mammal* OR swine OR pig* OR poaching OR "pet trade" OR pork OR "trade W/5 animal"

Block 2 – Public policy

policy OR law OR legal OR legislat* OR regulat* OR tariff OR subsidy OR tax OR ban OR "voluntary agreement" OR incentive OR fiscal OR guidelines OR govern* OR federal* OR closure OR closing OR state* OR "rest day*" OR "border control*" OR "habitat protection" OR "wetland protection" OR "supplement* fed" OR "supplement* feed*" OR "market size"

Block 3 – Prevention

Prevent* OR "ecological intervention*" OR "non-pharmaceutical intervention*" OR "public health" OR "risk management" OR "risk minimisation" OR "control strateg*" OR "outbreak risk" OR "reduc* W/5 transmission" OR "reduc* W/5 infection"

Block 4 - Zoonotic pathogens

Zika OR ebola OR covid-19 OR sars-cov-2 OR coronavirus OR sars OR mers OR h1n1 OR h7n9 OR h5n1 OR "one health" OR dengue OR "nipah virus" OR influenza OR zoonoses OR zoonosis OR zoonotic OR "West Nile" OR "HIV/AIDS" OR "avian flu" OR "hendra virus" OR "marburg virus" OR "yellow fever" OR "tick-borne encephalitis" OR "emerging infectious diseases" OR "emergent infectious diseases" OR brucellosis OR rabies OR chikungunya OR "bovine spongiform encephalopathy" OR "rift valley fever"

Search strings for all included academic databases

Scopus search

TITLE-ABS-KEY(Zika OR ebola OR covid-19 OR sars-cov-2 OR coronavirus OR sars OR mers OR h1n1 OR h7n9 OR h5n1 OR "one health" OR dengue OR "nipah virus" OR influenza OR zoonoses OR zoonosis OR zoonotic OR "West Nile" OR "HIV/AIDS" OR "avian flu" OR "hendra virus" OR "marburg virus" OR "yellow fever" OR "tick-borne encephalitis" OR "emerging

infectious diseases" OR "emergent infectious diseases" OR brucellosis OR rabies OR chikungunya OR "bovine spongiform encephalopathy" OR "rift valley fever") AND TITLE-ABS-KEY(Spillover OR "spill over" OR "cross-species transmission" OR poultry OR wildlife OR bushmeat OR "bush meat" OR livestock OR "animal market*" OR "wet market*" OR "bird market*" OR horse* OR waterfowl OR fowl OR bat OR bats OR mammal* OR swine OR pig* OR poaching OR "pet trade" OR pork OR "trade W/5 animal") AND TITLE-ABS-KEY(policy OR law OR legal OR legislat* OR regulat* OR tariff OR subsidy OR tax OR ban OR "voluntary agreement" OR incentive OR fiscal OR guidelines OR govern* OR federal* OR closure OR closing OR state* OR "rest day*" OR "border control*" OR "habitat protection" OR "wetland protection" OR "supplement* fed" OR "supplement* feed*" OR "market size") AND TITLE-ABS-KEY(Prevent* OR "ecological intervention*" OR "non-pharmaceutical intervention*" OR "public health" OR "risk management" OR "risk minimisation" OR "control strateg*" OR "outbreak risk" OR "reduc* W/5 transmission" OR "reduc* W/5 infection")

Medline search

(((Zika[Title/Abstract] OR ebola[Title/Abstract] OR covid-19[Title/Abstract] OR sars-cov-2[Title/Abstract] OR coronavirus[Title/Abstract] OR sars[Title/Abstract] OR mers[Title/Abstract] OR h1n1[Title/Abstract] OR h7n9[Title/Abstract] OR h5n1[Title/Abstract] OR "one health"[Title/Abstract] OR dengue[Title/Abstract] OR "nipah virus"[Title/Abstract] OR influenza[Title/Abstract] OR zoonoses[Title/Abstract] OR zoonosis[Title/Abstract] OR zoonotic[Title/Abstract] OR "West Nile"[Title/Abstract] OR "HIV/AIDS"[Title/Abstract] OR "avian flu"[Title/Abstract] OR "hendra virus"[Title/Abstract] OR "marburg virus"[Title/Abstract] OR "yellow fever"[Title/Abstract] OR "tick-borne encephalitis"[Title/Abstract] OR "emerging infectious diseases"[Title/Abstract] OR "emergent infectious diseases"[Title/Abstract] OR brucellosis[Title/Abstract] OR rabies[Title/Abstract] OR chikungunya[Title/Abstract] OR "bovine spongiform encephalopathy"[Title/Abstract] OR "rift valley fever"[Title/Abstract] OR zoonoses [mesh]) AND (Spillover[Title/Abstract] OR "spill over"[Title/Abstract] OR "cross-species transmission"[Title/Abstract] OR poultry[Title/Abstract] OR wildlife[Title/Abstract] OR bushmeat[Title/Abstract] OR "bush meat"[Title/Abstract] OR livestock[Title/Abstract] OR "animal market"[Title/Abstract] OR "animal markets"[Title/Abstract] OR "wet market"[Title/Abstract] OR "wet markets"[Title/Abstract] OR "bird market"[Title/Abstract] OR "bird markets"[Title/Abstract] OR horse[Title/Abstract] OR horses[Title/Abstract] OR waterfowl[Title/Abstract] OR fowl[Title/Abstract] OR bat[Title/Abstract] OR bats[Title/Abstract] OR mammal[Title/Abstract] OR

mammals[Title/Abstract] OR mammalian[Title/Abstract] OR swine[Title/Abstract] OR pig[Title/Abstract] OR pigs[Title/Abstract] OR poaching[Title/Abstract] OR "pet trade"[Title/Abstract] OR pork[Title/Abstract] OR animal N5 trade[Title/Abstract] OR disease reservoir [mesh])) AND (policy[Title/Abstract] OR law[Title/Abstract] OR legal[Title/Abstract] OR legislation[Title/Abstract] OR legislative[Title/Abstract] OR legislating[Title/Abstract] OR regulation[Title/Abstract] OR regulations[Title/Abstract] OR regulatory[Title/Abstract] OR tariff[Title/Abstract] OR subsidy[Title/Abstract] OR tax[Title/Abstract] OR ban[Title/Abstract] OR "voluntary agreement"[Title/Abstract] OR incentive[Title/Abstract] OR fiscal[Title/Abstract] OR guidelines[Title/Abstract] OR government[Title/Abstract] OR governments[Title/Abstract] OR federal[Title/Abstract] OR federally[Title/Abstract] OR closure[Title/Abstract] OR closing[Title/Abstract] OR state[Title/Abstract] OR "rest day"[Title/Abstract] OR "rest days"[Title/Abstract] OR "border control"[Title/Abstract] OR "border controls"[Title/Abstract] OR "habitat protection"[Title/Abstract] OR "wetland protection"[Title/Abstract] OR "supplemental feeding"[Title/Abstract] OR "market size"[Title/Abstract])) AND (Prevent[Title/Abstract] OR prevention[Title/Abstract] OR "ecological intervention"[Title/Abstract] OR "ecological interventions"[Title/Abstract] OR "non-pharmaceutical intervention"[Title/Abstract] OR "nonpharmaceutical interventions"[Title/Abstract] OR "public health"[Title/Abstract] OR "risk management"[Title/Abstract] OR "risk minimisation"[Title/Abstract] OR "control strategy"[Title/Abstract] OR "control strategies"[Title/Abstract] OR "outbreak risk"[Title/Abstract] OR reducing N5 transmission[Title/Abstract] OR reducing N5 infection[Title/Abstract])

Web of knowledge search

AB=(Zika OR ebola OR covid-19 OR sars-cov-2 OR coronavirus OR sars OR mers OR h1n1 OR h7n9 OR h5n1 OR "one health" OR dengue OR "nipah virus" OR influenza OR zoonoses OR zoonosis OR zoonotic OR "West Nile" OR "HIV/AIDS" OR "avian flu" OR "hendra virus" OR "marburg virus" OR "yellow fever" OR "tick-borne encephalitis" OR "emerging infectious diseases" OR "emergent infectious diseases" OR brucellosis OR rabies OR chikungunya OR "bovine spongiform encephalopathy" OR "rift valley fever")

AND

AB=(Spillover OR "spill over" OR "cross-species transmission" OR poultry OR wildlife OR bushmeat OR "bush meat" OR livestock OR "animal market*" OR "wet market*" OR "bird
market*" OR horse* OR waterfowl OR fowl OR bat OR bats OR mammal* OR swine OR pig* OR poaching OR "pet trade" OR pork OR trade NEAR animal)

AND

AB=(policy OR law OR legal OR legislat* OR regulat* OR tariff OR subsidy OR tax OR ban OR "voluntary agreement" OR incentive OR fiscal OR guidelines OR govern* OR federal* OR closure OR closing OR state* OR "rest day*" OR "border control*" OR "habitat protection" OR "wetland protection" OR "supplement* fed" OR "supplement* feed*" OR "market size")

AND

AB=(Prevent* OR "ecological intervention*" OR "non-pharmaceutical intervention*" OR "public health" OR "risk management" OR "risk minimisation" OR "control strateg*" OR "outbreak risk" OR reduc* NEAR transmission OR reduc* NEAR infection)

Ovid Global Health database search (all in abstract)

Zika OR ebola OR covid-19 OR sars-cov-2 OR coronavirus OR sars OR mers OR h1n1 OR h7n9 OR h5n1 OR one health OR dengue OR nipah virus OR influenza OR zoonoses OR zoonosis OR zoonotic OR West Nile OR HIV/AIDS OR avian flu OR hendra virus OR marburg virus OR yellow fever OR tick-borne encephalitis OR emerging infectious diseases OR emergent infectious diseases OR brucellosis OR rabies OR chikungunya OR bovine spongiform encephalopathy OR rift valley fever

AND

Spillover OR spill over OR cross-species transmission OR poultry OR wildlife OR bushmeat OR bush meat OR livestock OR animal market* OR wet market* OR bird market* OR horse* OR waterfowl OR fowl OR bat OR bats OR mammal* OR swine OR pig* OR poaching OR pet trade OR pork OR (trade adj5 animal)

AND

policy OR law OR legal OR legislat* OR regulat* OR tariff OR subsidy OR tax OR ban OR voluntary agreement OR incentive OR fiscal OR guidelines OR govern* OR federal* OR closure OR closing OR state* OR rest day* OR border control* OR habitat protection OR wetland protection OR supplement* fed OR supplement* feed* OR market size

AND

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4	Prevent" OR ecological intervention" OR non-pharmaceutical intervention" OR public health OR
5	risk management OR risk minimisation OR control strateg* OR outbreak risk OR (reduc* adj5
6 7	transmission) OR (reduc* adj5 infection)
8 9	List of organization websites searched for grey literature
10 11	1. World Organization for Animal Health (formerly OIE)
12	2. Food and Agriculture Organization
13 14	3. World Health Organization
15	4. Wildlife Disease Association
17	5. International Alliance against Health Risks in WildlifeTrade
18 10	6. United Nations Environment Program
20	7. United Nations Office for Drugs and Crime
21 22	8. Global Alliance for Rabies Control
22	9. EcoHealth Alliance
24 25	10 Network for EcoHealth and One Health
25 26	44. Jackson a line at a line and black that
27	11. International Livestock Research Institute
28	12. Preventing Pandemics at the Source
29 30	13. World Veterinary Association
31	14. CITES
32 33	15. TRAFFIC
34	16. One Health Commission
35	17. Martel Wildlife Fund
36 37	
38	18. World Trade Organization
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Supplementary file 2

Data charting form

Record	Author(s)				
	Year				
Policy	Country				
	World region (World Bank grouping) (44)				
	Country income (World Bank grouping) (44)				
	Disease				
	Stakeholders or sector responsible for implementing policy (retail,				
	agriculture, conservation, etc.)				
	Implementation date (start date, or range if the policy has been changed)				
	Intervention type				
	Location along spillover pathway adapted from Plowright et al. (7,21)				
	Policy level (local, national, regional, global)				
	Multi-sectoral initiative (Y/N)				
	Sector(s) responsible for policy				
Evaluation	Aim				
	Type (Process/outcome)				
	Study design				
	Theoretical framework and/or logic model underpinning evaluation (if				
	described)				
	Period of observation				
	Outcome measure(s) and change in measure(s)				
	Consideration of unintended consequences (Y/N)				
	If yes, which unintended consequences? (e.g., economic outcomes, food security)				

PRISMA-P 2015 Checklist

This checklist has been adapted for use with systematic review protocol submissions to BioMed Central journals from Table 3 in Moher D et al: Preferred reporting items for systematic review and meta-analysis protocols (PRISMA-P) 2015 statement. *Systematic Reviews* 2015 **4**:1

An Editorial from the Editors-in-Chief of *Systematic Reviews* details why this checklist was adapted - **Moher D, Stewart L & Shekelle P**: Implementing PRISMA-P: recommendations for prospective authors. *Systematic Reviews* 2016 **5**:15

Saation/tonio	#	Checklist item	Information reported		Line		
Section/topic			Yes	No	number(s)		
ADMINISTRATIVE INFORMATION							
Title							
Identification	1a	Identify the report as a protocol of a systematic review	\square		p.1 l.1-2		
Update	1b	If the protocol is for an update of a previous systematic review, identify as such		\boxtimes	N/A		
Registration	2	If registered, provide the name of the registry (e.g., PROSPERO) and registration number in the Abstract		\square	N/A		
Authors							
Contact	3а	Provide name, institutional affiliation, and e-mail address of all protocol authors; provide physical mailing address of corresponding author	\square		p.1 l.3-25		
Contributions	3b	Describe contributions of protocol authors and identify the guarantor of the review			p.12 l.16-18		
Amendments	4	If the protocol represents an amendment of a previously completed or published protocol, identify as such and list changes; otherwise, state plan for documenting important protocol amendments		\square	N/A		
Support							
Sources	5a	Indicate sources of financial or other support for the review	\square		p.12 l.14-16		
Sponsor	5b	Provide name for the review funder and/or sponsor			p.12 l.14-16		
Role of sponsor/funder	5c	Describe roles of funder(s), sponsor(s), and/or institution(s), if any, in developing the protocol	\square		p.12 l.16		
INTRODUCTION							
Rationale	6	Describe the rationale for the review in the context of what is already known	\square		p.4-6		
Objectives	7	Provide an explicit statement of the question(s) the review will address with reference to			p.6 l.12-27		



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		Checklist item	Information reported		Line	
Section/topic	#		Yes	No	number(s)	
		participants, interventions, comparators, and outcomes (PICO)				
METHODS						
Eligibility criteria	8	Specify the study characteristics (e.g., PICO, study design, setting, time frame) and report characteristics (e.g., years considered, language, publication status) to be used as criteria for eligibility for the review			p.8 l.2-8	
Information sources	9	Describe all intended information sources (e.g., electronic databases, contact with study authors, trial registers, or other grey literature sources) with planned dates of coverage	\boxtimes		p.7 l.19-20	
Search strategy	10	Present draft of search strategy to be used for at least one electronic database, including planned limits, such that it could be repeated	\boxtimes		p.11	
STUDY RECORDS					-	
Data management	11a	Describe the mechanism(s) that will be used to manage records and data throughout the review	\boxtimes		p.8 l.3-4; 20-24	
Selection process	11b	State the process that will be used for selecting studies (e.g., two independent reviewers) through each phase of the review (i.e., screening, eligibility, and inclusion in meta-analysis)	\square		p.8 l.9-16	
Data collection process	11c	Describe planned method of extracting data from reports (e.g., piloting forms, done independently, in duplicate), any processes for obtaining and confirming data from investigators	\boxtimes		p.8 l.19-25	
Data items	12	List and define all variables for which data will be sought (e.g., PICO items, funding sources), any pre-planned data assumptions and simplifications	\boxtimes		Table 2, p.8-9	
Outcomes and prioritization	13	List and define all outcomes for which data will be sought, including prioritization of main and additional outcomes, with rationale		\square	N/A – scoping review	
Risk of bias in individual studies	14	Describe anticipated methods for assessing risk of bias of individual studies, including whether this will be done at the outcome or study level, or both; state how this information will be used in data synthesis	\boxtimes		p.9 l.3-6	
DATA	_					
	15a	Describe criteria under which study data will be quantitatively synthesized	\boxtimes		Table 2, p.8-9	
Synthesis	15b	If data are appropriate for quantitative synthesis, describe planned summary measures, methods of handling data, and methods of combining data from studies, including any planned exploration of consistency (e.g., <i>I</i> ² , Kendall's tau)		\boxtimes	N/A	
-	15c	Describe any proposed additional analyses (e.g., sensitivity or subgroup analyses, meta-regression)		\boxtimes	N/A	
	15d	If quantitative synthesis is not appropriate, describe the type of summary planned			p.9 l.7-10, p.10 l.1-2	
Meta-bias(es)	16	Specify any planned assessment of meta-bias(es) (e.g., publication bias across studies, selective		\square	N/A – scoping	



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Saction/tonio	#	Checklist item	Informatio	Line	
Section/topic	#		Yes	No	number(s)
		reporting within studies)			review
Confidence in cumulative evidence	17	Describe how the strength of the body of evidence will be assessed (e.g., GRADE)		\square	N/A – scoping review

For peer review only

