



UK Health
Security
Agency

COVID-19 public health advice changes: scientific summary

Contents

Background.....	3
1. COVID-19 symptoms.....	4
2. Duration of infectiousness	5
SARS-CoV-2	5
Influenza.....	6
3. Conclusion.....	7
References.....	8

Background

The provision of free universal symptomatic and asymptomatic testing for the general public ended in England on 1 April 2022.

New [guidance](#) for the public was introduced on 1 April, advising people with symptoms of respiratory infections including COVID-19 to try to stay at home and avoid contact with other people if they have a high temperature or do not feel well enough to go to work or carry out normal activities. They are advised to resume normal activities once their temperature is resolved and they feel well enough to do so.

Over the course of the pandemic, a wide range of symptoms have been associated with COVID-19. These symptoms are similar to those seen in other respiratory viral infections, so in the absence of testing it is not possible to distinguish one viral illness from another based on symptoms alone.

This paper briefly summarises evidence that supports the guidance introduced on 1 April 2022.

1. COVID-19 symptoms

In the UK, the most commonly reported symptoms have consistently been cough, fatigue and headache, and the least commonly reported symptoms have consistently been abdominal pain, diarrhoea, and nausea or vomiting (1).

In the REACT study, cough, fever and anosmia have remained the symptoms with highest positive and negative predictive value for COVID-19 (2), although anosmia or ageusia was less prominent in infection with the Omicron variant. Cough and anosmia can persist for several weeks after the infectious period.

Other symptoms, such as sore throat, headache, runny nose and fatigue tend to manifest earlier (3) when individuals are likely to be more infectious, but are also very common in people without COVID-19 and are therefore poorly predictive of infection.

If there are other viruses circulating the predictive value of these symptoms decreases even further.

The symptom profile of COVID-19 has been shown to differ by age. In older patients, COVID-19 may be more likely to present with 'atypical' symptoms such as delirium, falls, generalized weakness, malaise, functional decline, and other less specific symptoms.

For most children and young people COVID-19 is a mild or asymptomatic infection and very few children become seriously ill. They generally present with similar symptoms to adults, though there are reports of more frequent presentation of fatigue and headaches (4).

Findings from a systematic review by Viner and others (5) showed that that fever and cough were the most common symptoms, occurring in around 40 to 60% of infected children and young people.

The symptom profile for the Omicron variant is similar to that of previous variants. A UKHSA analysis (6) comparing the symptom profile of Delta to Omicron variant showed that fever and cough were slightly more likely to be reported in Omicron compared to Delta infections, whereas loss of smell and taste were less likely to be reported.

There are no specific symptoms that predict a worse outcome of COVID-19 by themselves. Various risk scores have been developed in the last 2 years, however most of these require the addition of one or more laboratory tests. A recent pre-print of a simple clinical risk calculator showed that along with various comorbidities, dyspnoea, high respiratory rate and low (less than 95%) oxygen saturation on air were all associated with a significantly higher risk of hospitalization. The authors have now externally validated this risk score (7).

2. Duration of infectiousness

SARS-CoV-2

Epidemiological and contact tracing studies have shown that the infectiousness period of SARS-CoV-2 ranges from 9 days before symptom onset to 15 days after symptom onset, with most transmission occurring 3 days before symptom onset to 5 days after symptom onset (8). Limited viral RNA data on the Omicron variant show that the viral load peaks at 3 to 6 days post symptom onset, which is slightly shorter than that reported for wild-type lineages (9).

The one human challenge study to date in healthy young volunteers (10) with the original Wuhan virus strain showed that viral load peaked about 5 days post inoculation and viable virus was on average detectable until day 10 after the first positive test.

Positive lateral flow device (LFD) tests have been shown to be associated with high viral load in infectious cases (11). Recent data from the ATTACC study (12) and the human challenge study (10) confirm these findings and have shown a direct correlation between viral growth (as a proxy for infectiousness) and LFD tests in the course of an individual infection. A recent CDC (14) study assessing antigen test positivity in 729 individuals showed that 54% of individuals had a positive result between 5 to 9 days after symptom onset.

Recent real world data (unpublished) from NHS Test and Trace (13) on LFD positivity in individuals testing to reduce their 10-day isolation period show that 70% of individuals report a negative LFD result on both day 5 and 6, although the data are self-reported and there is likely to be reporting bias. There was an age gradient with a higher proportion (over 80%) of individuals aged under 20 reporting negative LFD results on days 5 and 6, compared to less than 60% in individuals aged over 60 years.

Individuals who are infected with SARS-CoV-2 and who are asymptomatic can still transmit virus to others (15). The evidence on whether the proportion of asymptomatic cases is higher in children than in adults remains limited (3,16). While there is evidence that asymptomatic children can transmit the disease (2,15), findings from a systematic review published in March 2022 suggest that asymptomatic cases (adults and children) are less infectious than symptomatic cases (16).

There is some evidence that children may be less infectious than adults (2,17). One rapid review suggested a shorter mean duration of viral RNA shedding in respiratory symptoms in children than adults (8). However, detection of viral RNA does not provide direct evidence of infectiousness. Early studies showed the median duration of fever in children to be 3 days compared to 10 days in adult patients (18), suggestive of a shorter illness in children. These results are also in line with the findings from a cohort study conducted in 258,790 UK school-

aged children that suggested COVID-19 in children tended to be of short duration and low symptom burden (4).

Influenza

Viral excretion data from influenza infected individuals indicate that the infectious period is shorter for influenza than for COVID-19. The mean infectious period has been estimated in the region of 1 day (95% CI 0.5 to 1.7), with 95% of cases infectious for less than 2.9 days (19).

Donnelly and others (20) estimated that only 5% of transmission events took place more than 3 days after the onset of clinical symptoms with the predominant influenza (H1N1) virus in 2009.

The proportion of true asymptomatic influenza infection is difficult to estimate due to varying study methods and populations.

In outbreak investigations (where infections were virologically confirmed) 16% of infections were found to be asymptomatic, whereas in longitudinal studies using serology, the point estimates of asymptomatic infections were estimated at around 65 to 85% (21), although there may be recall biases intrinsic to these studies.

A systematic review (22) assessing the relationship between viral shedding and disease transmission found limited evidence that asymptomatic or pre-symptomatic individuals plays an important role in influenza transmission. This is consistent with findings from Laue and others (23), which show that asymptomatic or pre symptomatic transmission may not be an important factor in influenza epidemics.

3. Conclusion

The similarity between the symptom profile of COVID-19 and other respiratory viral infections means that people presenting with acute respiratory viral symptoms are unable to distinguish one viral illness from another based on symptoms alone.

There is no strong evidence that any specific COVID-19 symptoms are associated with greater infectiousness.

References

1. Office for National Statistics (ONS). [Coronavirus \(COVID-19\) latest insights](#)
2. P Elliott and others. '[Twin peaks: the Omicron SARS-CoV-2 BA.1 and BA.2 epidemics in England](#)'. REACT-1, round 19 (preprint)
3. Khan SM, Farland LV, E, and others. '[Symptoms of COVID-19 in a population-based cohort study](#)'. medRxiv 2021 (preprint)
4. Molteni and others. '[Illness duration and symptom profile in symptomatic UK school-aged children tested for SARS-CoV-2](#)'. The Lancet Child and Adolescent Health, August 2021
5. Viner RM and others. 'Systematic review of reviews of symptoms and signs of COVID-19 in children and adolescents'. Archives of Disease in Childhood, 17 December 2020. archdischild-2020-320972. doi: 10.1136/archdischild-2020-320972. Epub ahead of print. PMID: 33334728; PMCID: PMC7747494
6. UK Health Security Agency. '[SARS-CoV-2 variants of concern and variants under investigation in England: Technical briefing 34. 2022](#)'
7. Ebell MH and others. 'Development and Validation of the COVID-NoLab and COVID-SimpleLab Risk Scores for Prognosis in 6 US Health Systems'. Journal of the American Board of Family Medicine, February 2021, volume 34, supplement pages S127 to S135. doi: 10.3122/jabfm.2021.S1.200464. PMID: 33622827; PMCID: PMC8343954
8. Ontario Agency for Health Protection and Promotion (Public Health Ontario). '[COVID-19 overview of the period of communicability: what we know so far](#)'. 2021
9. Ontario Agency for Health Protection and Promotion (Public Health Ontario). '[COVID-19 Omicron \(B.1.1.529\) Variant of Concern and Communicability...What We Know So Far](#)'. 2022
10. Killingley B, Mann AJ, Kalinova M and others. '[Safety, tolerability and viral kinetics during SARS-CoV-2 human challenge in young adults](#)'. Nature Medicine 2022
11. Lee LYW and others. 'Severe Acute Respiratory Syndrome Coronavirus 2 (SARS-CoV-2) Infectivity by Viral Load, S Gene Variants and Demographic Factors, and the Utility of Lateral Flow Devices to Prevent Transmission'. Clinical Infectious Diseases, 11 February

2022, volume 74, issue 3, pages 407 to 415. doi: 10.1093/cid/ciab421. PMID: 33972994; PMCID: PMC8136027

12. Unpublished NERVTAG paper: Kelly and others. 'Assessment of the relationship between lateral flow antigen tests and culture across the duration of infection for SARS-CoV-2 incident cases'
13. Unpublished UKHSA paper. Futschik and others. 'LFD positivity rates at Day 5 and 6'
14. Lefferts B, Blake I, Bruden D and others. '[Antigen Test Positivity After COVID-19 Isolation: Yukon-Kuskokwim Delta Region, Alaska, January to February 2022](#)'. Morbidity and Mortality Weekly Report 2022, volume 71, pages 293 to 298
15. Ravindra K and others. 'Asymptomatic infection and transmission of COVID-19 among clusters: systematic review and meta-analysis'. Public Health February 2022, volume 203, pages 100 to 109. doi: 10.1016/j.puhe.2021.12.003. Epub 2021 Dec 9. PMID: 35038628; PMCID: PMC8654597
16. Buitrago-Garcia and others. '[Occurrence and transmission potential of asymptomatic and presymptomatic SARS-CoV-2 infections: update of a living systematic review and meta-analysis](#)'. medRxiv (preprint)
17. World Health Organization (WHO). '[COVID-19 disease in children and adolescents: scientific brief](#)', 29 September 2021. 2021
18. Dhochak N, Singhal T, Kabra SK, Lodha R. 'Pathophysiology of COVID-19: Why Children Fare Better than Adults?' Indian Journal of Pediatrics. July 2020, volume 87, issue 7, pages 537 to 546. doi: 10.1007/s12098-020-03322-y. Epub 2020 May 14. PMID: 32410003; PMCID: PMC7221011
19. Cori and others. 'Estimating influenza latency and infectious period durations using viral excretion data'. August 2012: volume 4, issue 3, pages 132 to 138. doi: 10.1016/j.epidem.2012.06.001. Epub 2012 Jun 13
20. Donnelly C and others. 'Serial intervals and the temporal distribution of secondary infections within households of 2009 pandemic influenza A (H1N1): implications for influenza control recommendations'. Clinical Infectious Diseases 2011, volume 52, pages S123 to S130

21. Leung and others. 'Review Article: The Fraction of Influenza Virus Infections That Are Asymptomatic: A Systematic Review and Meta-analysis'. *Epidemiology* 2015, volume 26, issue 6, pages 862 to 872. doi:10.1097/EDE.0000000000000340
22. Patrozou E, Mermel LA. 'Does influenza transmission occur from asymptomatic infection or prior to symptom onset?'. *Public Health Reports* 2009, volume 124, issue 2, pages 193 to 196. doi:10.1177/003335490912400205
23. Lau LL, Cowling BJ, Fang VJ and others. 'Viral shedding and clinical illness in naturally acquired influenza virus infections'. *Journal of Infectious Diseases* 2010, volume 201, issue 10, pages 1509 to 1516. doi:10.1086/652241

About the UK Health Security Agency

UKHSA is responsible for protecting every member of every community from the impact of infectious diseases, chemical, biological, radiological and nuclear incidents and other health threats. We provide intellectual, scientific and operational leadership at national and local level, as well as on the global stage, to make the nation health secure.

[UKHSA](#) is an executive agency, sponsored by the [Department of Health and Social Care](#).

© Crown copyright 2022
Version 1.0

Published: April 2022
Publishing reference: GOV-12079



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.



UKHSA supports the
Sustainable Development Goals

