



PHE Weekly National Influenza Report

Summary of UK surveillance of influenza and other seasonal respiratory illnesses

13 April 2017 – Week 15 report (up to week 14 data)

This report is published weekly on the [PHE website](#). For further information on the surveillance schemes mentioned in this report, please see the [PHE website](#) and the [related links](#) at the end of this document.

| [Summary](#) | [Community surveillance](#) | [GP consultation rates](#) | [Hospitalisations](#) | [All-cause mortality](#) | [Microbiological surveillance](#) | [Vaccination](#) | [International](#) | [Acknowledgements](#) | [Related links](#) |

Summary

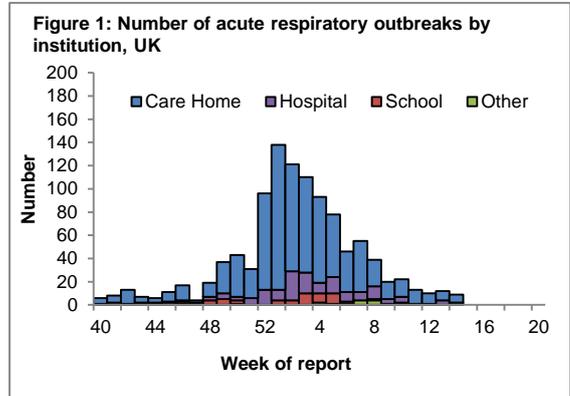
During week 14 (ending 09 April 2017), influenza activity continues to decrease across all indicators. The Department of Health has issued an [alert](#) on the prescription of antiviral medicines by GPs.

- [Community influenza surveillance](#)
 - Through the GP In Hours Syndromic Surveillance system, GP consultations for influenza-like illness (ILI) were decreasing in week 14.
 - Nine new acute respiratory outbreaks have been reported in the past 7 days. Seven outbreaks were from care homes, where two tested positive for influenza (one influenza A(not subtyped) and one influenza B). Two outbreaks were from hospitals where one tested positive for influenza A(not subtyped) and the other for influenza B.
- [Overall weekly influenza GP consultation rates across the UK](#)
 - In week 14, the overall weekly influenza-like illness (ILI) GP consultation rate was 4.3 per 100,000 in England compared to 3.8 per 100,000 in the previous week. This is below the baseline threshold of 14.3 per 100,000 for this season. In the devolved administrations, ILI rates have decreased in Wales and Scotland but increased in Northern Ireland compared to the previous week.
- [Influenza-confirmed hospitalisations](#)
 - In week 14, there were four admissions to ICU/HDU with confirmed influenza (two influenza B, one influenza A(unknown subtype) and one influenza A(H3N2)) were reported across the UK (119/156 Trusts in England) through the USSS mandatory ICU scheme with a rate of 0.01 per 100,000 compared to 0.02 per 100,000 in the previous week.
 - In week 14, there were four hospitalised confirmed influenza cases (two influenza A(H3N2), one influenza A(not subtyped) and one influenza B) reported through the USSS sentinel hospital network (14 NHS Trusts across England), with a rate of 0.08 per 100,000, compared to 0.19 per 100,000 in the previous week.
 - No confirmed influenza admissions have been reported from the six Severe Respiratory Failure centres in the UK in week 14.
- [All-cause mortality data](#)
 - In week 14 2017, no statistically significant excess all-cause mortality by week of death was seen through the EuroMOMO algorithm in England and in the devolved administrations.
- [Microbiological surveillance](#)
 - 11 samples tested positive for influenza (11 influenza B) through the UK GP sentinel swabbing schemes, with an overall positivity of 28.9% compared to 12.7% in week 13.
 - 47 influenza positive detections were recorded through the DataMart scheme (17 influenza A(H3N2), two influenza A(H1N1)pdm09 and 28 influenza B) in week 14. The overall positivity was at 3.5% in week 14 compared to 3.4% in week 13, which is below the threshold for 2016/17 season of 8.6%. The highest age-specific positivities were seen in the 65+ year olds (7.5%).
- [Vaccination](#)
 - Provisional data from the fifth monthly collection of influenza vaccine uptake by frontline healthcare workers show 63.4% were vaccinated by 28 February 2017, compared to 50.8% vaccinated in the previous season by 29 February 2016. The report provides uptake at Trust level.
 - Up to week 04 2017, in 85.0% of GP practices reporting weekly to Immform, the provisional proportion of people in England who had received the 2016/17 influenza vaccine in targeted groups was as follows: 48.5% in under 65 years in a clinical risk group, 44.9% in pregnant women and 70.5% in 65+ year olds. In 88.1% of GP practices reporting to Immform, the provisional proportion of children in England who had received the 2016/17 influenza vaccine was as follows: 39.0% in all 2 year olds, 41.6% in all 3 year olds and 33.8% in all 4 year olds.
 - Provisional data from the fourth monthly collection of influenza vaccine uptake in GP patients up to 31 January 2017 has been published. The [report](#) provides uptake at national, Area Team (AT), Clinical Commissioning Group (CCG) and by Local Authority (LA) levels.
 - Provisional [data](#) from the fourth monthly collection of influenza vaccine uptake for children of school years 1, 2 and 3 age show the provisional proportion of children in England who received the 2016/17 influenza vaccine via school, pharmacy or GP practice by 31 January 2017 in targeted groups was as follows: 57.6% in children of school Year 1 age (5-6 years); 55.3% in children of school Year 2 age (6-7 years); 53.3% in children of school Year 3 age (7-8 years).
- [International situation](#)
 - Globally, influenza activity in the temperate zone of the northern hemisphere continued to decrease. Worldwide, influenza A(H3N2) and influenza B viruses were predominant during this reporting period. In South Asia, influenza activity with mainly influenza A(H1N1) remained elevated. The [vaccine recommendation for the 2017-2018 northern hemisphere](#) influenza season has been made.

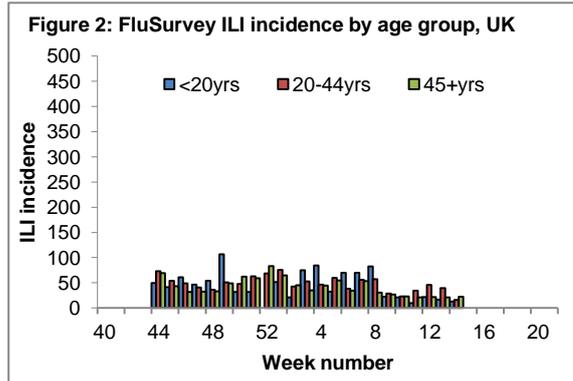
Through the GP In Hours Syndromic Surveillance system, GP consultations for influenza-like illness (ILI) were decreasing in week 14. Nine new acute respiratory outbreaks were reported in the past 7 days.

- PHE Real-time Syndromic Surveillance
 - During week 14, GP consultations for influenza-like illness were decreasing.
 - For further information, please see the syndromic surveillance [webpage](#).

- Acute respiratory disease outbreaks
 - Nine new acute respiratory outbreaks have been reported in the past 7 days. Seven outbreaks were from care homes, where two tested positive for influenza (one influenza A(not subtyped) and one influenza B). Two outbreaks were from hospitals where one tested positive for influenza A(not subtyped) and the other for influenza B.
 - Outbreaks should be recorded on HPZone and reported to the local Health Protection Teams and Respscidsc@phe.gov.uk.

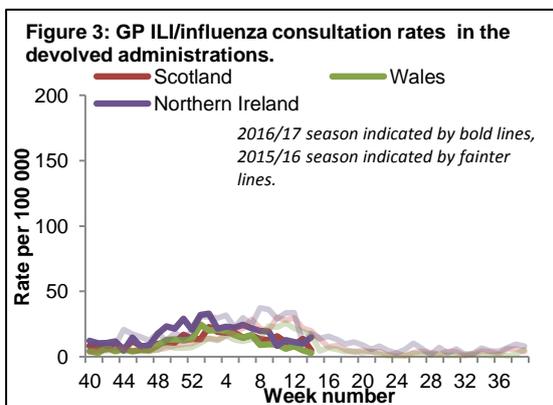


- FluSurvey
 - Internet-based surveillance of influenza-like illness in the general population is undertaken through the FluSurvey. A project run jointly by PHE and the London School of Hygiene and Tropical Medicine.
 - The overall ILI rate (all age groups) for week 14 was 20.3 per 1,000 (37/1,789 people reported at least 1 ILI), with the 45+ years age group reporting a higher rate of 22.2 per 1,000.
 - If you would like to become a participant of the FluSurvey project please do so by visiting the <https://flusurvey.org.uk/en/accounts/register/> website for more information.



In week 14, the overall weekly influenza-like illness GP consultation rate is low and is below the baseline threshold in England. In the devolved administrations, ILI rates have decreased in Wales and Scotland but increased in Northern Ireland compared to the previous week.

- Influenza/Influenza-Like-Illness (ILI)



Northern Ireland

- The Northern Ireland ILI rate has increased at 14.9 per 100,000 in week 14 compared to 10.0 per 100,000 in week 13. This remains below the baseline threshold (47.9 per 100,000).
- The highest rates were seen in the 65-74 year olds (29.0 per 100,000) and 45-64 year olds (16.2 per 100,000).

Wales

-The Welsh ILI rate decreased at 2.9 per 100,000 in week 14 compared to 4.9 per 100,000 in week 13 (Figure 3). This is below the baseline threshold (10.3 per 100,000).

- The highest rates were seen in the 1-4 year olds (6.8 per 100,000) and 15-44 year olds (4.2 per 100,000).

Scotland

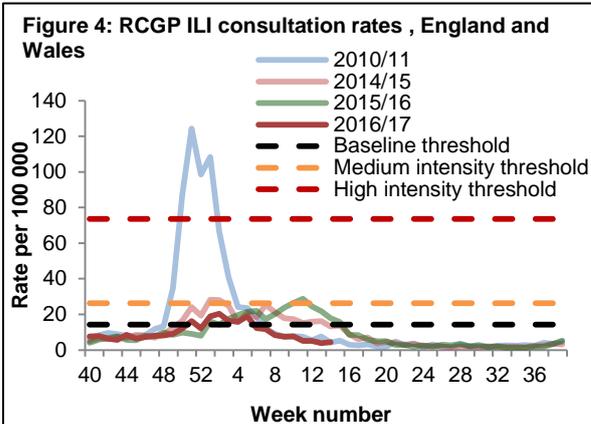
-The Scottish ILI rate decreased at 5.0 per 100,000 in week 14 compared to 13.6 per 100,000 in week 13 (Figure 3). This remains below the baseline threshold (36.1 per 100,000).

-The highest rates were seen in 45-64 year olds (7.0 per 100,000) and 75+ year olds (5.6 per 100,000).

RCGP (England and Wales)

- The weekly ILI consultation rate through the RCGP surveillance is at 4.3 per 100,000 in week 14 compared to 3.8 per 100,000 in week 13. This is below the baseline threshold (14.3 per 100,000) (Figure 4*). By age group, the highest rates were seen in 1-4 year olds (12.7 per 100,000) and 15-44 year olds (6.6 per 100,000).

**The Moving Epidemic Method has been adopted by the European Centre for Disease Prevention and Control to calculate thresholds for GP ILI consultations for the start of influenza activity in a standardised approach across Europe.*



GP In Hours Syndromic Surveillance System (England)

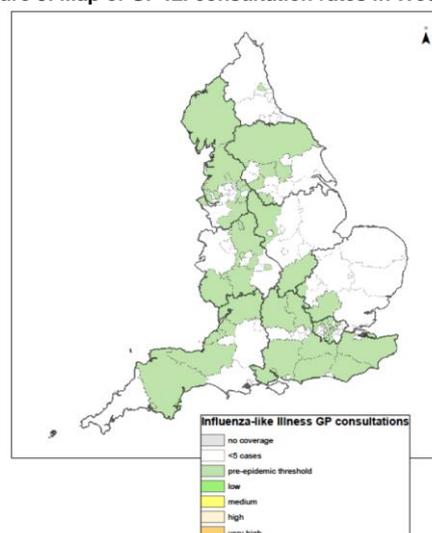
-The weekly ILI consultation rate through the GP In Hours Syndromic Surveillance system is at 3.1 per 100,000 in week 14 (Figure 5).

Figure 5 represents a map of GP ILI consultation rates in week 14 across England by Local Authorities, using influenza-like illness surveillance thresholds.

Thresholds are calculated using a standard methodology for setting ILI thresholds across Europe (the "Moving Epidemic Method" (MEM)) and are based on six previous influenza seasons (excluding the 2009/10 H1N1 pandemic)

-For further information, please see the syndromic surveillance [webpage](#).

Figure 5: Map of GP ILI consultation rates in Week 14



Influenza confirmed hospitalisations

[| Back to top |](#)

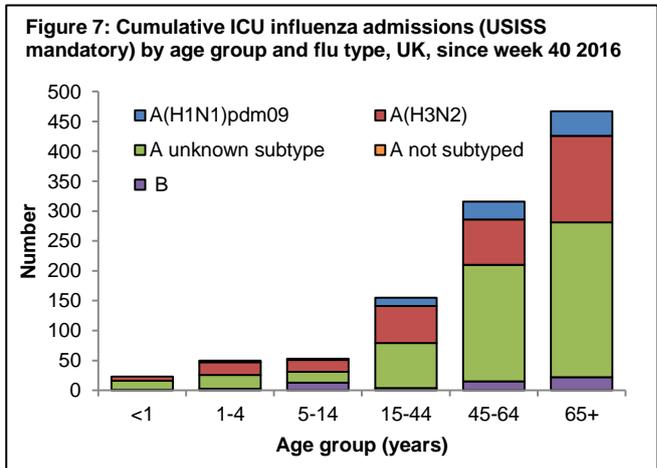
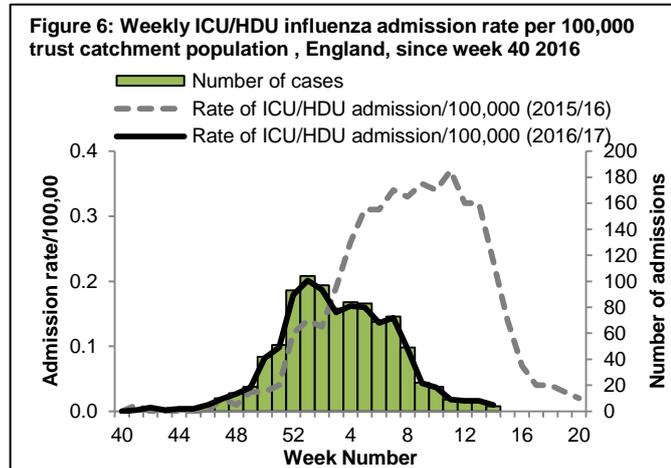
In week 14, there were four admissions to ICU/HDU with confirmed influenza (two influenza B, one influenza A(unknown subtype) and one influenza A(H3N2)) reported through the USISS mandatory ICU/HDU surveillance scheme across the UK (119 Trusts). Four hospitalised confirmed influenza cases (two influenza A(H3N2), one influenza A(not subtyped) and one influenza B) were reported through the USISS sentinel hospital network across England (14 Trusts).

A national mandatory collection (USISS mandatory ICU scheme) is operating in cooperation with the Department of Health to report the number of confirmed influenza cases admitted to Intensive Care Units (ICU) and High Dependency Units (HDU) and number of confirmed influenza deaths in ICU/HDU across the UK. A confirmed case is defined as an individual with a laboratory confirmed influenza infection admitted to ICU/HDU. In addition a sentinel network (USISS sentinel hospital network) of acute NHS trusts is established in England to report weekly laboratory confirmed hospital admissions. Further information on these systems is available through the [website](#). Please note data in previously reported weeks are updated and so may vary by week of reporting

- Number of new admissions and fatal confirmed influenza cases in ICU/HDU (USISS mandatory ICU scheme), UK (week 14)

- In week 14, there were four admissions to ICU/HDU with confirmed influenza (two influenza B, one influenza A(unknown subtype) and one influenza A(H3N2)) reported across the UK (119/156 Trusts in England) through the USISS mandatory ICU scheme, with a rate of 0.01 per 100,000 compared to a rate of 0.02 per 100,000 in week 13 (Figures 6 and 7). Two deaths were reported to have occurred in week 14.

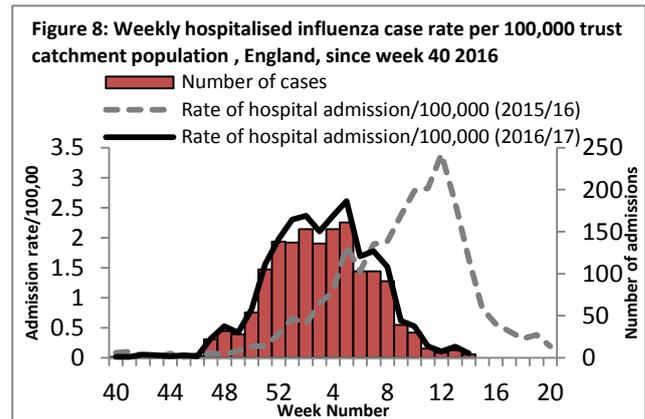
A total of 1,064 admissions (585 influenza A(unknown subtype), 331 influenza A(H3N2), 90 influenza A(H1N1)pdm09 and 58 influenza B) and 133 confirmed deaths have been reported since week 40 2016.



- USISS sentinel weekly hospitalised confirmed influenza cases, England (week 14)

- In week 14, there were four hospitalised confirmed influenza cases (two influenza A(H3N2), one influenza A(not subtyped) and one influenza B) reported through the USISS sentinel hospital network from 14 NHS Trusts across England (Figure 8), a rate of 0.08 per 100,000 compared to 0.19 per 100,000 in the previous week.

A total of 1,531 hospitalised confirmed influenza admissions (1,018 influenza A(H3N2), 418 influenza A(not subtyped), 69 influenza B and 26 influenza A(H1N1pdm09)) have been reported since week 40 2016.



- USISS Severe Respiratory Failure Centre confirmed influenza admissions, UK (week 14)

- In week 14, there were no confirmed influenza admissions reported from the six Severe Respiratory Failure (SRF) centres in the UK. There have been four confirmed influenza admissions (one influenza A(H3N2) and three influenza A(unknown subtype)) reported since week 40 2016.

All-cause mortality data

[| Back to top |](#)

In week 14 2017 in England, no statistically significant excess all-cause mortality by week of death was seen through the EuroMOMO algorithm in England. In the devolved administrations, no significant excess all-cause mortality was observed in week 14.

Seasonal mortality is seen each year in the UK, with a higher number of deaths in winter months compared to the summer. Additionally, peaks of mortality above this expected higher level typically occur in winter, most commonly the result of factors such as cold snaps and increased circulation of respiratory viruses, in particular influenza. Weekly mortality surveillance presented here aims to detect and report acute significant weekly excess mortality above normal seasonal levels in a timely fashion. Excess mortality is defined as a significant number of deaths reported over that expected for a given point in the year, allowing for weekly variation in the number of deaths. The aim is not to assess general mortality trends or precisely estimate the

excess attributable to different factors, although some end-of-winter estimates and more in-depth analyses (by age, geography etc.) are undertaken.

- All-cause death registrations, England and Wales

- In week 13 2017, an estimated 10,027 all-cause deaths were registered in England and Wales (source: [Office for National Statistics](#)). This is a decrease compared to the 10,325 estimated death registrations in week 12 2017.

- Excess all-cause mortality by age group, England, Wales, Scotland and Northern Ireland

- In week 14 2017 in England, no excess mortality by week of death above the upper 2 z-score threshold was seen overall, by age group or subnationally, after correcting ONS disaggregate data for reporting delay with the standardised [EuroMOMO](#) algorithm (Table 1). This data is provisional due to the time delay in registration; numbers may vary from week to week.

- In the devolved administrations, no significant excess mortality above the threshold was observed in week 14 (Table 2).

Table 1: Excess mortality by age group, England*

Age group (years)	Excess detected in week 14 2017?	Weeks with excess in 2016/17
<5	x	44
5-14	x	-
15-64	x	52-02
65+	x	45, 51-05

* Excess mortality is calculated as the observed minus the expected number of deaths in weeks above threshold

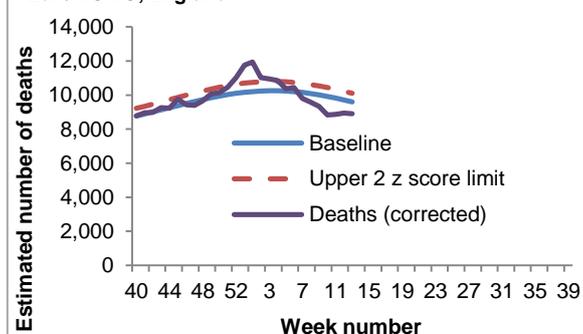
Table 2: Excess mortality by UK country, for all ages*

Country	Excess detected in week 14 2017?	Weeks with excess in 2016/17
England	x	52-05
Wales	x	03
Scotland	x	46,50,51,01,05
Northern Ireland	x	50-51,01-03,07-08

* Excess mortality is calculated as the observed minus the expected number of deaths in weeks above threshold

NB. Separate total and age-specific models are run for England which may lead to discrepancies between Tables 1 + 2

Figure 9: Excess mortality in all ages by week of death, EuroMOMO, England



Microbiological surveillance

[Back to top](#)

In week 14 2017, 11 samples tested positive for influenza (11 influenza B) through the UK GP sentinel schemes with an overall positivity of 28.9%. 47 positive detections were recorded through the DataMart scheme (17 influenza A(H3N2), two influenza A(H1N1)pdm09 and 28 influenza B) with a positivity of 3.5% in week 14.

- Sentinel swabbing schemes in England (RCGP) and the Devolved Administrations

-In week 14, 11 samples tested positive for influenza (11 influenza B) through the UK GP sentinel swabbing schemes, with an overall positivity of 28.9% compared to 12.7% in week 13 (Table 3).

Since week 40 2016, 932 samples (757 influenza A(H3N2), 60 influenza A(unknown subtype), 3 influenza A(H1N1)pdm09 and 112 influenza B) have tested positive for influenza through this scheme.

Table 3: Sentinel influenza surveillance in the UK

Week	England	Scotland	Northern Ireland	Wales
10	6/58 (10.3%)	13/36 (36.1%)	3/7 (-)	1/3 (-)
11	2/45 (4.4%)	10/53 (18.9%)	2/7 (-)	0/9 (-)
12	2/28 (7.1%)	10/43 (23.3%)	2/4 (-)	0/2 (-)
13	0/11 (-)	7/39 (18%)	0/1 (-)	0/4 (-)
14	0/4 (-)	5/23 (21.7%)	6/9 (-)	0/2 (-)

NB. Proportion positive omitted when fewer than 10 specimens tested

- Respiratory DataMart System (England)

In week 14 2017, out of the 1,355 respiratory specimens reported through the Respiratory DataMart System, 47 samples (3.5%) were positive for influenza (17 influenza A(H3N2), two influenza A(H1N1)pdm09 and 28 influenza B) (Figure 10), which is below the MEM threshold for this season of 8.6%. The highest positivity by age group was seen in the 65+ year olds (7.5%)(Figure 11). The overall positivity for RSV remained low at 0.7% in week 14. Positivity for rhinovirus positivity increased to 15.3% in week 14. Positivity for adenovirus remained at an increased level at 5.7% in week 14. Positivity for parainfluenza increased slightly from 7.6% in week 13 to 8.7% in week 14. Positivity for human metapneumovirus (hMPV) decreased to 1.5% in week 14.

Figure 10: DataMart samples positive for influenza, England

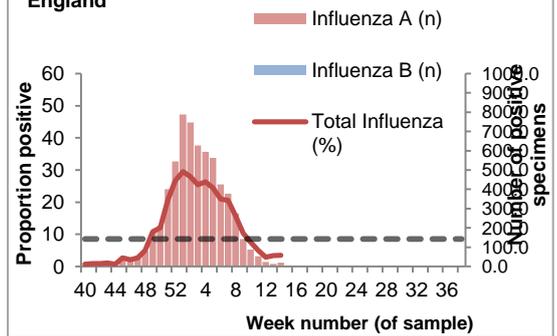


Figure 11: Datamart overall influenza % positive by age group, England

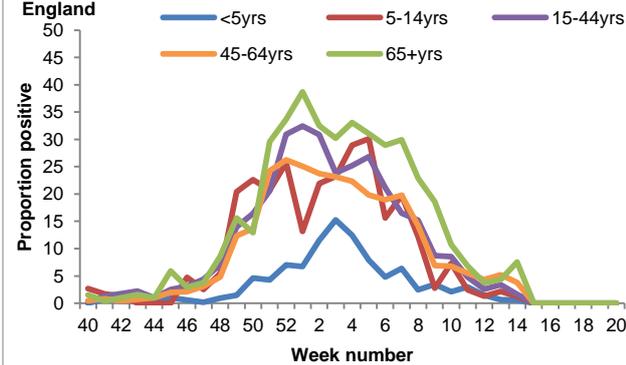
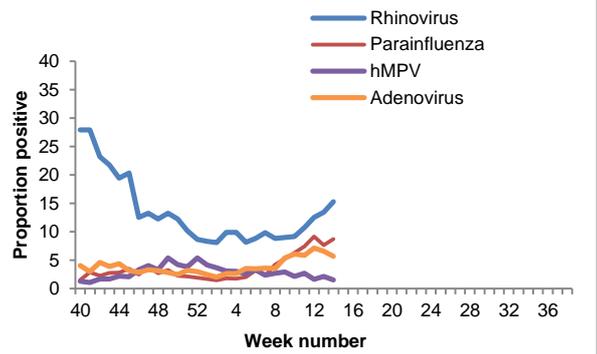


Figure 12: Datamart % positive for other respiratory viruses, England



**The Moving Epidemic Method has been adopted by the European Centre for Disease Prevention and Control to calculate thresholds for GP ILI consultations for the start of influenza activity in a standardised approach across Europe. The threshold to indicate a likelihood of influenza community circulation for Datamart % positive as calculated through the Moving Epidemic Method is 8.6% in 2016/17.*

- Virus characterisation

PHE characterises the properties of influenza viruses through one or more tests, including genome sequencing (genetic analysis) and haemagglutination inhibition (HI) assays (antigenic analysis). These data are used to compare how similar the currently circulating influenza viruses are to the strains included in seasonal influenza vaccines, and to monitor for changes in circulating influenza viruses. The interpretation of genetic and antigenic data sources is complex due to a number of factors, for example, not all viruses can be cultivated in sufficient quantity for antigenic characterisation, so that viruses with sequence information may not be able to be antigenically characterised as well. Occasionally, this can lead to a biased view of the properties of circulating viruses, as the viruses which can be recovered and analysed antigenically, may not be fully representative of majority variants, and genetic characterisation data does not always predict the antigenic characterisation

Since the start of the 2016/17 winter influenza season in week 40 2016, the PHE Respiratory Virus Unit has characterised four A(H1N1)pdm09 influenza viruses: one both genetically and antigenically and three antigenically. The A(H1N1)pdm09 virus genetically characterised belongs in the genetic subgroup 6B.1, which was the predominant genetic subgroup in the 2015/16 season. The three viruses antigenically analysed are similar to the A/California/7/2009 Northern Hemisphere 2016/17 (H1N1)pdm09 vaccine strain. Genetic characterisation of 318 A(H3N2) influenza viruses since week 40 showed that they all belong to genetic subclade 3C.2a, with 158 belonging to a cluster within this genetic subclade designated as 3C.2a1. The Northern Hemisphere 2016/17 influenza A(H3N2) vaccine strain A/HongKong/4801/2014 belongs in genetic subclade 3C.2a. This seasons A(H3N2) viruses are difficult to cultivate, and only 24 influenza A(H3N2) viruses have been isolated and antigenically characterised since week 40 2016, representing a minority of the detections, indicating the bias in antigenic data. The viruses antigenically analysed are similar to the A/HongKong/4801/2014 Northern Hemisphere 2016/17 A(H3N2) vaccine strain. Of the 24 antigenically characterised viruses, eight isolates have also been genetically characterised, with all belonging in genetic group 3C.2a, and six also belonging in the recently emerged 3C.2a1 cluster. 16 influenza B viruses have been analysed genetically since week 40/2015; 13 have been characterised as belonging to the B/Yamagata/16/88-lineage and 3 belonging to the B/Victoria/2/1987-lineage. Eighteen influenza B viruses have been isolated and antigenically characterised since week 40 2016. 13 viruses were characterised as belonging to the B/Yamagata/16/88-lineage and were antigenically similar to B/Phuket/3073/2013, the influenza B/Yamagata-lineage component of 2016/17 Northern Hemisphere quadrivalent vaccine. Five viruses were characterised as belonging to the B/Victoria/2/87-lineage and were antigenically similar to B/Brisbane/60/2008, the influenza B/Victoria-lineage component of 2016/17 Northern Hemisphere trivalent and quadrivalent vaccines.

- Antiviral susceptibility

Influenza positive samples are screened for mutations in the virus neuraminidase gene known to confer oseltamivir and/or zanamivir resistance. Additionally, testing of influenza A (H1N1)pdm09, A(H3N2), and influenza B virus isolates for neuraminidase inhibitor susceptibility (oseltamivir and zanamivir) is performed at PHE-RVU using a functional assay. The data summarized below combine the results of both testing methods. The samples tested are routinely obtained for surveillance purposes, but diagnostic testing of patients suspected to be infected with neuraminidase inhibitor-resistant virus is also performed.

Since week 40 2016, 305 influenza A(H3N2) have been tested for oseltamivir susceptibility; 300 are fully susceptible. 286 of the 305 were also tested for zanamivir susceptibility with 280 being fully susceptible. Four A(H3N2) viruses have been detected with an R292K amino acid substitution, which causes resistance to oseltamivir and a reduction in susceptibility to zanamivir, and one A(H3N2) virus with an E119V amino acid substitution was detected, causing resistance to oseltamivir but not tested for zanamivir susceptibility. All four R292K cases and the E119V case have been identified in patients with underlying medical conditions with some exposure to oseltamivir. Nine influenza A(H1N1)pdm09 and 17 influenza B (Yamagata) viruses have been tested for oseltamivir susceptibility and all were fully susceptible. One of the nine influenza A(H1N1)pdm09 virus and all 17 influenza B (Yamagata) virus have been tested for zanamivir susceptibility and all were fully susceptible.

- Antimicrobial susceptibility

-Table 4 shows in the 12 weeks up to 09 April 2017, the proportion of all lower respiratory tract isolates of *Streptococcus pneumoniae*, *Haemophilus influenzae*, *Staphylococcus aureus*, MRSA and MSSA tested and susceptible to antibiotics. These organisms are the key causes of community acquired pneumonia (CAP) and the choice of antibiotics reflects the British Thoracic Society empirical guidelines for management of CAP in adults.

Table 4: Antimicrobial susceptibility surveillance in lower respiratory tract isolates, 12 weeks up to 09 April 2017, E&W

Organism	Antibiotic	Specimens tested (N)	Specimens susceptible (%)
<i>S. pneumoniae</i>	Penicillin	3,995	90
	Macrolides	4,526	83
	Tetracycline	4,297	86
<i>H. influenzae</i>	Amoxicillin/ampicillin	19,574	69
	Co-amoxiclav	20,740	88
	Macrolides	7,663	10
	Tetracycline	20,439	98
<i>S. aureus</i>	Methicillin	6,749	90
	Macrolides	7,469	67
MRSA	Clindamycin	412	36
	Tetracycline	593	80
MSSA	Clindamycin	3,633	77
	Tetracycline	5,648	93

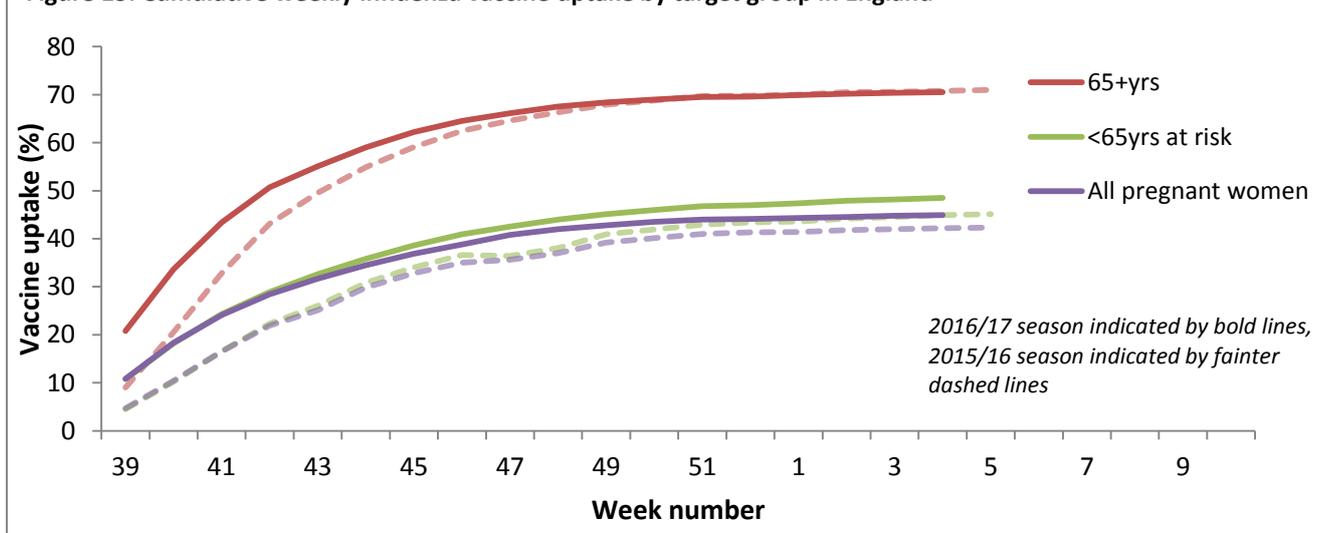
*Macrolides = erythromycin, azithromycin and clarithromycin

Vaccination

[| Back to top |](#)

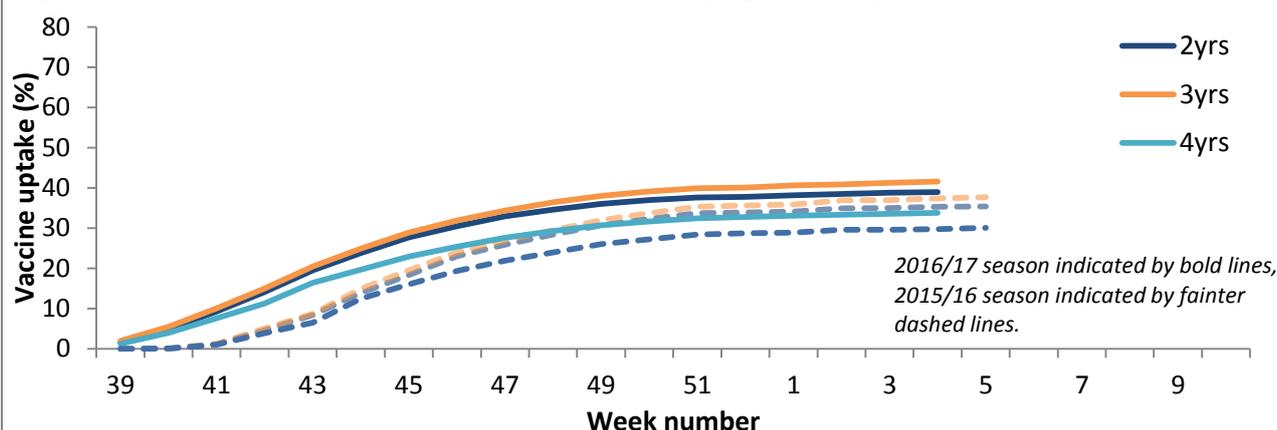
- Up to week 04 2017 in 85.0% of GP practices reporting weekly to Immform, the provisional proportion of people in England who had received the 2016/17 influenza vaccine in targeted groups was as follows, with vaccination activity starting earlier than last season (Figure 13):
 - 48.5% in under 65 years in a clinical risk group
 - 44.9% in pregnant women
 - 70.5% in 65+ year olds

Figure 13: Cumulative weekly influenza vaccine uptake by target group in England



- In 2016/17, all two-, three- and four-year-olds continue to be eligible for flu vaccination. In addition, the programme has been extended to children of school years 1, 2 and 3 age. Up to week 04 2017 in 88.1% of GP practices reporting weekly to Immform, the provisional proportion of children in England who had received the 2016/17 influenza vaccine in targeted groups was as follows (Figure 14):
 - 39.0% in all 2 year olds
 - 41.6% in all 3 year olds
 - 33.8% in all 4 year olds

Figure 14: Cumulative weekly influenza vaccine uptake by target group in England



- Provisional data from the fifth monthly collection of influenza vaccine uptake by frontline healthcare workers show 63.4% were vaccinated by 28 February 2017 from 98.9% of Trusts, compared to 50.8% vaccinated in the previous season by 29 February 2016. The report provides uptake at Trust level.
- Provisional data from the fourth monthly collection of influenza vaccine uptake in GP patients up to 31 January 2017 show that in 97.3% of all GP practices in England responding to the main GP survey, the proportion of people in England who received the 2016/17 influenza vaccine was as follows:
 - 48.7% in under 65 years in a clinical risk group
 - 44.8% in pregnant women
 - 70.4% in 65+ year olds
- Provisional data from the fourth monthly collection of influenza vaccine uptake in GP patients up to 31 January 2017 show that in 96.7% of all GP practices in England responding to the child GP survey, the proportion of people in England who received the 2016/17 influenza vaccine was as follows:
 - 38.9% in all 2 year olds
 - 41.5% in all 3 year olds
 - 33.9% in all 4 year olds
- Provisional data from the fourth monthly collection of influenza vaccine uptake for children of school years 1, 2 and 3 age (from a sample of 100% of all Local Authorities in England) show the proportion of children in England who received the 2016/17 influenza vaccine via school, pharmacy or GP practice by 31 January 2017 in targeted groups was as follows:
 - 57.6% in children of school Year 1 age (5-6 years)
 - 55.3% in children of school Year 2 age (6-7 years)
 - 53.3% in children of school Year 3 age (7-8 years)

International Situation

[Back to top](#)

Influenza activity in the temperate zone of the northern hemisphere continued to decrease. Worldwide, influenza A(H3N2) and influenza B viruses were predominant during this reporting period. In South Asia, influenza activity with mainly influenza A(H1N1) remained elevated.

- [Europe](#) updated on 07 April 2017 (Joint ECDC-WHO Influenza weekly update)

In week 13/2017, influenza activity across the region continued to decrease with all countries reporting low intensity of influenza activity. Of the 45 countries reporting on geographic spread, 3 reported widespread and 29 local or sporadic influenza activity indicating that influenza viruses are still circulating

This was the third week during the season that the proportion of type B viruses exceeded the proportion of type A viruses in sentinel detections. However, the overall number of type B virus detections remained low.

For week 13/2017, 94 (16%) of 598 sentinel specimens tested positive for influenza viruses. Of these, 90% were type B and 10% type A viruses. The proportion of type B viruses commonly increases in the second half of an influenza season. All subtyped influenza A viruses were A(H3N2). The lineage of 13 influenza B viruses was determined, of which 9 fell in B/Yamagata and 4 in B/Victoria lineages.

For week 13/2017, of the 11 countries that conduct sentinel surveillance of severe acute respiratory infection (SARI), 8 countries reported 581 SARI cases. Among these cases, 140 respiratory specimens were collected, 38 (27%) of which tested positive for influenza viruses.

For week 13/2017, 1,606 specimens from non-sentinel sources (such as hospitals, schools, non-sentinel primary care facilities, nursing homes and other institutions) tested positive for influenza viruses. Of these, 39% were type A (with 97% of the subtyped viruses being A(H3N2)), and 61% type B. The increase in proportion of type B viruses corresponds to the data seen in sentinel detections; however the number of B viruses detected remained low and similar to that seen in the previous 5 weeks.

The majority of participating European countries have had a marked excess in all-cause mortality since the end of 2016, in particular among the elderly aged 65 years and above. Currently, the mortality level seems to have decreased again. This season's excess mortality coincided with circulation of influenza A(H3N2), which usually leads to increased mortality among the elderly.

- [United States of America](#) updated on 07 April 2017 (Centre for Disease Control report)

During week 13, influenza activity decreased but remained elevated in the United States.

The most frequently identified influenza virus type reported by public health laboratories during week 13 was influenza B. The percentage of respiratory specimens testing positive for influenza in clinical laboratories decreased slightly.

A cumulative rate for the season of 57.2 laboratory-confirmed influenza-associated hospitalizations per 100,000 population was reported.

Nationwide during week 13, the proportion of outpatient visits for influenza-like illness (ILI) was 2.9%, which is above the national baseline of 2.2%.

- [Canada](#) updated on 07 April 2017 (Public Health Agency report)

Overall, influenza activity is slowly declining in Canada.

In week 13, all indicators (laboratory detections, influenza-like illness, outbreaks and hospitalizations) have either decreased or remained similar to the previous week.

Influenza activity due to influenza B is slowly increasing but is low compared to the same time period in the previous two seasons.

Influenza A activity is decreasing; however, influenza A continues to be the most common type of influenza affecting Canadians.

The majority of laboratory detections, hospitalizations and deaths have been among adults aged 65+ years

- [Global influenza update](#) updated on 03 April 2017 (WHO website)

Influenza activity in the temperate zone of the northern hemisphere continued to decrease. Worldwide, influenza A(H3N2) and influenza B viruses were predominant during this reporting period. In South Asia, influenza activity with mainly influenza A(H1N1) remained elevated.

In North America, overall influenza activity continued to decrease in Canada and United States of America, with influenza A(H3N2) virus predominating. In Mexico, influenza activity decreased slightly, but remained high, with influenza A(H1N1)pdm09 virus predominating.

In Europe, influenza activity continued to decrease to low levels in general, but especially in South Western Europe. In Northern Europe, some countries reported continued influenza activity, with influenza A (H3N2) and influenza B viruses. In some countries in Eastern Europe, influenza activity decreased but the proportion of influenza B virus detections increased in recent weeks.

In East Asia, low influenza activity was reported with influenza A(H3N2) virus predominant in the region.

In Western Asia, influenza activity continued to decrease with influenza B virus predominant in the region. In Armenia and Georgia, high levels of severe acute respiratory infection were reported in the recent weeks.

In Southern Asia, influenza activity continued to be reported in India, Maldives and Sri Lanka, with mainly influenza A(H1N1)pdm09 virus reported followed by influenza B virus. In South East Asia, influenza activity remained low.

In Northern Africa, low influenza activity was reported in Morocco and Tunisia, with influenza A(H3N2) and influenza B viruses circulating in the region.

In East and West Africa, low influenza activity was reported in the recent weeks, with influenza A(H1N1)pdm09, influenza A(H3N2) and influenza B viruses co-circulating.

In the Caribbean and Central America countries, influenza and other respiratory virus activity remained low in general.

In tropical South America, influenza and other respiratory virus activity remained low, although RSV activity remained elevated in Colombia. In the temperate zone of the Southern Hemisphere, influenza activity was at inter-seasonal levels.

Based on FluNet reporting, the WHO GISRS laboratories tested more than 132,143 specimens between 06 March 2017 and 19 March 2017. 23,560 were positive for influenza viruses, of which 15,164 (64.4%) were typed as influenza A and 8,396 (35.6%) as influenza B. Of the sub-typed influenza A viruses, 755 (15.1%) were influenza A(H1N1)pdm09 and 4247 (84.9%) were influenza A(H3N2). Of the characterized B viruses, 588 (77%) belonged to the B-Yamagata lineage and 176 (23%) to the B-Victoria lineage

The vaccine recommendation for the 2017-2018 northern hemisphere influenza season has been made. It is recommended that trivalent vaccines for use in the 2017-2018 northern hemisphere influenza season contain the following:

- an A/Michigan/45/2015 (H1N1)pdm09-like virus;
- an A/Hong Kong/4801/2014 (H3N2)-like virus; and
- a B/Brisbane/60/2008-like virus.

It is recommended that quadrivalent vaccines containing two influenza B viruses contain the above three viruses and a B/Phuket/3073/2013-like virus. The full report can be found [here](#).

- [Avian Influenza](#) latest update on 05 April 2017 (WHO website)

Influenza A(H5) viruses

Between [14 February and 16 March 2017](#), two new laboratory-confirmed human case of influenza A(H5N1) virus infection was reported to WHO from Egypt. Avian influenza A(H5N1) viruses are enzootic in poultry in Egypt.

Since 2003, a total of 858 laboratory-confirmed cases of human infection with avian influenza A(H5N1) virus, including 453 deaths, have been reported to WHO from 16 countries.

Influenza A(H5) subtype viruses have the potential to cause disease in humans and thus far, no human cases, other than those with influenza A(H5N1) and A(H5N6) viruses, have been reported to WHO. According to reports received by the World Organisation for Animal Health (OIE), various influenza A(H5) subtypes continue to be detected in birds in West Africa, Europe and Asia. There have also been numerous detections of influenza A(H5N8) viruses in wild birds and domestic poultry in several countries in Africa, Asia and Europe since June 2016, and influenza A(H5N5) in wild birds in Europe. For more information on the background and public health risk of these viruses, please see the WHO assessment of risk associated with influenza A(H5N8) virus [here](#).

Influenza A(H7N9)

On [31 March 2017](#), the National Health and Family Planning Commission of China (NHFPC) notified WHO of 17 additional laboratory-confirmed cases of human infection with avian influenza A(H7N9) virus in mainland China.

On [24 March 2017](#), the National Health and Family Planning Commission of China (NHFPC) notified WHO of 18 additional laboratory-confirmed cases of human infection with avian influenza A(H7N9) virus in mainland China.

A total of 1,364 laboratory-confirmed human infections with avian influenza A (H7N9) virus, including at least 489 deaths, have been reported through IHR notification since early 2013.

- [Middle East respiratory syndrome coronavirus \(MERS-CoV\)](#) latest update on 04 April 2017

On [21 March 2017](#) the national IHR focal point of Qatar reported one additional case of Middle East Respiratory Syndrome Coronavirus (MERS-CoV).

Between [23 February and 16 March 2017](#) the national IHR focal point of Saudi Arabia reported 18 additional cases of Middle East Respiratory Syndrome (MERS) including two fatal cases. Four deaths among previously reported MERS cases were also reported (case numbers 1, 4, 5 and 7 in the Disease Outbreak News published on 10 March 2017).

Up to 12 April 2017, a total of four cases of Middle East respiratory syndrome coronavirus, MERS-CoV, (two imported and two linked cases) have been confirmed in the UK. On-going surveillance has identified 950 suspect cases in the UK that have been investigated for MERS-CoV and tested negative.

Globally, since September 2012, WHO has been notified of 1,936 laboratory-confirmed cases of infection with MERS-CoV, including at least 690 related deaths. Further information on management and guidance of possible cases is available [online](#). The latest ECDC MERS-CoV risk assessment can be found [here](#), where it is highlighted that risk of widespread transmission of MERS-CoV remains low.

Acknowledgements

[| Back to top |](#)

This report was prepared by the Influenza section, Respiratory Diseases Department, Centre for Infectious Disease Surveillance and Control, Public Health England. We are grateful to all who provided data for this report including the RCGP Research and Surveillance Centre, the PHE Real-time Syndromic Surveillance team, the PHE Respiratory Virus Unit, the PHE Modelling and Statistics unit, the PHE Dept. of Healthcare Associated Infection & Antimicrobial Resistance, PHE regional microbiology laboratories, Office for National Statistics, the Department of Health, Health Protection Scotland, National Public Health Service (Wales), the Public Health Agency Northern Ireland, the Northern Ireland Statistics and Research Agency, QSurveillance[®] and EMIS and EMIS practices contributing to the QSurveillance[®] database.

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[| Back to top |](#)

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- Wales surveillance ([Public Health Wales](#))
- [Real time syndromic surveillance](#)
- MEM threshold [methodology paper](#) and [UK pilot paper](#)

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- [FluSurvey](#)
- [MOSA](#)

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- [USISS](#) system
- [EuroMOMO](#) mortality project

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- Seasonal influenza vaccine programme ([Department of Health Book](#))
- Childhood flu programme information for healthcare practitioners ([Public Health England](#))
- 2016/17 Northern Hemisphere seasonal influenza vaccine recommendations ([WHO](#))