nature portfolio

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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 <u>Editorial Policies</u> and the <u>Editorial Policy Checklist</u>.

Statistics

For all statistical analyses, confirm that the following items are present in the figure legend, table legend, main text, or Methods section.					
n/a	Cor	Confirmed			
	\square	The exact sample size (n) for each experimental group/condition, given as a discrete number and unit of measurement			
\boxtimes		A statement on whether measurements were taken from distinct samples or whether the same sample was measured repeatedly			
\boxtimes		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.			
\boxtimes		A description of all covariates tested			
\boxtimes		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)			
\boxtimes		For null hypothesis testing, the test statistic (e.g. F, t, r) with confidence intervals, effect sizes, degrees of freedom and P value noted Give P values as exact values whenever suitable.			
\boxtimes		For Bayesian analysis, information on the choice of priors and Markov chain Monte Carlo settings			
\boxtimes		For hierarchical and complex designs, identification of the appropriate level for tests and full reporting of outcomes			
\boxtimes		Estimates of effect sizes (e.g. Cohen's d, Pearson's r), indicating how they were calculated			
		Our web collection on statistics for biologists contains articles on many of the points above.			

Software and code

Policy information about availability of computer code

 Data collection
 The synthetic population used as input for the computational model has been generated from open source data (available from insee.fr) as described in a previously published work (the corresponding citation is reported in the paper).

 Data analysis
 The code used for the simulations is written in C++11 compiled with g++ 5.2.0. It is available on zenodo at the link https://zenodo.org/

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 <u>data availability statement</u>. 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 policy

The synthetic population used in the analysis are available on zenodo at the link https://zenodo.org/record/5910314#.Yf1b-vXMI-T

Field-specific reporting

Please select the one below 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

Ecological, evolutionary & environmental sciences For a reference copy of the document with all sections, see <u>nature.com/documents/nr-reporting-summary-flat.pdf</u>

Life sciences study design

All studies must disclose on these points even when the disclosure is negative.

Sample size	Analyses in Figure 2 and 3 of the main paper were obtained running 2000 stochastic simulations. This number was sufficiently large to lead to stable statistical estimates. Figure 4 was obtained by running 8000 stochastic simulations. In the parameter regime of Figure 4, the epidemic is close to extinction, thus stochastic fluctuations play an important role. For this reason an increased number of simulations was necessary.
Data exclusions	We only considered data for a mid-size french city and excluded census data not concerning the city. No data was excluded from the simulation results to compute the summary measurements in Tables and Figures.
Replication	Simulation of each scenario was executed independently and with a different seed of the random number generator. In Fig 2 & 3, each data point was based on 2000 replications. In Fig 4, each data point was based on 8000 replications. All simulations yielded data that could be used in the analysis.
Randomization	We used the minimum standard congruential random generator (rminstd_rand0) as implemented in libstdc++ 5.2.0 to simulate random numbers in stochastic simulations. No randomization was necessary in the investigation as we undertook a systematic numerical exploration of hypothetical scenarios.
Blinding	No blinding was necessary as the results of the simulations could not be affected by the investigator's knowledge of the scenarios under study.

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 & experimental systems

Methods

n/a	Involved in the study
\boxtimes	Antibodies
\boxtimes	Eukaryotic cell lines
\boxtimes	Palaeontology and archaeology
\boxtimes	Animals and other organisms
\boxtimes	Human research participants
\boxtimes	Clinical data

Dual use research of concern

- Involved in the study n/a
- \boxtimes ChIP-seq
- \boxtimes Flow cytometry
- MRI-based neuroimaging \boxtimes