Weekly national Influenza and COVID-19 surveillance report

Week 46 report (up to week 45 data)
12 November 2020
Executive summary

This report summarises the information from the surveillance systems which are used to monitor Coronavirus Disease 2019 (COVID-19), influenza, and other seasonal respiratory viruses in England. References to COVID-19 represent the disease name and SARS-CoV-2 represent the virus name. The report is based on data from week 45 (between 2 and 8 November 2020) and for some indicators daily data up to 10 November 2020.

Surveillance indicators suggest that COVID-19 activity at a national level has remained high during week 45. There is currently limited testing for other respiratory viruses, however, laboratory indicators suggest that influenza activity is low.

Week 44 was school half term in many parts of the country. During week 45 several social and physical distancing measures were re-introduced across England and mass testing was introduced in Liverpool. All of these factors are likely to impact on surveillance indicators.

Detections of COVID-19 cases in England remained high in week 45 with a slight increase on the previous week. However, overall positivity rates for both Pillars 1 and 2 decreased slightly. Incidence and positivity rates remain highest in the North of England. By age group, cases rates were highest in the 20 to 29 year olds. Positivity rates were highest in the 80+ year olds tested through Pillar 1 (NHS and PHE testing) and in the 10 to 19 year olds tested through Pillar 2 (community testing).

Through Respiratory Datamart, there were no influenza positive sample detected in week 45. Rhinovirus activity remains high but is decreasing in week 45.

The overall number of acute respiratory infection incidents reported to PHE Health Protection Teams have increased slightly from 1110 in the previous week to 1140 in week 45 in England. This mainly reflects an increase in incidents reported in care homes. It is important to note that an increasing number of outbreaks are being managed through other routes outside of Health Protection Teams. In the majority of reported incidents SARS-CoV-2 has been detected.

The majority of community and syndromic indicators decreased or remained stable during week 45. General practice (GP) influenza-like illness (ILI) consultations remained low in all UK schemes.

Through the UK GP swabbing scheme, SARS-CoV-2 positivity among patients contacting their GP with influenza like illness or lower respiratory tract infection symptoms increased from 16.3% in week 44 to 23.0% in week 45.

The overall COVID-19 confirmed hospital and ICU/HDU admission rates increased slightly whilst the influenza confirmed hospital and ICU/HDU admission rates remained low.

Emergency department attendances for COVID-19 like diagnosis have increased further in week 45 whilst those for acute respiratory infections remained stable.

The number of COVID-19 confirmed deaths was similar to the previous week. Overall excess all-cause mortality was observed in week 44; by age group in the 85+ year olds and subnationally in the North West and Yorkshire and Humber.
Overall estimated national seroprevalence based on blood donor samples was 5.9% with the highest seroprevalence by region seen in London and by age group in young adults.

Influenza vaccine uptake is the highest it has ever been at this point in the season for those aged 65+ and in 2 and 3 year olds. For those in at-risk groups uptake is higher than last season and comparable to seasons before that. For pregnant women uptake is lower than in previous seasons. Weekly vaccine coverage data are provisional. The weekly 2020/21 pregnant women and at-risk denominators are undergoing validation checks and so reported coverage in these groups may be underestimated.
# Contents

Executive summary ........................................................................................................ 2
Contents ......................................................................................................................... 2
Laboratory surveillance .................................................................................................. 6
  Confirmed COVID-19 cases (England) ........................................................................ 6
  Respiratory DataMart system (England) ..................................................................... 17
Community surveillance ................................................................................................. 19
  Acute respiratory infection incidents ........................................................................ 19
  COVID-19 cases by type of residence ....................................................................... 26
  Medical Officers of Schools Association (MOSA) & PHE surveillance scheme ........ 27
  FluSurvey ................................................................................................................... 28
  Google search queries ............................................................................................... 29
  NHS 111 ..................................................................................................................... 30
Primary care surveillance ................................................................................................. 34
  RCGP (England) ......................................................................................................... 34
  UK ............................................................................................................................... 36
  GP In Hours, Syndromic Surveillance ........................................................................ 37
  GP Out of Hours, Syndromic Surveillance ................................................................ 39
  Sentinel swabbing scheme in England and the Devolved Administrations .............. 41
Secondary care surveillance ............................................................................................ 42
  SARI Watch ............................................................................................................... 42
  Hospitalisations, SARI Watch .................................................................................. 43
  ICU/HDU admissions, SARI Watch ........................................................................... 47
  ECMO, SARI Watch .................................................................................................. 51
  Emergency Department attendances, Syndromic surveillance ................................ 52
Mortality surveillance ...................................................................................................... 54
  Cumulative COVID-19 deaths .................................................................................. 54
  Daily excess all-cause mortality (England) ................................................................. 59
Microbiological surveillance ........................................................................................... 62
  Virus characterisation ............................................................................................... 62
Antiviral susceptibility ........................................................................................................62
Antimicrobial susceptibility .................................................................................................63
COVID-19 sero-prevalence surveillance .............................................................................64
Influenza vaccination ...........................................................................................................67
  Influenza vaccine uptake in GP patients ........................................................................67
  Influenza vaccine uptake in school age children ............................................................69
  Influenza vaccine uptake in healthcare workers .............................................................69
International update ..........................................................................................................70
  Global COVID-19 update ...............................................................................................70
  Global influenza update ................................................................................................72
Other respiratory viruses .....................................................................................................74
Related links .........................................................................................................................75
Laboratory surveillance

Confirmed COVID-19 cases (England)

As of 09:00 on 10 November 2020, a total of 1,053,330 have been confirmed positive for COVID-19 in England under Pillars 1 and 2.

Overall case numbers remained high in week 45. Overall positivity in both Pillars 1 and 2 have decreased slightly. The highest case rates were seen in the 20 to 29 year olds in Pillars 1 and 2. The highest positivity rates were noted in the 80+ year olds in Pillar 1 and in the 10 to 19 year olds in Pillar 2. Cases rates and positivity continue to be highest in the North of England.

From the week 42 report onwards, case rates in Figures 3,4,7 and 9 have been calculated using mid-2019 ONS population estimates.

**Figure 1: Laboratory confirmed COVID-19 cases tested under Pillar 1 and Pillar 2, based on sample week with overall weekly positivity for Pillars 1 and 2 (%)**

*The data are shown by the week the specimen was taken from the person being tested. This gives the most accurate analysis of this time progression, however, for the most recent week results for more samples are expected therefore this should be interpreted with caution.*
*Positivity data was previously deduplicated across the course of the pandemic to prevent persistent infections being counted as new cases. Since week 40, positivity is calculated as the number of individuals testing positive during the week divided by the number of individuals tested during the week. This approach accounts for the increasing number of individuals who will have been tested multiple times as the pandemic progresses.


Age and sex

Figure 2: Age/sex pyramids for laboratory confirmed COVID-19 cases tested under Pillars 1 and 2 (a) cumulative number since week 27 (n=799,651), and (b) in weeks 44 and 45 (n=263,503)

(a)

(b)
Figure 3: Weekly laboratory confirmed COVID-19 case rates per 100,000, tested under Pillar 1 and Pillar 2, by sex

Figure 4: Weekly laboratory confirmed COVID-19 case rates per 100,000, tested under Pillar 1 and Pillar 2, by age group
Figure 5: Weekly positivity (%) of laboratory confirmed COVID-19 cases tested overall and by sex under (a) Pillar 1 and (b) Pillar 2, (SGSS and Respiratory DataMart)
Figure 6: Weekly positivity (%) of laboratory confirmed COVID-19 cases tested under Pillar 1, (a) by male and age group and (b) by female and age group and; under Pillar 2, (c) by male and age group and (d) by female and age group, (SGSS and Respiratory DataMart)

(a) Pillar 1 - Male

(b) Pillar 1 - Female
Weekly National Influenza & COVID-19 Report: week 46 report (up to week 45 data)

(c) Pillar 2 - Male

(d) Pillar 2 - Female
Geography

Table 1: Cumulative number of cases under Pillars 1 and 2 (n=1,036,248) and cumulative number of cases since week 27 under Pillar 1 and 2 (n=801,300)

<table>
<thead>
<tr>
<th>PHE Centres</th>
<th>Cumulative Pillar 1 + 2 cases</th>
<th>Cumulative since week 27, Pillar 1 + 2 cases</th>
</tr>
</thead>
<tbody>
<tr>
<td>North East</td>
<td>74,234</td>
<td>59,195</td>
</tr>
<tr>
<td>North West</td>
<td>253,107</td>
<td>210,863</td>
</tr>
<tr>
<td>Yorkshire and Humber</td>
<td>163,903</td>
<td>135,207</td>
</tr>
<tr>
<td>West Midlands</td>
<td>115,294</td>
<td>90,151</td>
</tr>
<tr>
<td>East Midlands</td>
<td>101,244</td>
<td>80,598</td>
</tr>
<tr>
<td>East of England</td>
<td>66,466</td>
<td>42,340</td>
</tr>
<tr>
<td>London</td>
<td>116,883</td>
<td>83,231</td>
</tr>
<tr>
<td>South East</td>
<td>90,583</td>
<td>57,859</td>
</tr>
<tr>
<td>South West</td>
<td>54,534</td>
<td>41,856</td>
</tr>
</tbody>
</table>

Figure 7: Weekly laboratory confirmed COVID-19 case rates per 100,000 population tested under Pillar 1 and Pillar 2, by PHE Centres and sample week
Figure 8: Weekly positivity of laboratory confirmed COVID-19 cases tested under (a) Pillar 1 (%) and (b) Pillar 2 (%), by PHE Centres and sample week, (SGSS and Respiratory DataMart)
Figure 9: Weekly rate of COVID-19 cases per 100,000 population tested under Pillars 1 and 2, by upper-tier local authority, England (box shows enlarged map of London area)
Ethnicity

Figure 10: Weekly incidence per 100,000 population by ethnicity, England

*the incidence rates on Figure 10 have been calculated using the mid-2018 ONS population estimates
Respiratory DataMart system (England)

The Respiratory Datamart system was initiated during the 2009 influenza pandemic to collate all laboratory testing information in England. It is now used as a sentinel laboratory surveillance tool, monitoring all major respiratory viruses in England. 16 laboratories in England will be reporting data for this season. As this is based on a sample of labs - SARS-CoV-2 positivity figures quoted here will differ from those quoted in the Confirmed COVID-19 cases section, however, they are included to facilitate comparison with data on other respiratory viruses.

In week 45 2020, out of the 97,718 respiratory specimens reported through the Respiratory DataMart System (based on data received from 12 out of 16 laboratories), 5,214 samples were positive for SARS-CoV-2 with an overall positivity of 5.3%. The highest positivity was noted in the 65+ year olds at 7.2% in week 45. The overall influenza positivity was low at 0.0% in week 45, with no sample testing positive for influenza (out of 939 tested) (Figure 11).

Rhinovirus positivity decreased slightly at 13.6% in week 45 compared to 18.1% in the previous week (Figure 12). The highest positivity by age group for rhinovirus was noted in the less than 5 year olds in week 45 (Figure 13). Respiratory syncytial virus (RSV), adenovirus, parainfluenza and human metapneumovirus (hMPV) positivity all remained low at 0.0%, 2.2%, 0.0% and 0.4% respectively in week 45 (Figure 12).

Figure 11: DataMart samples positive for influenza and weekly positivity (%) for influenza and SARS-CoV-2, England
Figure 12: DataMart weekly positivity (%) for other respiratory viruses, England

Figure 13: DataMart weekly positivity (%) for rhinovirus by age, England
Community surveillance

Acute respiratory infection incidents

Information on acute respiratory infection (ARI) incidents is based on situations reported to PHE Health Protection Teams (HPTs). These include:

- confirmed outbreaks of acute respiratory infections ie two or more laboratory confirmed cases (SARS-CoV-2, influenza or other respiratory pathogens) linked to a particular setting
- setting situations where an outbreak is suspected

All suspected outbreaks are further investigated by the HPT in liaison with local partners and a significant proportion do not meet the criteria of a confirmed outbreak. For example if suspected cases test negative for COVID19 or other respiratory pathogens, or cases are subsequently found not to have direct links to the setting. Since Pillar 2 testing became open to everyone during week 21 more incidents of mild disease have been detected in settings with healthy young populations.

Processes for reporting ARI incidents vary between PHE Centres.

The situations captured on HPZone represent a subset of all ongoing clusters and outbreaks in England rather than an exhaustive listing. A variety of arrangements are in place with local authorities and other stakeholders supporting HPTs, however data are not routinely documented on HPZone. As a result, the number of outbreaks reported for some of the regions are underestimates.

The denominator (the overall number of settings in each category) will differ by the setting category, for example there are fewer hospitals than workplaces, as will the propensity to report incidents to PHE. Therefore these data are more useful for monitoring trends over time than making comparisons across setting categories.

The number of incidents in each setting with at least one laboratory confirmed case of COVID19 are reported below. A national school helpline started operating on 17 September 2020 and a Universities helpline started operating on 7 October. This is likely to have had an impact on the number of situations/outbreaks being reported to HPTs in these settings

1226 new ARI incidents have been reported in week 45 in the UK (Figure 14):

- 442 incidents were from care homes where 309 had at least one linked case that tested positive for SARS-CoV-2
• 88 incidents were from hospitals where 72 had at least one linked case that tested positive for SARS-CoV-2
• 188 incidents were from educational settings where 146 had at least one linked case that tested positive for SARS-CoV-2
• 6 incidents were from prisons where 5 had at least one linked case that tested positive for SARS-CoV-2
• 275 incidents were from workplace settings where 182 had at least one linked case that tested positive for SARS-CoV-2
• 10 incidents were from food outlet/restaurant settings where 8 had at least one linked case that tested positive for SARS-CoV-2
• 217 incidents were from the other settings category where 151 had at least one linked case that tested positive for SARS-CoV-2

Figure 14: Number of acute respiratory infection (ARI) incidents by institution, UK

*excludes data from Wales
Figure 15: Number of acute respiratory infection (ARI) incidents by institution, England

Figure 16: Number of acute respiratory infection (ARI) incidents in care homes by virus type from week 27, England
Figure 17: Number of acute respiratory infection (ARI) incidents in hospitals by virus type from week 27, England

![Hospital chart showing number of ARI incidents by virus type from week 27 to week 53.]

Figure 18: Number of acute respiratory infection (ARI) incidents in educational settings by virus type from week 27, England

![Educational settings chart showing number of ARI incidents by virus type from week 27 to week 53.]

---

Weekly National Influenza & COVID-19 Report: week 46 report (up to week 45 data)
Figure 19: Number of acute respiratory infection (ARI) incidents in prisons by virus type from week 27, England

Prisons

<table>
<thead>
<tr>
<th>Virus Type</th>
<th>Number of ARI Incidents</th>
</tr>
</thead>
<tbody>
<tr>
<td>Influenza A</td>
<td></td>
</tr>
<tr>
<td>Influenza B</td>
<td></td>
</tr>
<tr>
<td>SARS-CoV-2</td>
<td></td>
</tr>
<tr>
<td>Rhinovirus</td>
<td></td>
</tr>
<tr>
<td>RSV</td>
<td></td>
</tr>
<tr>
<td>Other respiratory viruses</td>
<td></td>
</tr>
<tr>
<td>No organism reported</td>
<td></td>
</tr>
</tbody>
</table>

Date of report week

Figure 20: Number of acute respiratory infection (ARI) incidents in workplace settings by virus type from week 27, England

Workplace settings

<table>
<thead>
<tr>
<th>Virus Type</th>
<th>Number of ARI Incidents</th>
</tr>
</thead>
<tbody>
<tr>
<td>Influenza A</td>
<td></td>
</tr>
<tr>
<td>Influenza B</td>
<td></td>
</tr>
<tr>
<td>SARS-CoV-2</td>
<td></td>
</tr>
<tr>
<td>Rhinovirus</td>
<td></td>
</tr>
<tr>
<td>RSV</td>
<td></td>
</tr>
<tr>
<td>Other respiratory viruses</td>
<td></td>
</tr>
<tr>
<td>No organism reported</td>
<td></td>
</tr>
</tbody>
</table>

Date of report week
Figure 21: Number of acute respiratory infection (ARI) incidents in food outlet/restaurants settings by virus type from week 27, England

Figure 22: Number of acute respiratory infection (ARI) incidents in other settings settings by virus type from week 27, England
Table 2: Total number of situations/incidents by institution and PHE Centres over the past four weeks with the total number in the last week in brackets

<table>
<thead>
<tr>
<th>PHE Centres</th>
<th>Care home</th>
<th>Hospital</th>
<th>Educational settings</th>
<th>Prisons</th>
<th>Workplace settings</th>
<th>Food outlet/restaurant settings</th>
<th>Other settings</th>
<th>Total</th>
</tr>
</thead>
<tbody>
<tr>
<td>East of England</td>
<td>122(27)</td>
<td>18(3)</td>
<td>26(4)</td>
<td>2(0)</td>
<td>48(13)</td>
<td>2(0)</td>
<td>29(9)</td>
<td>247(56)</td>
</tr>
<tr>
<td>East Midlands</td>
<td>130(36)</td>
<td>50(4)</td>
<td>153(40)</td>
<td>5(3)</td>
<td>112(39)</td>
<td>10(1)</td>
<td>45(13)</td>
<td>505(136)</td>
</tr>
<tr>
<td>London</td>
<td>78(26)</td>
<td>66(25)</td>
<td>165(29)</td>
<td>7(0)</td>
<td>203(36)</td>
<td>19(2)</td>
<td>45(15)</td>
<td>583(133)</td>
</tr>
<tr>
<td>North East</td>
<td>93(43)</td>
<td>2(0)</td>
<td>9(0)</td>
<td>0(0)</td>
<td>23(2)</td>
<td>0(0)</td>
<td>62(20)</td>
<td>189(65)</td>
</tr>
<tr>
<td>North West</td>
<td>222(51)</td>
<td>26(5)</td>
<td>53(2)</td>
<td>3(1)</td>
<td>98(19)</td>
<td>6(1)</td>
<td>188(34)</td>
<td>596(113)</td>
</tr>
<tr>
<td>South East</td>
<td>155(47)</td>
<td>30(9)</td>
<td>232(42)</td>
<td>4(1)</td>
<td>132(38)</td>
<td>22(3)</td>
<td>84(25)</td>
<td>659(165)</td>
</tr>
<tr>
<td>South West</td>
<td>169(51)</td>
<td>7(4)</td>
<td>101(14)</td>
<td>3(0)</td>
<td>97(19)</td>
<td>17(2)</td>
<td>58(13)</td>
<td>452(103)</td>
</tr>
<tr>
<td>West Midlands</td>
<td>146(51)</td>
<td>71(23)</td>
<td>159(28)</td>
<td>3(1)</td>
<td>261(58)</td>
<td>14(1)</td>
<td>142(33)</td>
<td>796(195)</td>
</tr>
<tr>
<td>Yorkshire and Humber</td>
<td>201(65)</td>
<td>11(4)</td>
<td>122(20)</td>
<td>0(0)</td>
<td>173(49)</td>
<td>8(0)</td>
<td>145(36)</td>
<td>660(174)</td>
</tr>
<tr>
<td><strong>Total</strong></td>
<td>1316(397)</td>
<td>281(77)</td>
<td>1020(179)</td>
<td>27(6)</td>
<td>1147(273)</td>
<td>98(10)</td>
<td>798(198)</td>
<td>4687(1140)</td>
</tr>
</tbody>
</table>
COVID-19 cases by type of residence

Table 3 shows the proportion of confirmed COVID-19 cases according to their type of residence. Property classifications are derived from Ordnance Survey AddressBase and are matched to address details within the laboratory data. Properties are identified by unique property reference number (UPRN) and basic land property unit (BLPU). Cases with poor or no address data which failed the address matching and are classed as ‘undetermined’. No fixed abode and overseas addresses identified by recording in the laboratory data.

In week 44, the highest percentage of confirmed COVID-19 cases by type of residence was seen in residential dwelling (Table 3).

Week 45 data is not available for this week’s report.

Table 3: Type of residence of confirmed COVID-19 cases by percentage of total weekly cases

<table>
<thead>
<tr>
<th>Type of residence</th>
<th>week 39</th>
<th>week 40</th>
<th>week 41</th>
<th>week 42</th>
<th>week 43</th>
<th>week 44</th>
</tr>
</thead>
<tbody>
<tr>
<td>Residential dwelling (including houses, flats, sheltered accommodation)</td>
<td>79.3</td>
<td>78.1</td>
<td>79.5</td>
<td>81.3</td>
<td>81.4</td>
<td>85.8</td>
</tr>
<tr>
<td>Undetermined</td>
<td>16.2</td>
<td>15.2</td>
<td>14.4</td>
<td>13.3</td>
<td>14.2</td>
<td>10.0</td>
</tr>
<tr>
<td>Care/Nursing home</td>
<td>1.4</td>
<td>1.5</td>
<td>1.4</td>
<td>1.7</td>
<td>1.8</td>
<td>2.1</td>
</tr>
<tr>
<td>Residential institution (including residential education)</td>
<td>1.1</td>
<td>2.4</td>
<td>2.3</td>
<td>1.6</td>
<td>0.9</td>
<td>0.6</td>
</tr>
<tr>
<td>House in multiple occupancy (HMO)</td>
<td>1.0</td>
<td>1.6</td>
<td>0.9</td>
<td>0.8</td>
<td>0.6</td>
<td>0.6</td>
</tr>
<tr>
<td>Medical facilities (including hospitals and hospices, and mental health)</td>
<td>0.3</td>
<td>0.3</td>
<td>0.3</td>
<td>0.3</td>
<td>0.2</td>
<td>0.2</td>
</tr>
<tr>
<td>Other property classifications</td>
<td>0.5</td>
<td>0.7</td>
<td>0.9</td>
<td>0.8</td>
<td>0.7</td>
<td>0.6</td>
</tr>
<tr>
<td>Prisons, detention centres, secure units</td>
<td>0.2</td>
<td>0.2</td>
<td>0.1</td>
<td>0.1</td>
<td>0.2</td>
<td>0.2</td>
</tr>
<tr>
<td>No fixed abode</td>
<td>0.0</td>
<td>0.0</td>
<td>0.0</td>
<td>0.0</td>
<td>0.0</td>
<td>0.0</td>
</tr>
<tr>
<td>Overseas address</td>
<td>0.0</td>
<td>0.0</td>
<td>0.0</td>
<td>0.0</td>
<td>0.0</td>
<td>0.0</td>
</tr>
</tbody>
</table>
Medical Officers of Schools Association (MOSA) & PHE surveillance scheme

Boarding schools in England within the MOSA network are recruited each season to report various respiratory related illnesses including influenza like illnesses (ILI).

Data will be reported from week 46.

If you are a MOSA school and would like to participate in this scheme, please email mosa@phe.gov.uk for more information.
FluSurvey

An internet based surveillance system has been developed based on FluSurvey. FluSurvey is a web tool survey designed to monitor trends of influenza like illness (ILI) in the community using self-reported respiratory symptoms from registered participants. The platform has been adapted to capture respiratory symptoms, exposure risk and healthcare seeking behaviours among registered participants to contribute to national surveillance of COVID-19 activity as well as influenza activity since week 44.

A total of 3,490 participants completed the weekly COVID-19 surveillance survey in week 45, of which 116 (3.3%) reported fever or cough and 59 (1.7%) reporting ILI. The most commonly reported method of access to healthcare services continue to be through telephoning a GP practice in week 45 (Figure 23).

Figure 23: Rate of contact with different healthcare services among FluSurvey participants reporting fever or cough symptoms, England

![Rate of contact with different healthcare services](image-url)
Google search queries

This is a web-based syndromic surveillance system which uses daily search query frequency statistics obtained from the Google Health Trends API [1]. This model focuses on search queries about COVID-19 symptoms as well as generic queries about “coronavirus” (e.g. “covid-19”). The search query frequency time series has been weighted based on symptom frequency as reported in other data sources. Frequency of searches for symptoms is compared with a baseline calculated from historical daily data.

During week 44, the overall and media-debiasing weighted Google search scores decreased further (Figure 24).

Week 45 data is not available for this week’s report.

**Figure 24: Normalised Google search score for COVID-19 symptoms, with weighted score for media-debiasing and historical trend, England**
The NHS 111 service monitors daily trends in phone calls made to the service in England, to capture trends in infectious diseases such as influenza and norovirus.

Up to 91 November 2020, the daily percentage of NHS 111 ‘potential COVID-19-like’ calls (as a percentage of total NHS 111 calls) and the number of online assessments remained stable. The daily percentage of cold/flu calls (as a percentage of total NHS 111 calls) and cold/flu completed online assessments remained stable (Figure 25 and 26). The daily percentage of loss of taste or smell calls increased slightly whereas online assessments remained stable.

Please note that NHS 111 callers (from 11 May 2020) and NHS 111 online users (from 11 June 2020), who are assessed as having probable COVID-19 symptoms are now triaged using symptom specific pathways e.g. cold/flu, which are included in routine syndromic indicators.

Further information about these caveats is available from the PHE Remote Health Advice Syndromic Surveillance bulletin.

Figure 25: NHS 111 telephony indicators (and 7-day moving average) for (a) daily potential COVID-19 calls, (b) daily cold/flu calls and (c) daily loss of taste or smell calls, as a percentage of total calls for all ages, England
Figure 26: NHS 111 completed online assessments (and 7-day moving average) for (a) daily potential COVID-19 online assessments, (b) daily cold/flu online assessments and (c) daily loss of taste or smell online assessments, as the number of completed online assessments for all ages, England.
Primary care surveillance

RCGP (England)

The weekly ILI consultation rate through the RCGP surveillance was 1.5 per 100,000 registered population in participating GP practices in week 45 compared to the 1.4 per 100,000 in the previous week. This is below the baseline threshold (12.2 per 100,000) (Figure 27). By age group, the highest rates were seen in the under 1 year olds (6.1 per 100,000) and in the 45 to 64 year olds (1.9 per 100,000). The Lower Respiratory Tract Infections (LRTI) consultation rate was at 19.5 per 100,000 in week 45, which was similar to from the rate of 19.6 per 100,000 from the previous week. The COVID-19-like indicator consultation rate decreased at 46.7 per 100,000 in week 45 compared to 101.5 per 100,000 in the previous week (Figure 28).

Figure 27: RCGP ILI consultation rates, all ages, England
Figure 28: RCGP ILI, LRTI and COVID-19-like indicator consultation rates, England
Overall, weekly ILI consultations rates were below baseline levels in all UK schemes (Table 4).

By age group, the highest rates were seen in the 45 to 64 year olds in Scotland (1.3 per 100,000), in the 45 to 64 year olds in Wales (1.9 per 100,000) and in the 45 to 64 year olds in Northern Ireland (3.7 per 100,000).

Table 4: GP ILI consultations in the UK for all ages with MEM thresholds applied

<table>
<thead>
<tr>
<th>GP ILI consultation rates (all ages)</th>
<th>Week number</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>40</td>
</tr>
<tr>
<td>England (RCGP)</td>
<td>2.1</td>
</tr>
<tr>
<td>Wales</td>
<td>1.0</td>
</tr>
<tr>
<td>Scotland</td>
<td>0.5</td>
</tr>
<tr>
<td>Northern Ireland</td>
<td>1.3</td>
</tr>
</tbody>
</table>

The Moving Epidemic Method (MEM) 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 (based on 10 seasons excluding 2009/10), in a standardised approach across Europe. For MEM threshold values for each country, please visit: https://www.gov.uk/guidance/sources-of-uk-flu-data-influenza-surveillance-in-the-uk#clinical-surveillance-through-primary-care
GP In Hours, Syndromic Surveillance

The GP In Hours (GPIH) syndromic surveillance system monitors the number of GP visits during regular hours of known clinical indicators.

Up to 9 November 2020, GPIH consultations for potential COVID-19-like consultations and ILI consultations remained stable (Figure 29). Please note that the GPIH COVID-19-like indicator presented in this report is derived from a reduced denominator population, compared to ILI.

Please note GP data should be interpreted with caution due to changes in advice regarding accessing GP surgeries due to COVID-19. Further information about these caveats is available from the PHE GP In Hours Syndromic Surveillance bulletin.

Figure 29: GPIH clinical indicators for (a) potential COVID-19 GP consultations and (b) influenza-like illness GP consultations, England

(a)
GP Out of Hours, Syndromic Surveillance

The GP Out of Hours (GPOOH) syndromic surveillance system monitors the numbers of daily unscheduled visits and calls to GPs during evenings, overnight, on weekends and on public holidays. Both systems cover around 55% of England’s population.

Up to 9 November 2020, GP out-of-hours and unscheduled care consultations for acute respiratory infections and influenza-like illness and difficulty breathing/asthma/wheeze remained stable (Figure 30).

Figure 30: GPOOH daily contacts (%) for (a) difficulty breathing/wheeze/asthma, (b) influenza-like illness and (c) acute respiratory infections, England
Sentinel swabbing scheme in England and the Devolved Administrations

In week 45 2020, 55 samples tested positive for SARS-CoV-2 with an overall positivity of 23.0% (55/239) compared to 16.3% (43/263) in the previous week, through the UK GP sentinel swabbing schemes (Figure 31).

Samples up to week 41 were only tested for SARS-CoV-2.

Figure 31: Number of influenza and COVID-19 positive samples and weekly positivity (%), UK GP sentinel swabbing scheme

*For the most recent week, more samples are expected to be tested therefore the graph in Figure 31 should be interpreted with caution

*Positivity (%) is not calculated when the total number tested is less than 10
Secondary care surveillance

SARI Watch

The Severe Acute Respiratory Infection (SARI) Watch surveillance system was established in 2020 to report the number of laboratory confirmed influenza and COVID-19 cases admitted to hospital and critical care units (ICU/HDU) in NHS acute trusts across England. This has replaced the USISS Mandatory and Sentinel data collections for influenza surveillance used in previous seasons, and the COVID-19 hospitalisations in England surveillance system (CHESS) collections for COVID-19 surveillance.

The weekly rate of new admissions of COVID-19 and influenza cases is based on the trust catchment population of those NHS Trusts who made a new return. This may differ from other published figures such as the total number of people currently in hospital with COVID-19.

Trends in hospital and critical care admission rates need to be interpreted in the context of testing recommendations.
Hospitalisations, SARI Watch

In week 45, the weekly hospital admission rate for COVID-19 increased further whilst the hospital admission rate remained low for influenza.

The hospitalisation rate for COVID-19 was at 14.03 per 100,000 in week 45 compared to 13.53 per 100,000 in the previous week. The hospitalisation rate for influenza was at 0.01 per 100,000 in week 45 compared to 0.00 per 100,000 in the previous week; and there was one new confirmed influenza (one influenza A(H3N2)) hospital admissions reported.

From the week 46 report, regional observations for SARI Watch will be by PHE Centres. By PHE centre, the highest hospital admission rate for COVID-19 was observed in the North East. By age groups, the highest hospital admission rate for confirmed COVID-19 was in the 85+ year olds.

**Figure 32: Weekly overall hospital admission rates per 100,000 of new COVID-19 and influenza positive cases reported through SARI Watch, England**

* influenza hospital admission rate is reported from week 40 2020 onwards
* influenza hospital admission rate based on 29 sentinel NHS trusts for week 45
* COVID-19 hospital admission rate based on 121 NHS trusts for week 45
Figure 33: Weekly overall influenza hospital admission rates per 100,000 trust catchment population with MEM thresholds, SARI Watch, England

* the MEM thresholds used are those from the 2019/20 season due to the pandemic

Figure 34: Weekly influenza hospital admissions by influenza type, SARI Watch, England
Figure 35: Weekly hospital admission rate by PHE Centre for new (a) COVID-19 positive cases and (b) influenza reported through SARI Watch

(a)

(b)
Figure 36: Weekly hospital admission rate by age group for new (a) COVID-19 positive cases and (b) influenza reported through SARI Watch
ICU/HDU admissions, SARI Watch

In week 45, the weekly ICU/HDU admission rates for COVID-19 were high but stable whilst the ICU/HDU admission rate remained low for influenza.

The ICU/HDU rate for COVID-19 was at 0.97 per 100,000 in week 45 (based on data reported from 116 NHS Trusts) compared to 0.92 per 100,000 in the previous week. The ICU/HDU rate for influenza was at 0.00 per 100,000 in week 45 compared to the same rate in the previous week. There was no new influenza confirmed ICU/HDU admissions.

From the week 46 report, regional observations for SARI Watch will be by PHE Centres. By PHE Centre, the highest ICU/HDU admission rates for COVID-19 were observed in East Midlands. By age groups, the highest ICU/HDU admission rates for COVID-19 were observed in the 65 to 74 year olds.

Figure 37: Weekly overall ICU/HDU admission rates per 100,000 of new COVID-19 and influenza positive cases reported through SARI Watch, England

* influenza ICU/HDU admission rate is reported from week 40 2020 onwards
* influenza ICU/HDU admission rate based on 108 NHS trusts for week 45
* COVID-19 ICU/HDU admission rate based on 116 NHS trusts for week 45
Figure 38: Weekly overall influenza ICU/HDU admission rates per 100,000 trust catchment population with MEM thresholds, SARI Watch, England

Figure 39: Weekly influenza ICU/HDU admissions by influenza type, SARI Watch, England
Figure 40: Weekly ICU/HDU admission rate by PHE Centre for new (a) COVID-19 positive cases and (b) influenza reported through SARI Watch

(a) [Graph showing ICU/HDU admission rate per 100,000 for different regions over weeks 27 to 53.]

(b) [Graph showing ICU/HDU admission rate per 100,000 for different regions over weeks 40 to 25.]
Figure 41: Weekly ICU/HDU admission rate by age group for new (a) COVID-19 positive cases and (b) influenza reported through SARI Watch

(a)

(b)
ECMO, SARI Watch

From week 27 2020, a total of 37 laboratory confirmed COVID-19 admissions have been reported from the 6 Severe Respiratory Failure (SRF) centres in the UK.

There were 12 new laboratory confirmed COVID-19 admissions reported in week 45 (Figure 42).

**Figure 42: Laboratory confirmed ECMO admissions (COVID-19, influenza and non-COVID-19 confirmed) to Severe Respiratory Failure centres in the UK**

*From the week 45 report (this report), data on ECMO admissions is being presented for the UK (including retrospective data from week 27 onwards).*
Emergency Department attendances, Syndromic surveillance

The Emergency Department Syndromic Surveillance System (EDSSS) monitors the daily visits in a network of emergency departments across England.

Up to 9 November 2020, the daily number of ED attendances for all ages as reported by 68 EDs, for COVID-19-like attendances increased while attendances for acute respiratory infections remained stable (Figure 43).

Please note: the COVID-19-like ED indicator is an underestimation of the number of COVID-19 attendances as it only includes attendances with a COVID-19-like diagnosis as their primary diagnosis. The EDSSS COVID-19-like indicator should therefore be used to monitor trends in ED attendances and not to estimate actual numbers of COVID-19 ED attendances. Further information about these caveats is available from the PHE Emergency Department Syndromic Surveillance bulletin.

Figure 43: Daily ED attendances for (a) COVID-19-like and (b) acute respiratory infections, all ages, England
Mortality surveillance

Cumulative COVID-19 deaths

Changes to the definitions of COVID-19 related deaths in England are described in more detail in an accompanying PHE technical summary.

The current definitions used for mortality surveillance of COVID-19 in England are:

(a) 28 day definition: A death in a person with a laboratory-confirmed positive COVID-19 test and died within (equal to or less than) 28 days of the first positive specimen date
(b) 60 day definition: A death in a person with a laboratory-confirmed positive COVID-19 test and either: died within 60 days of the first specimen date OR died more than 60 days after the first specimen date only if COVID-19 is mentioned on the death certificate

The introduction of these definitions will affect the numbers which have been presented in past reports and therefore Figure 44 represents these differences by definition.

Figure 44: Number of deaths since week 27 by week of death and time since laboratory confirmation of COVID-19, England

*The data are shown by the week of death. This gives the most accurate analysis of this time progression, however, for the most recent weeks’ numbers more deaths are expected to be registered therefore this should be interpreted with caution.
Figure 45: Age/sex pyramid of laboratory confirmed COVID-19 deaths, since week 27

Table 5: Ethnic group (%) of COVID-19 deaths and time since laboratory confirmation of COVID-19, England

<table>
<thead>
<tr>
<th>Ethnicity</th>
<th>28 day definition</th>
<th>60 day definition</th>
</tr>
</thead>
<tbody>
<tr>
<td>White</td>
<td>88.5</td>
<td>89.1</td>
</tr>
<tr>
<td>Asian / Asian British</td>
<td>8.2</td>
<td>7.6</td>
</tr>
<tr>
<td>Black / African / Caribbean / Black British</td>
<td>1.8</td>
<td>1.8</td>
</tr>
<tr>
<td>Mixed / Multiple ethnic groups</td>
<td>0.3</td>
<td>0.3</td>
</tr>
<tr>
<td>Other ethnic group</td>
<td>1.2</td>
<td>1.2</td>
</tr>
</tbody>
</table>
### Table 6: Cumulative number of COVID-19 deaths since week 27 and time since laboratory confirmation of COVID-19 by PHE Centres

<table>
<thead>
<tr>
<th>PHE Centres</th>
<th>28 day definition</th>
<th>60 day definition</th>
</tr>
</thead>
<tbody>
<tr>
<td>North East</td>
<td>694</td>
<td>797</td>
</tr>
<tr>
<td>North West</td>
<td>2,554</td>
<td>2,901</td>
</tr>
<tr>
<td>Yorkshire &amp; Humber</td>
<td>1,227</td>
<td>1,429</td>
</tr>
<tr>
<td>West Midlands</td>
<td>827</td>
<td>998</td>
</tr>
<tr>
<td>East Midlands</td>
<td>776</td>
<td>931</td>
</tr>
<tr>
<td>East of England</td>
<td>508</td>
<td>693</td>
</tr>
<tr>
<td>London</td>
<td>477</td>
<td>598</td>
</tr>
<tr>
<td>South East</td>
<td>573</td>
<td>829</td>
</tr>
<tr>
<td>South West</td>
<td>263</td>
<td>326</td>
</tr>
</tbody>
</table>
Figure 46: Cumulative mortality rate of COVID-19 cases per 100,000 population tested under Pillars 1 and 2 for the past four weeks by (a) 28 day definition and (b) 60 day definition

(a)
* Figure 46 has been calculated using mid-2019 ONS population estimates
Daily excess all-cause mortality (England)

Deaths occurring from 1 January to 04 November 2020 were assessed to calculate the daily excess above a baseline using age-group and region specific all cause deaths as provided daily by the General Register Office (GRO). The deaths were corrected to allow for delay to registration based on past data on these delays and the baseline was from the same day of the year in the previous 5 years +/- 7 days with an extrapolated time trend, and with 2 and 3 standard deviation (SD) limits shown (Figure 47).

Weeks in which at least 2 days exceeded the 3SD threshold are shown in Table 7 and the daily difference from the baseline by age and region is given in Figure 47. Note that as these data are by date of death with delay corrections, numbers are subject to change each week, particularly for more recent days.

Significant excess all-cause mortality was observed in week 44 overall, by age group in the 75 to 84 year olds and subnationally in the North West and Yorkshire and Humber. The excess noted in week 33 coincides with a heat wave (Figure 47, 48 and Table 7).

**Figure 47: Daily excess all-cause deaths in all ages, England, 1 January 2020 to 4 November 2020**

![Daily excess all-cause deaths graph](image)

^ based on same day in previous 5 years +/- 1 week with a linear trend projected
* corrected for delay to registration from death
Table 7: Excess all-cause deaths by (a) age group and (b) PHE centres, England

(a)

<table>
<thead>
<tr>
<th>Age Group</th>
<th>Excess detected in week 44 2020?</th>
<th>Weeks in excess since week 10 2020</th>
</tr>
</thead>
<tbody>
<tr>
<td>All</td>
<td>✓</td>
<td>13 to 21, 23, 33, 43, 44</td>
</tr>
<tr>
<td>under 25</td>
<td>x</td>
<td>None</td>
</tr>
<tr>
<td>25 to 44</td>
<td>x</td>
<td>14 to 16, 32, 38</td>
</tr>
<tr>
<td>45 to 64</td>
<td>x</td>
<td>12 to 19</td>
</tr>
<tr>
<td>65 to 74</td>
<td>x</td>
<td>13 to 19</td>
</tr>
<tr>
<td>75 to 84</td>
<td>✓</td>
<td>13 to 21, 33, 43, 44</td>
</tr>
<tr>
<td>85+</td>
<td>x</td>
<td>13 to 21, 33</td>
</tr>
</tbody>
</table>

(b)

<table>
<thead>
<tr>
<th>PHE Centres</th>
<th>Excess detected in week 44 2020?</th>
<th>Weeks in excess since week 10 2020</th>
</tr>
</thead>
<tbody>
<tr>
<td>East of England</td>
<td>x</td>
<td>14 to 19</td>
</tr>
<tr>
<td>East Midlands</td>
<td>x</td>
<td>13 to 19</td>
</tr>
<tr>
<td>London</td>
<td>x</td>
<td>12 to 19, 33</td>
</tr>
<tr>
<td>North East</td>
<td>x</td>
<td>14 to 21</td>
</tr>
<tr>
<td>North West</td>
<td>✓</td>
<td>13 to 20, 33, 42, 43, 44</td>
</tr>
<tr>
<td>South East</td>
<td>x</td>
<td>13 to 21, 33</td>
</tr>
<tr>
<td>South West</td>
<td>x</td>
<td>14 to 19, 33</td>
</tr>
<tr>
<td>West Midlands</td>
<td>x</td>
<td>13 to 20</td>
</tr>
<tr>
<td>Yorkshire and Humber</td>
<td>✓</td>
<td>14 to 21, 23, 43, 44</td>
</tr>
</tbody>
</table>
Figure 48: Daily excess all-cause deaths by (a) age group and (b) PHE centres, England, 1 March 2020 to 04 November 2020
Microbiological surveillance

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.

In week 45, no influenza viruses were characterised by PHE Respiratory Virus Unit (RVU).

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.

In week 45, no influenza viruses were tested for antiviral susceptibility.
Antimicrobial susceptibility

Table 8 shows in the 12 weeks up to week 45 2020, the proportion of all lower respiratory tract isolates of Streptococcus pneumoniae, Haemophilus influenza, 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 8: Antimicrobial susceptibility surveillance in lower respiratory tract

<table>
<thead>
<tr>
<th>Organism</th>
<th>Antibiotic</th>
<th>Specimens tested (N)</th>
<th>Specimens susceptible (%)</th>
</tr>
</thead>
<tbody>
<tr>
<td>S. pneumoniae</td>
<td>Penicillin</td>
<td>769</td>
<td>86</td>
</tr>
<tr>
<td></td>
<td>Macrolides</td>
<td>825</td>
<td>76</td>
</tr>
<tr>
<td></td>
<td>Tetracycline</td>
<td>818</td>
<td>77</td>
</tr>
<tr>
<td>H. influenzae</td>
<td>Amoxicillin/ampicillin</td>
<td>3,552</td>
<td>62</td>
</tr>
<tr>
<td></td>
<td>Co-amoxiclav</td>
<td>3,867</td>
<td>73</td>
</tr>
<tr>
<td></td>
<td>Macrolides</td>
<td>760</td>
<td>5</td>
</tr>
<tr>
<td></td>
<td>Tetracycline</td>
<td>3,965</td>
<td>97</td>
</tr>
<tr>
<td>S. aureus</td>
<td>Methicillin</td>
<td>2,789</td>
<td>94</td>
</tr>
<tr>
<td></td>
<td>Macrolides</td>
<td>3,027</td>
<td>70</td>
</tr>
<tr>
<td>MRSA</td>
<td>Clindamycin</td>
<td>106</td>
<td>38</td>
</tr>
<tr>
<td></td>
<td>Tetracycline</td>
<td>141</td>
<td>72</td>
</tr>
<tr>
<td>MSSA</td>
<td>Clindamycin</td>
<td>1,855</td>
<td>75</td>
</tr>
<tr>
<td></td>
<td>Tetracycline</td>
<td>2,520</td>
<td>93</td>
</tr>
</tbody>
</table>

* Macrolides = erythromycin, azithromycin and clarithromycin

Data source: PHE’s SGSS CDR module. Please note that this is different to the data source used during the 2019/20 influenza season when the SGSS AMR module was used, and so the results are not directly comparable.
There has been a reduction in the total number of bacterial positive lower respiratory tract clinical samples reported to PHE since mid-March 2020.
COVID-19 sero-prevalence surveillance

The results from testing samples provided by healthy adult blood donors aged 17 years and older, supplied by the NHS Blood and Transplant (NHS BT collection) between weeks 17-44 are summarised. This programme has previously involved testing approximately 1000 donor samples from two different NHS regions each week. In this week’s report, the data presented reflects a change in the sampling strategy as of week 44, with approximately 250 samples from each geographic NHS region being tested each week. Since week 26, an exclusion of donors aged 70 years and older donating throughout lockdown was lifted, and therefore data since then include donors in this older age group.

Seroprevalence in Adults aged 17 years and older (Blood Donors)

The results presented here are based on testing using the Euroimmun assay for blood donor samples collected between weeks 17-44. This report presents seropositivity estimates using a 4-week rolling prevalence for national and regional estimates. Seroprevalence estimates reported are based on seropositivity which are unadjusted for the sensitivity and specificity of the assays used. This is because assay sensitivity will change according to the time since infection in these cohorts due to waning of antibodies.

National Prevalence

Overall population weighted (by age group, sex and NHS region) antibody prevalence using the Euroimmun assay among blood donors aged 17 years and older in England was 5.9% (95% CI 4.9% - 7.1%) for the period 7th October– 1st November (weeks 41-44). Estimates are based on 7708 samples, of which 466 were positive. This compares with 5.7% (95% CI 5.1% - 6.2%) for the period of 16th September – 11th October (weeks 38-41). Changes in prevalence over time need to take into account demographic changes in the donor population, with later data including donors aged 70 years and older who were previously excluded from donating during lockdown. Waning immunity is also likely to be a contributing factor.

Regional Prevalence over Time

Seropositivity (weighted by age group and sex) vary across the country and over time. Figure 1 shows the overall 4-weekly rolling proportion seropositive in each region over time. Seropositivity estimates are plotted on the mid-point of the 4-weekly period.

In London where estimates have consistently been highest, the 4-weekly rolling seropositivity increased from 11.9% (week 16-19) to 13.7% (weeks 20-23). From week 24 seropositivity declined and plateaued with estimates at 7.8% in weeks 30-33. This was then followed by a rise in seropositivity to 10.4% (95% CI 9.1% - 12%) in weeks 34-37 and has plateaued to 8.2% (95% CI 7.2% - 9.4%) in weeks 41-44. Contributory factors to this fluctuation are likely to include variability in the precise locations of
sampling within London and changes in exposure of donors. Increases in seropositivity observed in weeks 34-37 in part may reflect samples being tested from donors who were likely to be returning to donate having donated in earlier parts of the epidemic when incidence was high.

Data from the North West show that seropositivity increased from 5.3% (95% CI 4.1 - 6.9%) in weeks 39-42 to 6.6% (95% CI 5.1% - 8.1%) in weeks 41-44.

In the East of England seropositivity was 4.4% (95% CI 3.4% - 5.7%) in the most recent data (weeks 41-44) fluctuating between 4.1% (95% CI 3.0% - 5.7%) in weeks 32-35 and 5.9% (95% CI 4.6% - 7.6%) in weeks 35-38.

Seropositivity in the South East region was 3.8% (95% CI 2.9% - 5.1%) for weeks 41-44 lower than the 5.1% (95% CI 3.3% - 7.7%) observed in weeks 33-36.

Seropositivity in the South West region was 3.9% (95% CI 2.7 – 5.6%) in weeks 41-44 lower than the 4.1% (95% CI 2.9% - 5.8%) observed in the previous survey in weeks 37-40.

In the North East and Yorkshire NHS region the seropositivity has increased to 5.8% (95% CI 3.5%-9.5%) in weeks 41-44 compared with 3.7% (95% CI 2.7% - 5.0%) in week 36-39.

Data from the Midlands also show a higher proportion seropositive at 7.3% (95% CI 3.4% - 15.0%) in weeks 41-44. This compares to 5.5% (95% CI 4.3% - 7.0%) in weeks 36-39. Due to the recent change in sampling strategy, there are relatively few samples from the Midlands included in the latest rolling 4-week window, resulting in a larger uncertainty for this most recent estimate. This is expected to narrow in subsequent weeks.

The change in proportion seropositive observed in some regions is likely to be driven by changes in the precise locations of sample collection. However, the most recent increases observed in the Midlands, North West and North East regions cannot be fully explained by this and are likely to reflect increased transmission, consistent with other surveillance data. Increases in seropositivity reflect transmission occurring at least two to three weeks previously given the time taken to generate an antibody response following infection.

Declines in prevalence observed during the summer months can be partially explained by demographic differences in the donor population as lockdown measures were relaxed. Examples include a reduction in attendance of regular donors in August and that donors aged 70 years and above were not allowed to donate during lockdown, but
this exclusion was lifted from week 26. Waning immunity will also be a contributing factor to the lower prevalence.

Prevalence by age group is not reported in this week’s report due to the recent change in sampling strategy. The overlap in sampling methodologies in the current reporting period has resulted in a small number of samples from older age groups in some regions which makes interpretation of trends difficult.

**Figure 49: 4-weekly rolling SARS-CoV-2 antibody seroprevalence (% seropositive) in blood donors by region, using Euroimmun test; error bars show 95% confidence intervals**
Influenza vaccination

Influenza vaccine uptake in GP patients

Up to week 45 2020 in 87.2% of GP practices reporting weekly to Immform for the main collection, the provisional proportion of people in England who had received the 2020/21 influenza vaccine in targeted groups was as follows (Figure 50):

- 35.2% in under 65 years in a clinical risk group
- 27.9% in pregnant women
- 70.7% in 65+ year olds

Figure 50: Cumulative weekly influenza vaccine uptake by target group in England

In 2020/21, all 2 and 3 year olds continue to be eligible for influenza vaccination through their GPs. Up to week 43 2020, in 97.3% of GP practices reporting weekly to Immform for the childhood collection, the provisional proportion of children in England who had received the 2020/21 influenza vaccine in targeted groups was as follows (Figure 51):

- 41.9% in 2 year olds
- 43.4% in 3 year olds
Figure 51: Cumulative weekly influenza vaccine uptake in 2 and 3 year olds, in England

2020/21 season indicated by bold lines, 2019/20 season indicated by fainter lines
Influenza vaccine uptake in school age children

The first report on influenza vaccine uptake in school age children (Year Reception to Year 7) will be published in November 2020.

Influenza vaccine uptake in healthcare workers

The first report on influenza vaccine uptake in healthcare workers will be published in November 2020.
**International update**

**Global COVID-19 update**

Globally, up to 10 November 2020, a total of 51,103,820 cases of COVID-19 infection have been reported worldwide, including 1,265,862 COVID-19 related deaths.

For further information on the global COVID-19 situation please see the WHO COVID-19 situation reports.

**Figure 52: Global map of cumulative COVID-19 cases**
Figure 53: Global map of weekly COVID-19 case incidence rate per 100,000, week 45 2020
Global influenza update

Updated on 11 November 2020 (based on data up to 25 October 2020) (WHO website)

In the temperate zone of the northern hemisphere, influenza activity remained below inter-seasonal levels, though sporadic detections were reported in some countries. In the temperate zones of the southern hemisphere, no influenza detections were reported across countries. Worldwide, of the very low numbers of detections reported, seasonal influenza A(H3N2) viruses accounted for the majority of detections.

In the countries of North America, influenza activity indicators, including the percent of tests positive for influenza, were at very low levels.

In Europe, influenza activity remained at inter-seasonal levels though sporadic detections were reported across reporting countries.

In Central Asia and Northern Africa, there were no influenza updates for this reporting period.

In Western Asia, sporadic influenza detections were reported in recent weeks.

In East Asia, influenza illness indicators and influenza activity remained at inter-seasonal levels in most reporting countries.

In the Caribbean and Central American countries, sporadic influenza detections were reported.

In tropical South America, there were no influenza detections across reporting countries.

In tropical Africa, influenza activity was reported in West Africa in Côte d’Ivoire and Niger, and in East Africa in Kenya.

In Southern Asia, influenza activity of predominately influenza A(H3N2) was reported in Bangladesh and India in recent weeks.

In South East Asia, influenza detections continued to be reported in Cambodia and Lao People’s Democratic Republic (PDR).

In Oceania, ILI and other influenza activity indicators remained below usual levels for this time of year in general, despite continued testing.

The WHO GISRS laboratories tested more than 94241 specimens between 12 October 2020 and 25 October 2020. 140 were positive for influenza viruses, of which 80 (57.1%)
were typed as influenza A and 60 (42.9%) as influenza B. Of the sub-typed influenza A viruses, 2 (5.4%) were influenza A(H1N1pdm09) and 35 (94.6%) were influenza A (H3N2). Of the characterized B viruses, 1 (6.7%) belonged to the B-Yamagata lineage and 14 (93.3%) to the B-Victoria lineage.

**Influenza in Europe**

Updated on 9 November 2020 ([Joint ECDC-WHO Europe Influenza weekly update](https://www.ecdc.europa.eu/en/healthtopics/influenza-weekly-updates)

For week 44 2020, influenza activity remained at interseasonal levels throughout Europe.

Of 31 countries and areas that reported on the intensity indicator, 28 reported activity at baseline levels, and 3 reported low intensity (Azerbaijan, Serbia and Slovakia) for week 44 2020. Of 32 countries and areas that reported on geographic spread, 27 reported no activity and 5 reported sporadic spread (Azerbaijan, Denmark, Portugal, Slovakia and United Kingdom (Scotland)) for week 44 2020.

For week 44 2020, of 260 sentinel specimens tested for influenza viruses, none were positive.

There were no hospitalized laboratory-confirmed influenza cases in ICUs for week 44 2020 and since the start of the season.

There were no laboratory-confirmed influenza cases in wards outside ICUs for week 44 2020 and since the start of the season.

**Influenza in the Northern Hemisphere**

Influenza activity remains low in the United States of America and in Canada.

For further information on influenza in the United States of America please see the [Centre for Disease Control weekly influenza surveillance report](https://www.cdc.gov/flu/weekly/).

For further information on influenza in Canada please see the [Public Health Agency weekly influenza report](https://www.canada.ca/en/public-health/services/publications/influenza-report.html).
Other respiratory viruses

Avian influenza

Latest update on 9 November 2020 (WHO website)

Influenza A(H5) viruses:
According to reports received by the World Organisation for Animal Health (OIE), various influenza A(H5) subtypes continue to be detected in birds in Africa, Europe and Asia.

Influenza A(H7N9) viruses:
There have been no publicly available reports from animal health authorities in China or other countries on influenza A(H7N9) virus detections in animals in recent months.

Influenza A(H9N2) viruses:
Since the last update on 10 July 2020, one new laboratory-confirmed human case of influenza A(H9N2) virus infection was reported from China to WHO on 28 August 2020.

Middle East respiratory syndrome coronavirus (MERS-CoV)

Latest update on 10 November 2020 (WHO website)

Up to 10 November 2020, a total of five cases of Middle East respiratory syndrome coronavirus, MERS-CoV, (three imported and two linked cases) have been confirmed in the UK. On-going surveillance has identified 1,816 suspected cases in the UK since September 2012 that have been investigated for MERS-CoV and tested negative.

From 1 April to 31 May 2020, the National IHR Focal Point of Saudi Arabia reported 9 new cases of MERS-CoV infection, including five deaths.

Globally, since September 2012, WHO has been notified of 2,562 laboratory-confirmed cases of infection with MERS-CoV, including 881 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 very low.
Related links

Previous national COVID-19 reports
Previous weekly influenza reports
Annual influenza reports
Sources of influenza surveillance data
Sources of COVID-19 surveillance data

PHE has delegated authority, on behalf of the Secretary of State, to process Patient Confidential Data under Regulation 3 The Health Service (Control of Patient Information) Regulations 2002 http://www.legislation.gov.uk/uksi/2002/1438/regulation/3/made. Regulation 3 makes provision for the processing of patient information for the recognition, control and prevention of communicable disease and other risks to public health.
About Public Health England

Public Health England exists to protect and improve the nation’s health and wellbeing, and reduce health inequalities. We do this through world-leading science, research, knowledge and intelligence, advocacy, partnerships and the delivery of specialist public health services. We are an executive agency of the Department of Health and Social Care, and a distinct delivery organisation with operational autonomy. We provide government, local government, the NHS, Parliament, industry and the public with evidence-based professional, scientific and delivery expertise and support.

Public Health England
Wellington House
133-155 Waterloo Road
London SE1 8UG
Tel: 020 7654 8000

---

You may re-use this information (excluding logos) free of charge in any format or medium, under the terms of the Open Government Licence v3.0. To view this licence, visit OGL. Where we have identified any third party copyright information you will need to obtain permission from the copyright holders concerned.

PHE supports the UN Sustainable Development Goals