nature portfolio

Corresponding author(s):	Kramer, SC
Last updated by author(s):	May 10, 2024

Reporting Summary

Nature Portfolio wishes to improve the reproducibility of the work that we publish. This form provides structure for consistency and transparency in reporting. For further information on Nature Portfolio policies, see our Editorial Policies and the Editorial Policy Checklist.

For all statistical analyses, confirm that the following items are present in the figure legend, table legend, main text, or Methods section.

~ .			
SΤ	at:	121	ICS

n/a	Confirmed
	$oxed{x}$ The exact sample size (n) for each experimental group/condition, given as a discrete number and unit of measurement
x	A statement on whether measurements were taken from distinct samples or whether the same sample was measured repeatedly
	The statistical test(s) used AND whether they are one- or two-sided Only common tests should be described solely by name; describe more complex techniques in the Methods section.
X	A description of all covariates tested
×	A description of any assumptions or corrections, such as tests of normality and adjustment for multiple comparisons
	A full description of the statistical parameters including central tendency (e.g. means) or other basic estimates (e.g. regression coefficient) AND variation (e.g. standard deviation) or associated estimates of uncertainty (e.g. confidence intervals)
	For null hypothesis testing, the test statistic (e.g. <i>F</i> , <i>t</i> , <i>r</i>) with confidence intervals, effect sizes, degrees of freedom and <i>P</i> value noted <i>Give P values as exact values whenever suitable.</i>
x	For Bayesian analysis, information on the choice of priors and Markov chain Monte Carlo settings
x	For hierarchical and complex designs, identification of the appropriate level for tests and full reporting of outcomes
	$oxed{x}$ Estimates of effect sizes (e.g. Cohen's d , Pearson's r), indicating how they were calculated
	Our web collection on <u>statistics for biologists</u> contains articles on many of the points above.

Software and code

Policy information about availability of computer code

Data collection [

NA

Data analysis

All analyses were conducted in R version 4.2.3. The model was coded and run using the package pomp (version 5.4), and fit using the sbplx algorithm via the package nloptr (version 2.0.3). All code can be found at: https://github.com/sarahckramer/resp_virus_interactions

For manuscripts utilizing custom algorithms or software that are central to the research but not yet described in published literature, software must be made available to editors and reviewers. We strongly encourage code deposition in a community repository (e.g. GitHub). See the Nature Portfolio guidelines for submitting code & software for further information.

Data

Policy information about availability of data

All manuscripts must include a data availability statement. This statement should provide the following information, where applicable:

- Accession codes, unique identifiers, or web links for publicly available datasets
- A description of any restrictions on data availability
- For clinical datasets or third party data, please ensure that the statement adheres to our <u>policy</u>

Links to all publicly-available data used in this study can be found, along with the code used for the analyses, at: https://github.com/sarahckramer/resp_virus_interactions

Research involving human participants, their data, or biological material

-	race, ethnicity and racism.		
Reporting on sex and gende	NA		
Reporting on race, ethnicity other socially relevant group	NA		
Population characteristics	NA		
Recruitment	NA		
Ethics oversight	NA		
Note that full information on t	the approval of the study protocol must also be provided in the manuscript.		
Field-specific	c renorting		
•	w that is the best fit for your research. If you are not sure, read the appropriate sections before making your selection.		
Life sciences			
	Behavioural & social sciences		
or a reference copy or the accum-	internal and sections, see internal sections, see internal sections and sections and sections are sections.		
Ecological, e	volutionary & environmental sciences study design		
	n these points even when the disclosure is negative.		
Study description	We fit mathematical models of influenza and respiratory syncytial virus cocirculation to observed data from Hong Kong and Canada, norder to estimate the strength and duration of the interaction effect between the two viruses. There were no covariates to control or in this study.		
Research sample	olicly-available, weekly data on 1) the total number of tests performed for influenza and RSV, 2) the number of tests positive for uenza or RSV, and 3) the proportion of all-cause consultations due to influenza-like illness were obtained from the Centre for alth Protection of the Government of the Hong Kong Special Administrative Region and the Public Health Agency of Canada. These as were chosen for their availability, and because both datasets contain several years of data on both viruses.		
Sampling strategy	As previously-collected data were used for this study, we did not conduct any sampling.		
Data collection	ients in public hospitals (Hong Kong), or in either hospitals or emergency departments (Canada) were tested for influenza and/or / if diagnostic information could potentially aid treatment. The number of patients with influenza-like illness were reported by epatient clinics (Hong Kong) and primary care providers (Canada).		
Timing and spatial scale	evant data in Hong Kong cover the period from January 2014 through mid-February 2020; data in Canada cover from September 10 through August 2014. In both locations, data were available at a weekly time scale, and were collected from patients from oughout the country.		
Data exclusions	id not consider data collected during the 2009 influenza pandemic or during the recent pandemic of SARS-CoV-2. This is use typical circulation patterns of both influenza and RSV were strongly affected by both pandemics.		
Reproducibility	code to repeat the analyses has been made publicly available.		
Randomization	dels were fit to the data described above; no randomization was conducted.		
Blinding	s is a modeling study, in which a model was fit to data in order to estimate several key parameters of interest; blinding is not evant to this study design.		
Did the study involve fiel	d work? Yes X No		

Reporting for specific materials, systems and methods

We require information from authors about some types of materials, experimental systems and methods used in many studies. Here, indicate whether each material, system or method listed is relevant to your study. If you are not sure if a list item applies to your research, read the appropriate section before selecting a response.

Materials & experime	ntal systems N	Methods		
n/a Involved in the study		/a Involved in the study		
Antibodies	[▼ ChIP-seq		
x Eukaryotic cell lines	[Flow cytometry		
Palaeontology and a	ırchaeology	MRI-based neuroimaging		
Animals and other o	organisms			
Clinical data	Clinical data			
Dual use research of concern				
Plants				
'				
Plants				
FIGITIS				
Seed stocks	NA			
Novel plant genotypes	NA			
Authentication	NA			