# Epidemiological Data Source Assessment

This tool helps practitioners rate different data sources on things like quality, timeliness and usefulness for modelling SARS-CoV-2. By pooling expert opinions, we can create visual comparisons showing the trade-offs between different data types. This makes it easier to decide which data to use when estimating transmissibility, and helps identify when different sources might conflict with each other.

Please think about a candidate dataset that you might consider for modelling SARS-CoV-2 and answer the following questions which are grouped into six main categories: basic meta-data, scope, resolution, data quality, data utility, and practical considerations with each then having further subcategories. For each of these subcategories you will assign a value between 0 and 5 (for numeric entries), assign a category, or enter free text.

1. Email *	
Basic meta-data	
Dasic meta-data	
What kind of data is it and why was it collected? The essentials about the data sour	ce.
2. Source type *	
The category/classification of data	
Mark only one oval.	
Confirmed cases time series	
Hospitalisation time series	
Wastewater	
Contact tracing	
Other:	

3.	Study design *
	Type of study
	Mark only one oval.
	Sentinel surveillance
	Cohort study
	Routine surveillance
	Contact survey
	Other:
1	Description *
4.	Description *  Brief explanation of what the data contains
	bher explanation of what the data contains
5.	Primary purpose *
	Original intent of data collection
	Mark only one oval.
	Clinical management
	Other:
S	cope
	ho's included in the data and who isn't? Covers the population represented, any ubgroups, and how close it is to actual infection timing.
6.	The source population *
	Overall population from which the data are drawn, eg hospital catchment areas

7.	Target population *
	Who does the data aim to cover
	Tick all that apply.
	Specific age groups
	Risk-specific
	Convenience
	Geographic structure
	Healthcare workers
	Whole population
	Other:
8.	Stratification/covariates (except spatial-temporal see Resolution section) *
	Tick all that apply.
	Demographic
	Clinical
	None
	Other:
9.	Collection type *
	Mark only one oval.
	Routine
	Triggered
	One-off
	One-on
10.	If triggered, potential triggers
	Mark only one oval.
	Greater than a threshold
	New variant/pathogen detected
	Other:

#### Resolution

How detailed is the data? Looks at whether it's individual or aggregated, and how fine-grained the time and location information is.

11.	Data aggregation *
	Mark only one oval.
	Individual
	Aggregated
10	Temporal data *
12.	Does your data have a temporal component
	Mark only one oval.
	Yes Skip to question 13
	No Skip to question 16
Re	solution - temporal data
Que	estions related to temporal data
13.	Collection frequency
	How often data are collected
	Mark only one oval.
	Continuously
	Daily
	Multiple times a week
	Weekly
	Fortnightly
	Monthly
	Less frequently than monthly

14.	Reporting frequency
	How often releases/updates occur
	Mark only one oval.
	Continuously
	Daily
	Multiple times a week
	Weekly
	Fortnightly
	Monthly
	Less frequently than monthly
15.	Time period covered
	Tick all that apply.
	Early outbreak Peak Endemic Continuous
	Other:
Res	solution - spatial data
Que	estions related to spatial data
16.	Spatial data *
	Mark only one oval.
	Yes
	No Skip to section 7 (Data quality)

17.	Spatial resolution
	Lowest spatial resolution across geographical areas
	Mark only one oval.
	International
	National
	Regional
	Local
	Hyper-local
18.	Geographic coverage
	Completeness of target geographical areas
	Mark only one oval.
	1 2 3 4 5
	Very Very high coverage
Da	ata quality
	w trustworthy is it? Rates things like measurement accuracy, potential biases, and porting delays.
Da	ata quality - measurement quality
10	
19.	Quality of case definitions
	How standardised is the case definition? For example does it vary across time or space or both. Leave blank if not applicable.
	Mark only one oval.
	1 2 3 4 5
	Very Very variable
	.,

#### 20. Test sensitivity

Sensitivity of any test used to obtain information on infection. Leave blank if not applicable.

Mark only one oval.



#### 21. Test specificity

Specificity of any test used to obtain information on infection. Leave blank if not applicable.

Mark only one oval.



### 22. Potential for unexplained variability \*

Mark only one oval.



#### 23. Reporting delay \*

Lag between event occurrence and reporting relative to granularity of time considered

Mark only one oval.



Data quality - sources of bias

#### 24. Outages

Potential for missing reports of information	Potential	for	missina	reports	of	info	rmatior
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Mark only one oval.



# 25. Censoring \*

Potential for censored data (events are known to have happened but the precise time of events is unknown i.e. to within an interval).

Mark only one oval.



#### 26. Truncation \*

Potential for truncated data (events are not known to have occurred but may be reported due to lack of observation such as conditioning on case report for onset data).

Mark only one oval.



27. Selection (ie unrepresentative relative to target population), including ascertainment bias/underreporting

Potential for these

Mark only one oval.



# 28. If there is the potential for selection bias what kind

	Tick all that apply.
	Ascertainment/underreporting Only more severe events observed Response is age (or other factor) dependent Location specific observation Socio-economic factors Confounding from external factors (i.e. environmental factors) Other:
Da	ta quality - bias characteristics
If b	ias of any kind of present what characteristics does it have?
29.	Time-varying
	Potential for biases to vary over time
	Mark only one oval.
	1 2 3 4 5
	Non
30.	Direction of bias
	Put 3 if unclear
	Mark only one oval.
	1 2 3 4 5
	Larg

# Data utility

What can you actually use it for? Identifies which transmission metrics it can inform and how directly it provides that information.

#### 31. Quantities informed \*

What target quantities does the data source inform. Does it inform it directly or indirectly (e.g. wastewater indirectly informs prevalence and community prevalence surveys with random sampling directly inform prevalence estimates).

Tick all that apply.

	Direct	Indirect	Not informed
Basic reproduction number			
Time-varying reproduction number			
Time-varying / basic reproduction number ratio			
Incidence			
Prevalence			
Heterogeneity in transmission (e.g. superspreading)			
Drivers of transmission			

32. Relationship of target population to general population \*

Can information on the target population be generalised to the general population

Mark only one oval.

	1	2	3	4	5		
It ca						Very	easily

Why can't the information on the target population be generalised population	to the general
actical considerations	
w feasible is it to use? Covers accessibility, sustainability, cost, and wheth Il with other data sources.	er it plays
Scalability *	Dropdown
How does the data collection scale with outbreak size	
Mark only one oval.	
independent	
Sub-linearly	
Linearly	
Sub-exponentially	
Exponentially	
More than exponentially	
Sustainability *	
Likelihood of continued collection/effort to collect	
Mark only one oval.	
1 2 3 4 5	
Very Very unlikely	
	population  actical considerations  w feasible is it to use? Covers accessibility, sustainability, cost, and wheth a sources.  Scalability * How does the data collection scale with outbreak size  Mark only one oval.  independent  Sub-linearly  Linearly  Sub-exponentially  Exponentially  More than exponentially  Sustainability *  Likelihood of continued collection/effort to collect  Mark only one oval.  1 2 3 4 5

36.	Cost *
	How resource-intensive is the collection?
	Mark only one oval.
	1 2 3 4 5
	Very O Very high
37.	Accessibility *
	How easily data can be obtained. For example due to privacy concerns.
	Mark only one oval.
	1 2 3 4 5
	Very Very easily
38.	Linkage potential * Ability to connect with other datasets  Mark only one oval.
	1 2 3 4 5
	Very Very easily
39.	What data sources if any have been linked to? The category/classification of data
	Tick all that apply.
	Confirmed cases time series
	Hospitalisation time series     Wastewater
	Contact tracing
	Other:

40.	Data format *
	The level of structure in the data source
	Mark only one oval.
	1 2 3 4 5
	Unst
41.	How generalisable are these findings for other pathogens? *
	Mark only one oval.
	1 2 3 4 5
	Not Very generalisable
42.	Which pathogens if any could your answers be generalised to?

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