

### Influenza and COVID-19 Surveillance Security graphs Agency

UKHSA publishes a national influenza and COVID-19 surveillance report which summaries the information from the surveillance systems which are used to monitor influenza, COVID-19 and other seasonal respiratory viruses in England.

Additional figures based on these surveillance systems are included in this slide set.

The figures presented in this slide set are based on data from week 48 (between) 27 November and 3 December 2023).

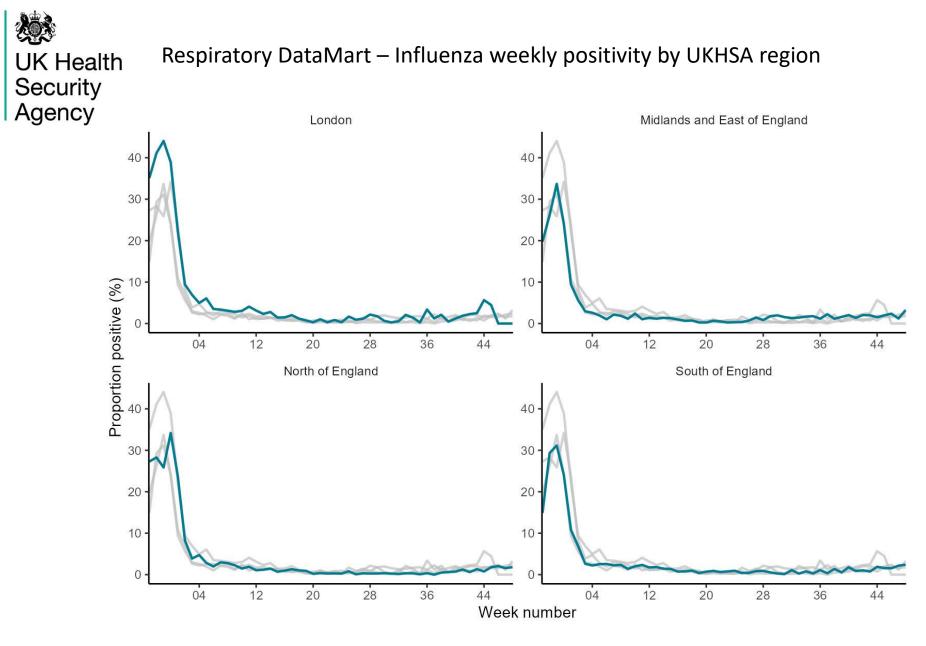


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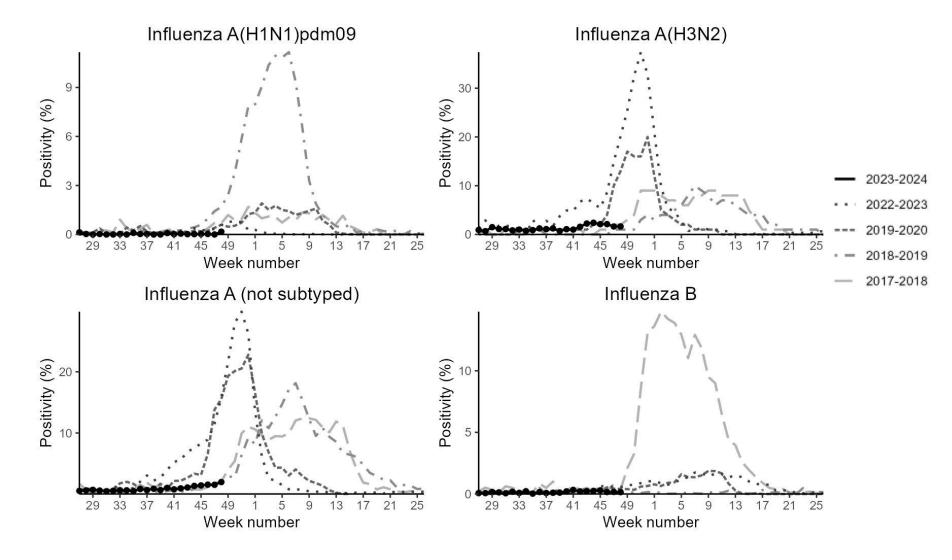
## Respiratory Datamart system (England)

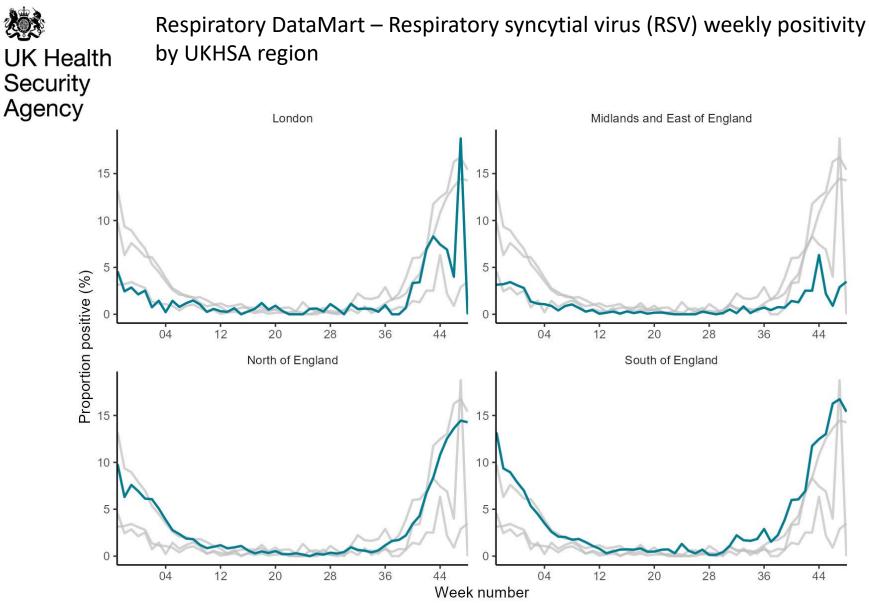


\*Changes in positivity in London should be interpreted with caution as there was a low number of samples this week and is subject to retrospective updates



### Respiratory DataMart – Influenza subtypes

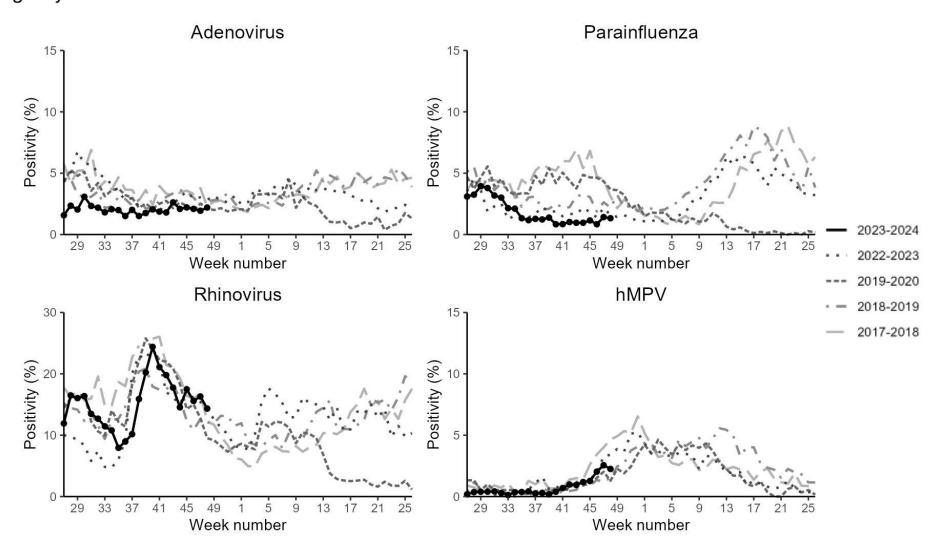




\*Changes in positivity in London should be interpreted with caution this week and is subject to retrospective updates

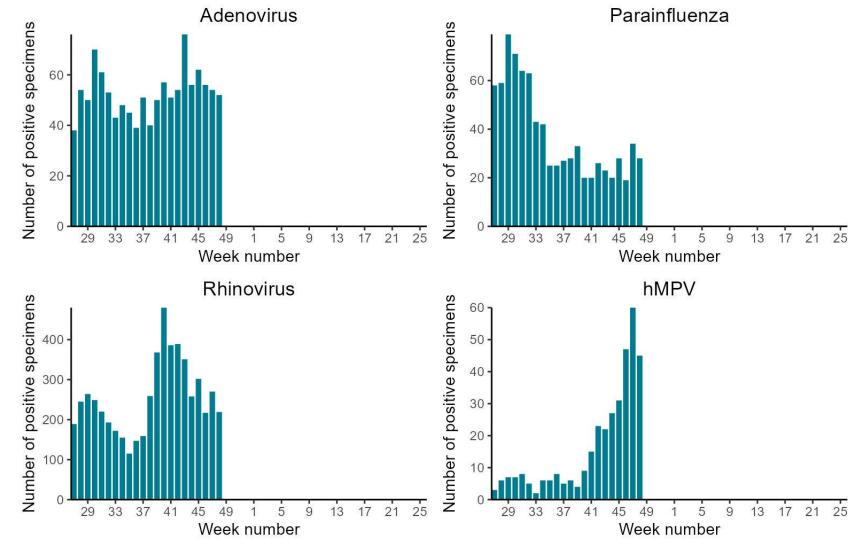
UK Health Security Agency

### Respiratory DataMart – other respiratory viruses

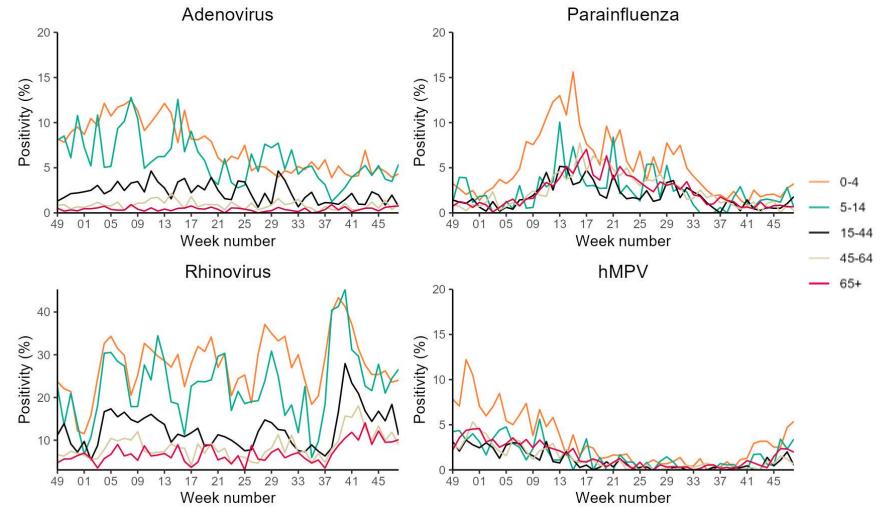




### Respiratory DataMart – other respiratory viruses



### WK Health Respiratory DataMart – other respiratory viruses Security Agency





## Confirmed COVID-19 episodes in England

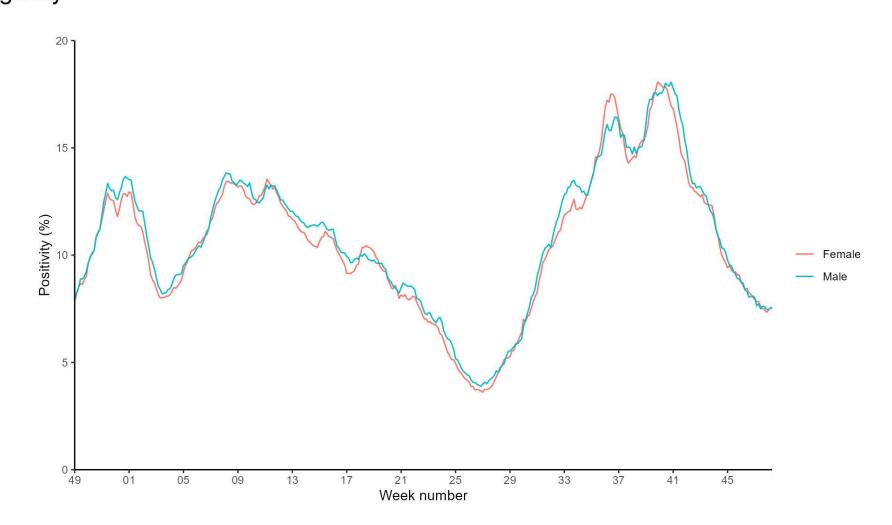


### Confirmed COVID-19 episodes in England

#### **Data Information**

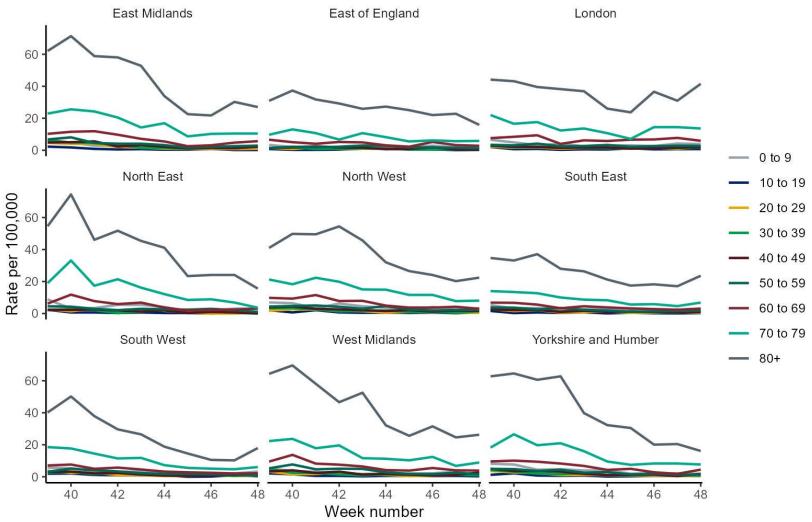
- From the week 32 report onwards, case rates have been updated to use the latest ONS population estimates for mid-2020. Previously case rates were calculated using the mid-2019 population estimates
- From 11 January 2022 the requirement for <u>confirmatory PCR testing in individuals who test positive using a lateral flow device was</u> <u>temporarily removed</u>.
- Rates by ethnicity and IMD quantile will continue to be presented using the mid-2019 estimates, until the mid-2020 estimates become available.
- From 31 January 2022, UKHSA moved all COVID-19 case reporting in England to use a new episode-based definition which includes
  possible reinfections. Each infection episode is counted separately if there are at least 91 days between positive test results (PCR or
  LFD). Each infection episode begins with the earliest positive specimen date. Further information can be found on the UK COVID-19
  dashboard.
- Since 1 April 2022, free universal symptomatic and asymptomatic testing for the general public in England is no longer available, as outlined in the plan for <u>living with COVID-19</u>. As such, there will be a reduction in the reporting of data obtained through Pillar 2 from April 2022 onwards. Data in this report should be interpreted in the context of this change to testing. <u>Public health guidance</u> remains in place for cases and their close contacts. Additionally, further changes in <u>testing policy</u> are in effect since 1 April 2023, which may affect case rates and positivity rates.

UK Health Security Agency Seven-day rolling average PCR positivity (%) of confirmed COVID-19 cases tested by sex under Pillar 1



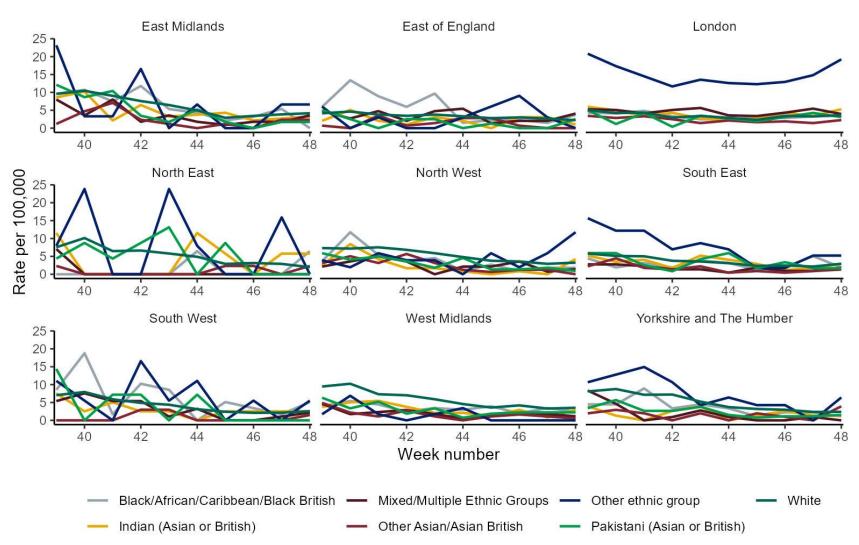


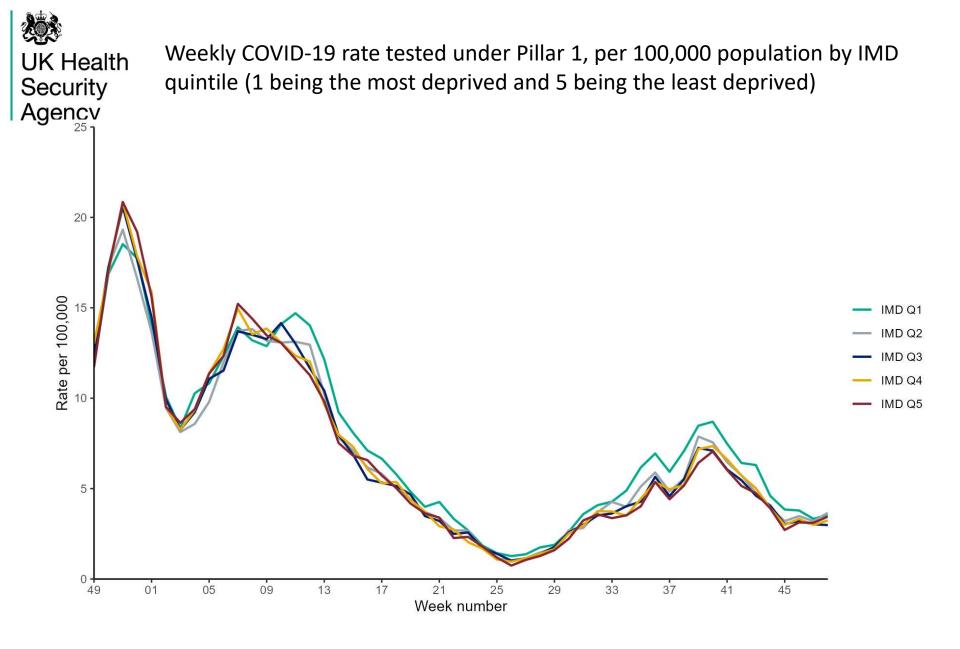
## Weekly COVID-19 episodes tested under Pillar 1, per 100,000 population by age group and region, weeks 39 to 48



UK Health Security Agency

Weekly COVID-19 episodes tested under Pillar 1, per 100,000 population by ethnicity and region, weeks 39 to 48



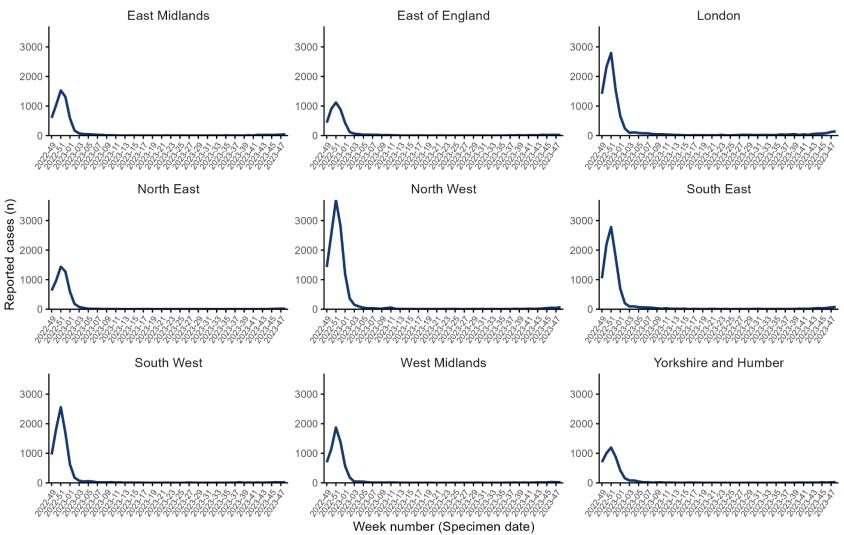




# Second generation surveillance system (SGSS)

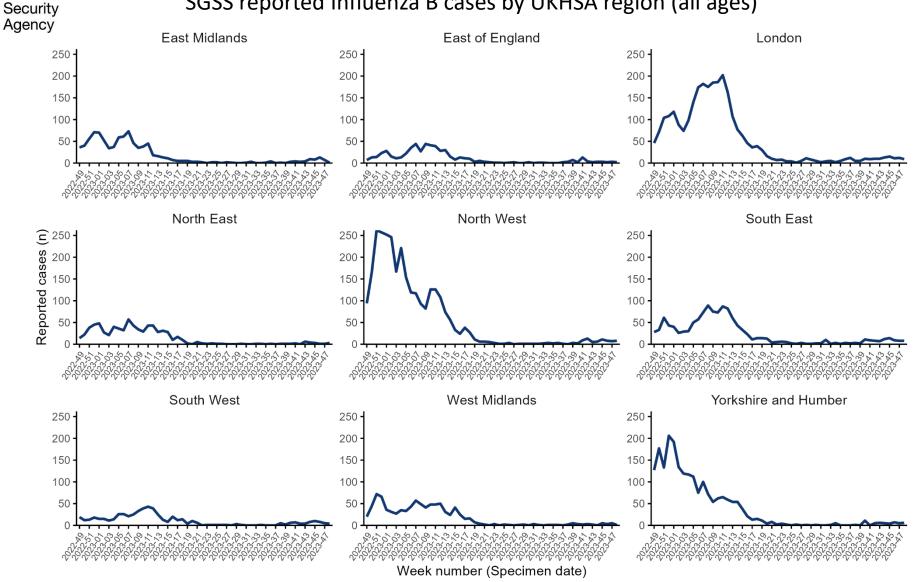


### SGSS reported Influenza A cases by UKHSA region (all ages)



The presented figures are based on laboratory reports through SGSS. Testing and reporting procedures vary by virus, UKHSA region and over time, including short-term trends in testing. Therefore comparisons should be done with caution.

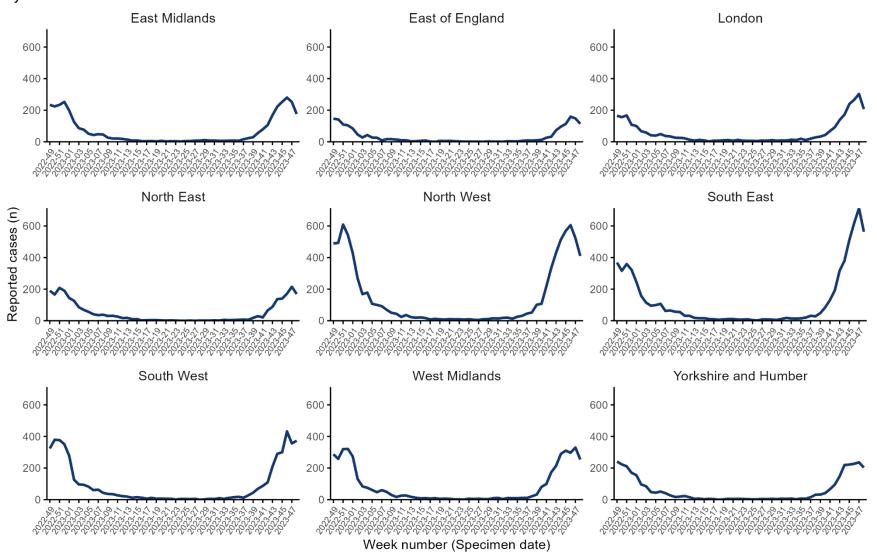
### SGSS reported Influenza B cases by UKHSA region (all ages)



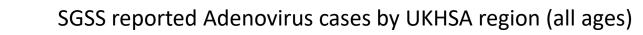
The presented figures are based on laboratory reports through SGSS. Testing and reporting procedures vary by virus, UKHSA regions and over time, including short-term trends in testing. Therefore comparisons should be done with caution.

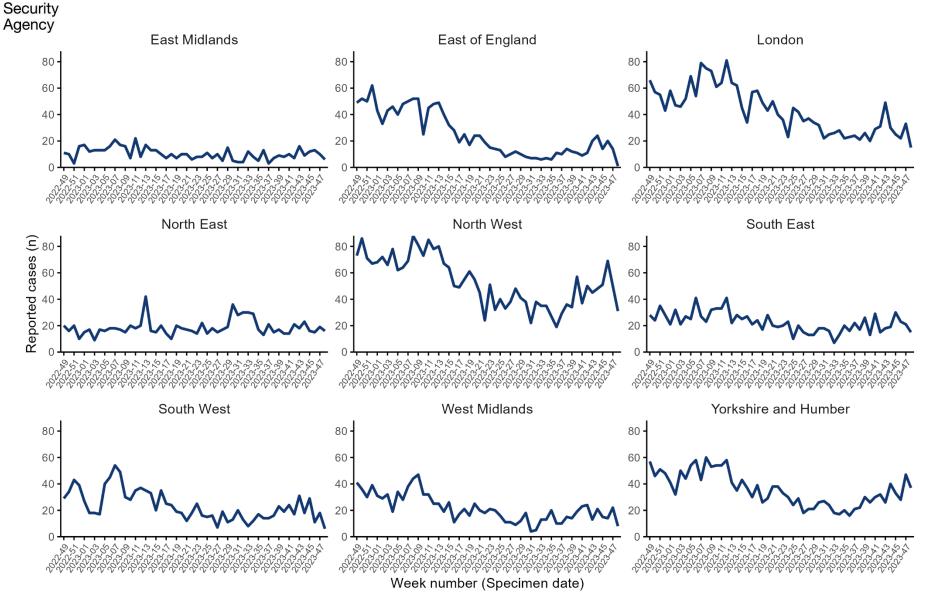
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### SGSS reported RSV cases by UKHSA region (all ages)



The presented figures are based on laboratory reports through SGSS. Testing and reporting procedures vary by virus, UKHSA region and over time, including short-term trends in testing. Therefore comparisons should be done with caution.

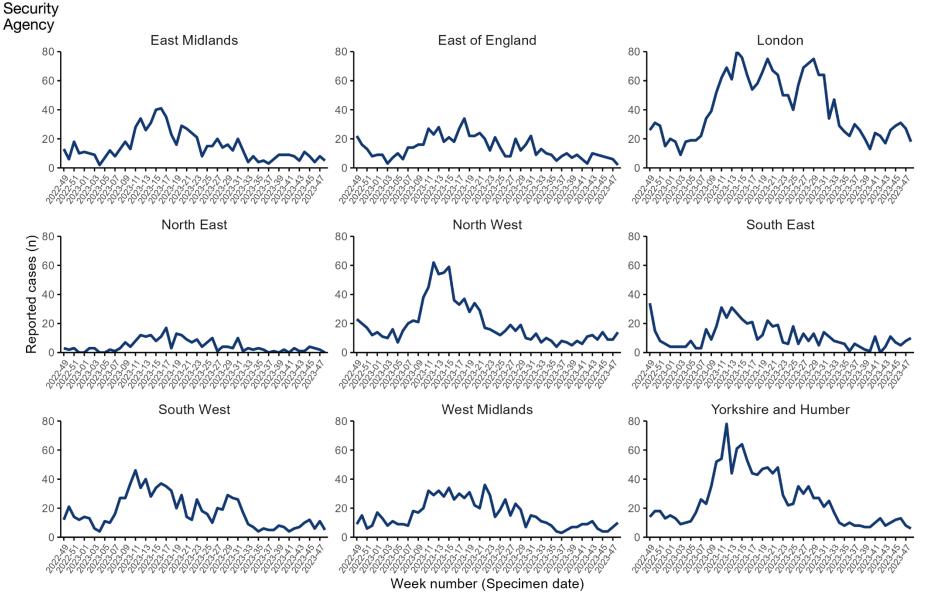




The presented figures are based on laboratory reports through SGSS. Testing and reporting procedures vary by virus, UKHSA region and over time, including short-term trends in testing. Therefore comparisons should be done with caution.

**M** 

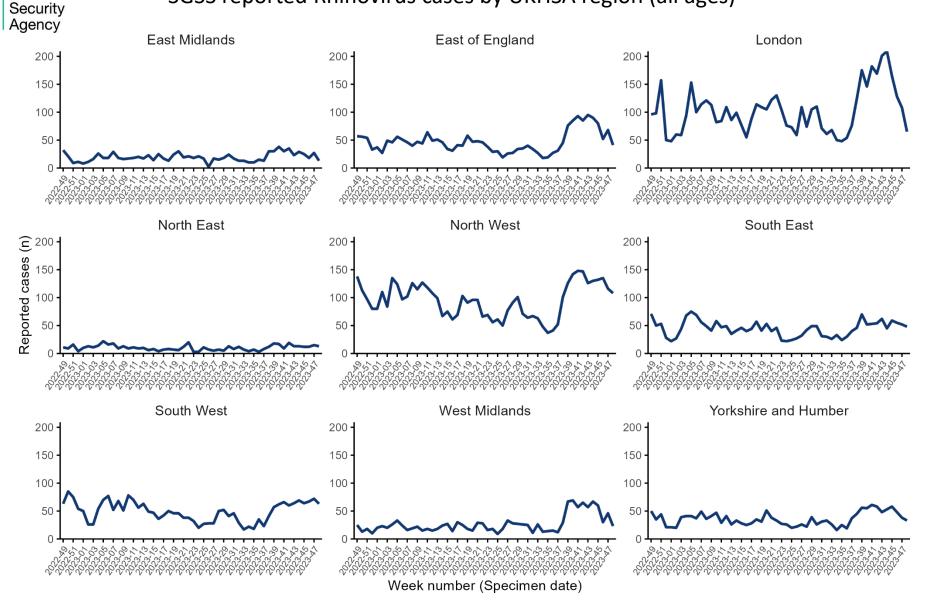
### SGSS reported Parainfluenza cases by UKHSA region (all ages)



The presented figures are based on laboratory reports through SGSS. Testing and reporting procedures vary by virus, UKHSA region and over time, including short-term trends in testing. Therefore comparisons should be done with caution.

**M** 

### SGSS reported Rhinovirus cases by UKHSA region (all ages)

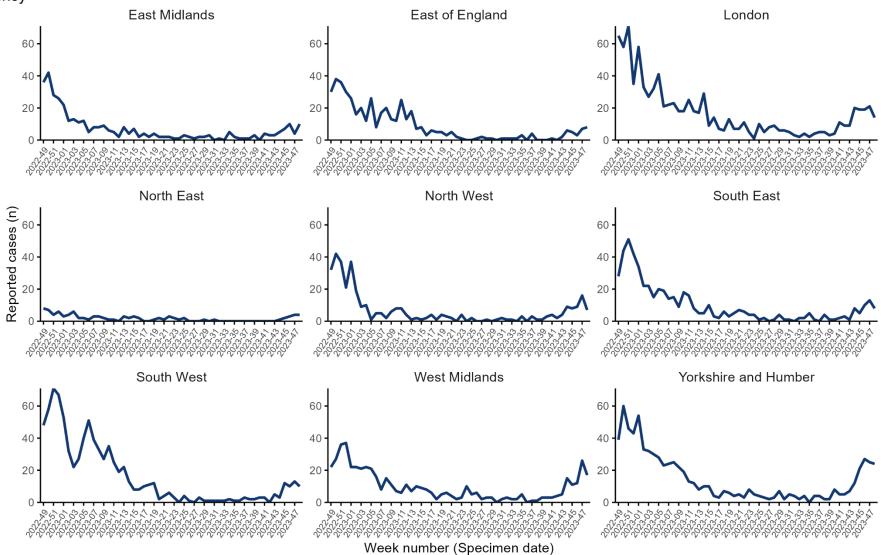


The presented figures are based on laboratory reports through SGSS. Testing and reporting procedures vary by virus, UKHSA region and over time, including short-term trends in testing. Therefore comparisons should be done with caution. 22

**M** 



### SGSS reported hMPV cases by UKHSA region (all ages)



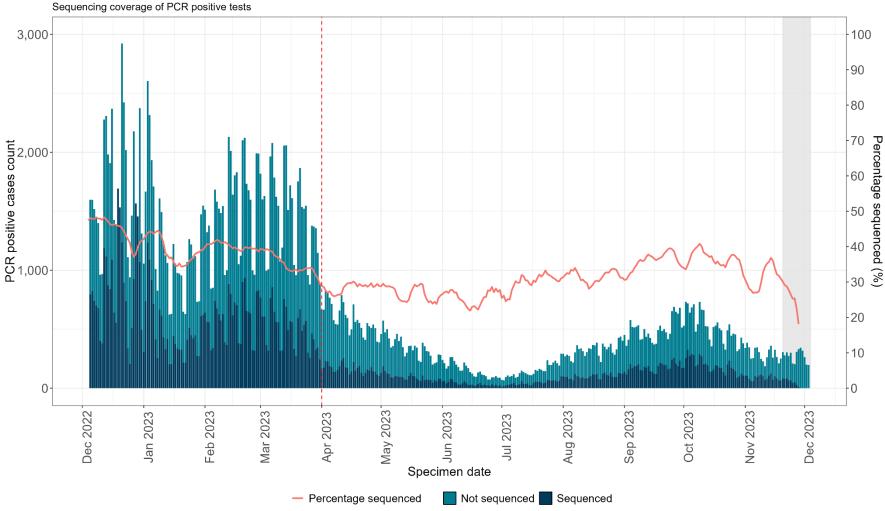
The presented figures are based on laboratory reports through SGSS. Testing and reporting procedures vary by virus, UKHSA region and over time, including short-term trends in testing. Therefore comparisons should be done with caution.



## SARS-CoV-2 Whole Genome Sequencing (WGS) coverage, England



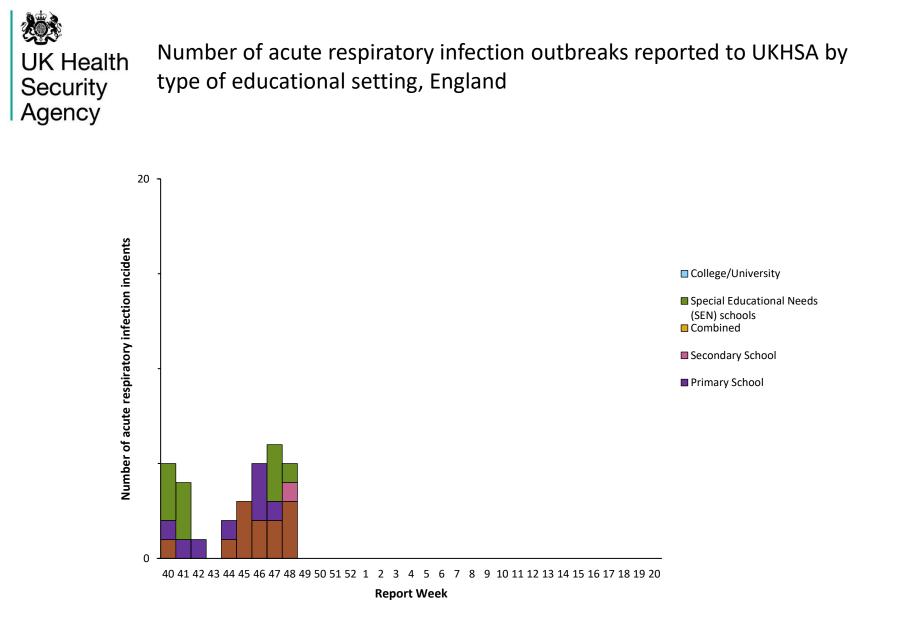
## SARS-CoV-2 coverage of sequencing with a valid result and genotyping over time



Grey shading was applied to the previous 14 days to account for reporting delays in sequencing data. Cases where the individual only tested using a lateral flow device are not included in the percentage denominator.



## **Community surveillance**





## Primary Care surveillance



7 December 2023

UK Health Security Agency

General practice Influenza-like-illness consultation rates per 100,000 population, UK administrations

GP ILI consultation rates (all ages)	Week number								
	40	41	42	43	44	45	46	47	48
England (RCGP)	3.5	3.2	3.5	3.2	3.3	3.8	3.4	3.8	4.6
Wales	5.0	3.1	1.7	2.9	3.6	4.0	3.3	4.2	4.3
Scotland	1.5	0.7	2.7	2.6	1.9	7.0	2.3	4.3	4.7
Northern Ireland	3.3	3.2	3.6	3.4	2.9	4.2	3.7	3.7	4.2



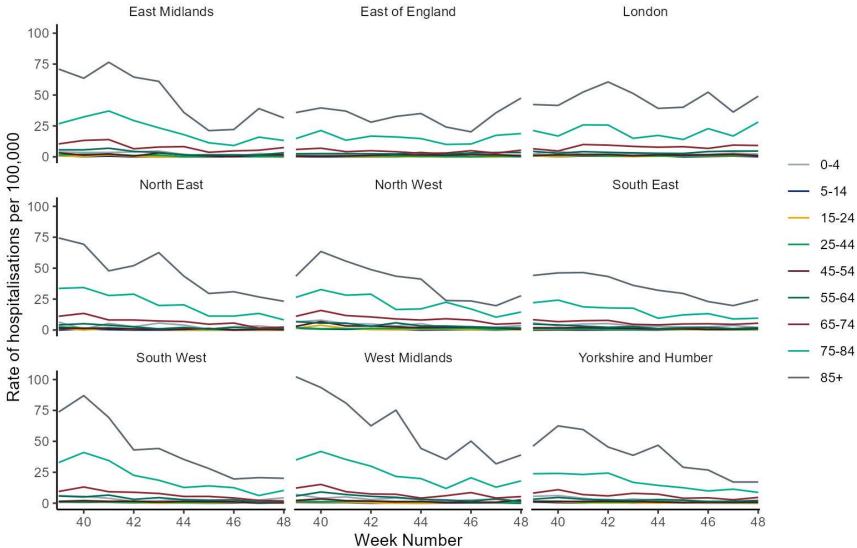
## Secondary Care surveillance



7 December 2023



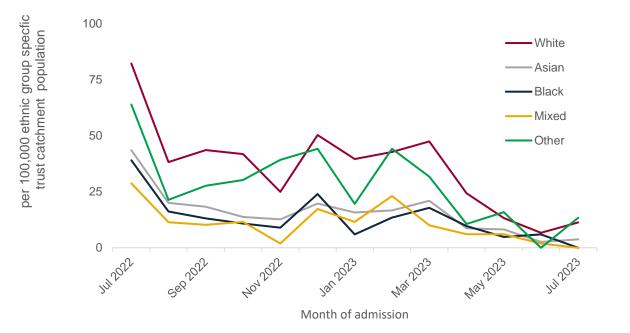
## Weekly COVID-19 hospitalisation rate per 100,000 trust catchment population by age group and region, weeks 39 to 48



Data extracted on 23 August 2023

UK Health Security Agency

Rate of COVID-19 hospitalisation (to all levels of care including ICU-HDU) by ethnic group, per 100,000 ethnic group specific trust catchment population, England





### Preceding, co- and secondary infections in persons with COVID-19 and influenza in England, Jul 2022 – 6<sup>th</sup> Dec 2023

HCAI, Fungal, AMR, AMU & Sepsis Division

### Preceding/co-/secondary infections with COVID-19

### Background

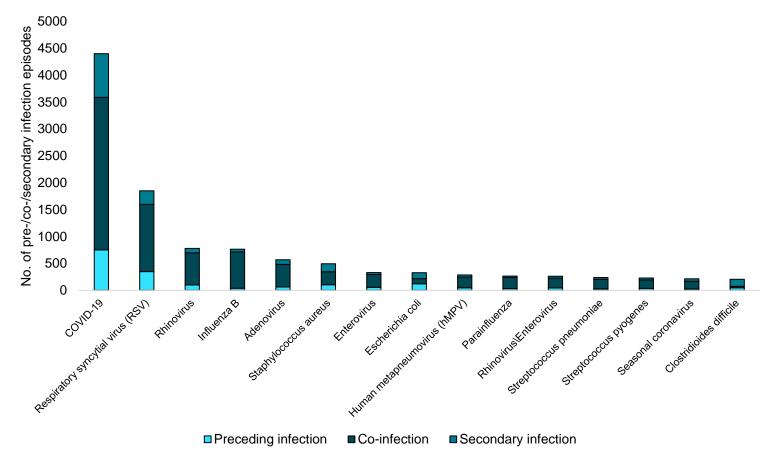
- Numbers of preceding/co-/secondary infection remain low across UKHSA surveillance systems.
- Free community testing ended 31 March 2022 as part of the government's Living with COVID-19 plan, with asymptomatic testing continuing in some settings. As of 31 August 2022, asymptomatic testing in all settings, including hospitals, has been paused. Please use caution when comparing incidence of bacterial, fungal and viral preceding/co-/secondary infections with COVID-19 over time due to these differences in testing strategies.
- Published data analysed from pandemic wave 1 indicated increased mortality associated with COVID-19 and <u>influenza</u>, <u>key bacterial and fungal infections</u> and <u>invasive pneumococcal disease</u> (IPD) in comparison to persons without co/secondary infection.
- <u>Data analysis</u> from wave 1 indicated that *Aspergillus* and *candidemia* cases had increased risk of mortality in comparison to patients without co/secondary infection.

## Surveillance of bacterial, fungal and respiratory viral infections in persons with COVID-19 in England

### **Data information**

- Data are provisional and subject to change due to possible delayed reporting of microbiological samples
- Relative undertesting for other pathogens may result in an underestimate of preceding/co-/secondary infection cases. In addition, testing varies between pathogens therefore caution should be used in comparing preceding/co-/secondary infection rates between different pathogens
- Preceding/co-/secondary infections refers to when a person has a COVID-19 infection with one or more other pathogen (Please see Appendix 1 – Preceding/co-/secondary infection definitions.)
  - Preceding infection: SARS-CoV-2 detected after another pathogen
  - Co-infection: SARS-CoV-2 and other pathogen detected at the same time
  - Secondary infection: SARS-CoV-2 detected before another pathogen
- The following outputs included in this section have been produced via the Unified Infection Dataset (UID)
- Bacterial, fungal and respiratory viral infection data sources:
  - Fungal, bacterial and respiratory viral data (excluding *Clostridioides difficile*): Second Generation Surveillance System (SGSS)
  - Respiratory viral data: Respiratory Datamart
  - *Clostridioides difficile*: HCAI Data Capture System

Most frequent bacterial/fungal/respiratory viral infections, by timing of diagnosis, in persons with influenza in England from ISO week 27 of 2022



### Key findings:

From ISO week 27 of 2022, the most frequent organisms identified were COVID-19, RSV, and rhinovirus.

\*The baseline infection is any type of influenza (influenza A or B or both) for all bacterial/fungal/respiratory viral preceding/co-/secondary infections except for influenza B where the baseline infection is influenza A.

### Appendix 1: Pre-/co-/secondary infection definitions

The day pertains to the date of the sample collection that yielded a positive result. These definitions do not apply to persistent COVID-19 patients. Patients with persistent COVID-19 require independent clinical assessment.

Influenza A         +/- 1d         2-28d^           Influenza B         +/- 1d         2-28d^           RSV         +/- 1d         2-28d	
RSV +/- 1d 2-28d	
Adenovirus +/- 1d 2-28d	
Enterovirus +/- 1d 2-28d	
Human metapneumovirus +/- 1d 2-28d	
Parainfluenza (any subtype) +/- 1d 2-28d	
Seasonal coronavirus +/- 1d * 2-28d	
Rhinovirus +/- 1d 2-28d	
Co-infections in ECMO patient (patients with most severe clinical respiratory signs)	
ECMO patients Individual case review Individual case review	
Blood stream and respiratory infections (bacterial and fungal)	
Achromobacter xylosoxidans +/- 1d 2-28d	
Acinetobacter spp., +/- 1d 2-28d	
Aspergillus +/- 1d 2-28d (pre) 2-60d (post, continually hospitalised patients	only)
Bordetella pertussis       +/- 28 d Culture/PCR (based on pertussis sample date)       N/A (Pertussis presentation is often delayed)         +/- 28 Serology/Oral fluid (anti-pertussis toxin Ig) (based on pertussis symptom onset date, excluding cases without onset date)       N/A (Pertussis presentation is often delayed)	
Burkholderia cepacia +/- 1d 2-28d	
Candida spp +/- 1d 2-28d (pre) 2-60d (post, continually hospitalised patients	only)
Chlamydia pneumoniae 0-7d PCR 0-7d PCR within 14-28 d (8-13d PCR*)	
Enterobacter spp., +/- 1d 2-28d	
Enterococcus spp. +/- 1d 2-28d	
E. coli +/- 1d 2-28d	
Haemophilus influenzae +/- 2d 3-28d	Continued overlaaf

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### Appendix 1 continued: Pre-/co-/secondary infection definitions

Organism	Definition co-infection with SARS-CoV-2†	Definition of infection pre-SARS-CoV-2 infection (other pathogen is primary infection) or Definition of post SARS-CoV-2 secondary infection (SARS-CoV-2 is primary infection)				
Blood stream and respiratory infections (ba	cterial and fungal)					
Klebsiella spp.	+/- 1d	2-28d				
Legionella pneumophila/species	Individual case review	Individual case review				
Mycoplasma pneumoniae	0-7d PCR, IgM serology 0-21d <16y	PCR within 14-28 d (8-13d PCR*)				
Neisseria meningitidis	+/- 2d	3-28d				
Pseudomonas spp.,	+/- 1d	2-28d				
Serratia spp.,	+/- 1d	2-28d				
Staphylococcus aureus	+/- 1d	2-28d				
Coag-neg Staphylococcus (S. haemolyticus)	+/- 1d	2-28d				
Stenotrophomonas spp., (S. maltophilia)	+/- 1d	2-28d				
Streptococcus spp. <b>‡</b>	+/- 1d	2-28d				
Streptococcus pneumoniae	+/- 2d	3-28d				
Tuberculosis						
Mycobacterium tuberculosis	Individual case review	Individual case review				
Pathogens of the immunocompromised (eg	I HIV)					
HIV	Individual case review	Individual case review				
Gastrointestinal infections						
Listeria	0-5d *	Individual case review				
Campylobacter	0-5d *	Individual case review				
Shiga toxin-producing E. coli (STEC)	0-5d *	Individual case review				
Norovirus	0-5d *	Individual case review				
Salmonella	0-5d *	Individual case review				
Shigella	0-5d *	Individual case review				
Anaerobes						
C. difficile	+/- 1d	2-28d				
Bacteroides sp. ( <i>B. fragilis</i> and non-fragilis Bacteroides)	+/- 1d	2-28d				

See next slides for notes

### Appendix 1 continued: Pre-/co-/secondary infection definitions

### Notes

+ From the first specimen date of a SARS-CoV-2 infection episode.

\* Additional data check required. (Resistance is not detailed, data for MERS is not currently available).

^ Definition post- SARS-CoV-2 secondary infection (SARS-CoV-2 is primary infection). This has been extended from prior 14d secondary infection definition for influenza used by UKHSA to account for disparities in testing throughout the 28d period after SARS-CoV-2 detection.

**‡** Streptococcus species includes the following groups and species:

Group	Species/other names		
Anginosus Group	Streptococcus anginosus; Streptococcus constellatus (Streptococcus constellatus subspecies constellatus Streptococcus		
	constellatus subspecies pharynges); Streptococcus Group F; Streptococcus intermedius; Streptococcus milleri group;		
	Streptococcus sinensis		
Bovis Group	Streptococcus alactolyticus; Streptococcus bovis untyped; Streptococcus equinus; Streptococcus gallolyticus subspecies		
	gallolyticus (Streptococcus bovis biotype I); Streptococcus infantarius (Streptococcus infantarius sp infantarius; Streptococcus		
	bovis biotype II); Streptococcus lutetiensis; Streptococcus infantarius subspecies coli (Streptococcus bovis biotype II);		
	Streptococcus pasteurianus (Streptococcus bovis biotype II)		
Closely Related Genera	Abiotrophia spp.; Aerococcus spp.; Faklamia spp.; Gemella spp.; Globicatella sanguinis; Granulicatella spp.; Leuconostoc		
	spp.; Pedicoccus spp.; Peptostreptococcus spp.		
Mitis Group	Streptococcus cristatus; Streptococcus mitior; Streptococcus mitis; Streptococcus oralis; Streptococcus pseudopneumoniae;		
	Streptococcus infantis; Streptococcus peroris		
Mutans Group	Streptococcus mutans; Streptococcus sobrinus		
Other streptococci (including but not	Anaerobic streptococcus; Streptococcus acidominimus; Streptococcus spp., other named/not fully identified; Streptococcus		
limited to)	suis; Streptococcus uberis		
Salivarius Group	Streptococcus vestibularis; Streptococcus thermophilus		
Sanguinis Group	Streptococcus gordonii; Streptococcus massiliensis; Streptococcus parasanguinis; Streptococcus sanguinis		
Streptococcus Group A	Group A; Streptococcus pyogenes; Streptococcus dysgalactiae subspecies equisimilis		
Streptococcus Group B	Group B; Streptococcus agalactiae		
Streptococcus Group C	Group C; Streptococcus dysgalactiae subspecies equisimilis; Streptococcus equi subspecies zooepidemicus		
Streptococcus Group G	Group G; Streptococcus canis; Streptococcus dysgalactiae subspecies equisimilis		