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COVID-19 - SARS-CoV-2

NOTIFIABLE

The disease

COVID-19 is a disease of the respiratory tract caused by the SARS-CoV-2 virus, an RNA virus of the family of Coronaviridae and genus Betacoronavirus (Zhu *et al*, 2020). As with other coronaviruses, SARS-CoV-2 encodes four major structural proteins, spike (S), membrane (M), envelope (E) and a helical nucleocapsid (N) (Dhama *et al*, 2020) The S glycoprotein is considered the main antigenic target, including the receptor binding domain (RBD) which binds to angiotensin converting enzyme 2 (ACE2) on host cells (Kaur *et al*, 2020, Amanat *et al*, 2020).

SARS-CoV-2 is primarily spread through respiratory droplets and aerosols and from direct person-to-person contact. The role of fomites appears to play a minor role in transmission. (Goldman *et al*. 2020). Viral RNA can persist in respiratory samples for 7-12 days after symptom onset, and viral loads are highest soon after symptom onset.

Secondary attack rates within households are high (Lopez Bernal *et al*, 2020). The estimated reproductive number (R0) for the original wild-type strain was 2.8, with subsequent strains becoming more infectious. The Delta strain had an estimated R0 of 5.1 and the Omicron strain, which emerged in late 2021, had an estimated R0 of 9.5. (Liu *et al*. 2022)

Common symptoms include headache, fatigue, cough, and muscle aches. In severe cases, COVID-19 can lead to pneumonia, acute respiratory distress syndrome, multiple organ failure and death. Compared to previous variants, Omicron is less likely to cause loss of sense of smell (anosmia) and more likely to cause a sore throat. (Menni *et al*. 2022, Pachetti *et al*, 2020).

The long-term sequelae of COVID-19 infection, dubbed Long COVID or Post-Acute Sequelae of SARS-CoV-2 (PASC) infection, are an area of ongoing study. In the UK, 4.5% of cases report long-term symptoms 12-16 weeks after the initial infection. Reported symptoms are varied, involving most organ systems and affecting both physical and mental health. (Crook *et al*. 2022)

Natural immunity due to previous infection lasts up to 1 year before beginning to wane, (Hall *et al*, 2022), although new strains and variants, such as Omicron, appear to exhibit greater immune escape, making reinfection more common.

History and epidemiology of the disease

Initial reports of severe respiratory infections of an unknown origin first arose in Wuhan, China, in late 2019 (WHO, 2020), with sequenced lower respiratory tract samples subsequently detecting a novel coronavirus (Huang *et al.*, 2020). In March 2020, the World Health Organization (WHO) declared a SARS-CoV-2 pandemic (WHO Director- General, 2020). As of October 2022, there have been over twenty million confirmed episodes of COVID-19 in England, and over 600 million worldwide.

The first COVID cases were initially detected in the UK in January 2020, with further cases being detected in February. Cases continued to rise throughout March until a national lockdown was introduced on 23 March 2020.

During the first wave in the UK, infection fatality ratios (IFR) for COVID-19, derived from combining mortality data with infection rates in seroprevalence studies, showed markedly higher IFR in the oldest age groups (Ward *et al.*, 2020). In the first year of the pandemic, cumulative death rates after COVID infection have therefore been highest in those aged over 75; death rates in males exceeded those in females (table 1).

During autumn 2020, the Alpha variant, noted for its increased transmissibility over the wild type, was first detected in Kent. By December 2020, Alpha had become the dominant strain in the UK. In April 2021, the Delta variant, first observed in India, was detected in the UK, and became dominant by July 2021. On 3 December 2021, the Omicron variant, first observed in South Africa, reached the UK, becoming the dominant variant by the 17th December 2021. Overall, Omicron has been shown to cause less severe disease than the previous strains, albeit on a background of a population with immunity due to vaccination and previous infection. Compared with Delta, Omicron was 40% as likely to cause hospital admission and 30% as likely to cause death. Nyberg *et al.*, 2022).

Successive sub-lineages (BA.1, BA.2, BA.4, BA.5) of the Omicron variant circulated during 2022, often associated with an increase in incidence rates (figure).

Information on new variants under investigation is available at: <https://www.gov.uk/government/publications/investigation-of-sars-cov-2-variants-technical-briefings>

Table 1: Cumulative deaths within 28 days of a positive test for SARS-CoV-2 in England, 2020

	Male		Female		Total	
	Count (n)	Rate (per 100,000)	Count (n)	Rate (per 100,000)	Count (n)	Rate (per 100,000)
Under 18	15	0.24	17	0.29	32	0.26
18 to 49	809	6.94	525	4.56	1334	5.76
50 to 64	3751	70.34	2008	36.50	5759	53.16
65 to 74	6822	253.45	3772	129.77	10594	189.23
75 to 84	13307	845.38	8773	465.39	22080	638.30
85 and over	13134	2498.24	13820	1569.24	26954	1916.51
Total	37838	135.22	28915	101.22	66753	118.04

COVID-19 - SARS-CoV-2

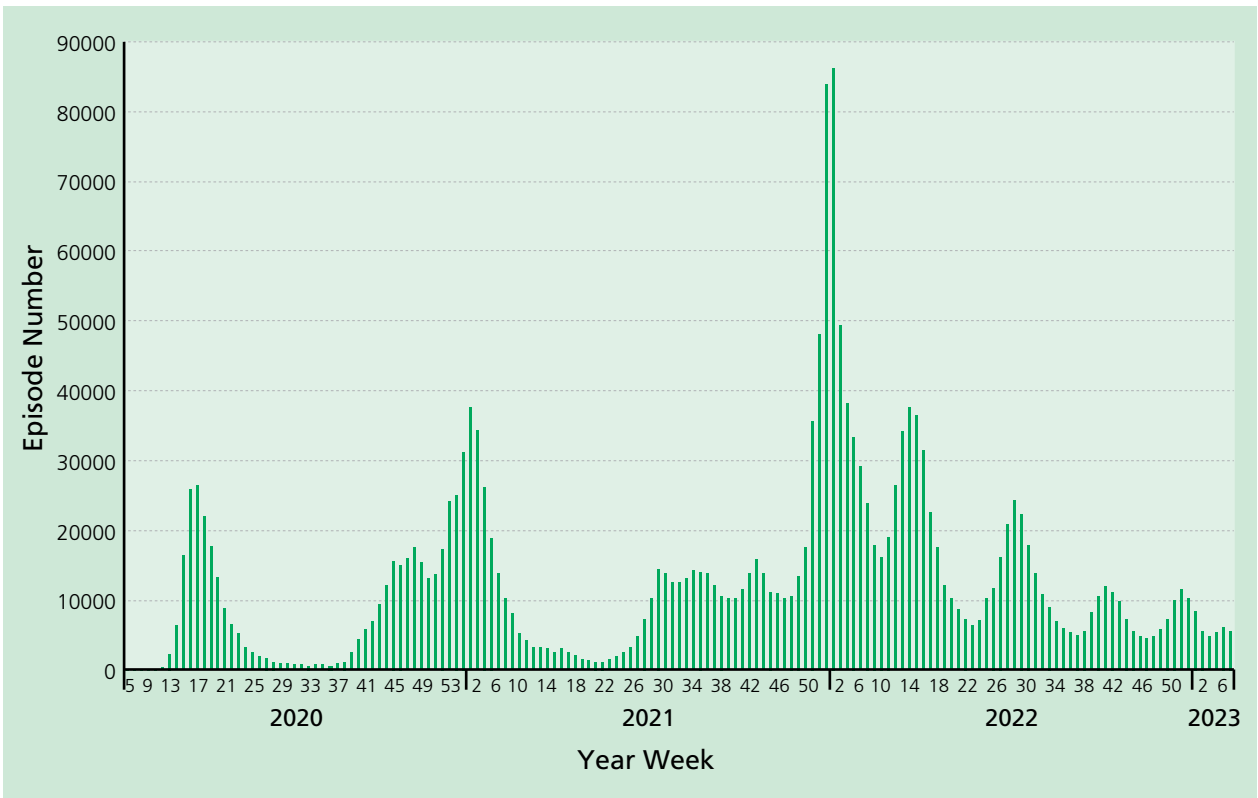


Figure: Number of confirmed episodes of SARS-CoV-2 infection: by test type and week 2020-2023. (Source: pillar 1 testing in English NHS laboratories)

Children

In general, children with SARS-CoV-2 infection remain asymptomatic or develop mild disease, often with upper respiratory symptoms, but symptoms may be non-specific and atypical, affecting other organ systems. Following the emergence and rapid spread of the Omicron variant and subvariants since November 2021, nearly all children in the UK now have antibodies against SARS-CoV-2; mostly due to natural infection in the youngest age groups.

Severe COVID-19 requiring hospitalisation is rare in children and deaths even more so, with an infection fatality ratio of less than 1 in 100,000 infections in 0-19 year-olds (Bertran *et al.*, 2022).

Paediatric multisystem inflammatory syndrome temporally associated with SARS-CoV-2 infection (PIMS-TS), known in the USA as multisystem inflammatory syndrome in children, (MIS-C), a presentation similar to Kawasaki disease, was first identified in April 2020 and estimated to affect around 1 in 3,000 children. This syndrome was reported most commonly in male children aged 6 to 12 years, and the appearance of cases of lagged behind cases of COVID-19 by around 4 weeks. (Feldstein *et al.* 2021).

PIMS-TS risk in the UK declined during the Delta wave, falling further after the Omicron wave, likely because of natural and vaccine-induced immunity against the virus in children and key mutations in the more recent SARS-CoV-2 variants (Shingleton *et al.*, 2022; Cohen *et al.*, 2022).

The majority of children recover completely after acute SARS-CoV-2 infection and any persistent systems will improve with time (Rytter *et al.*, 2021). Serious long-term complications after recovering from acute SARS-CoV-2 infection, too, are rare in children and studies are ongoing to assess risks and outcomes in longitudinal follow-up studies.

Pregnant women and neonates

Pregnant and recently pregnant women with COVID-19 are more likely to be admitted to an intensive care unit, have invasive ventilation or extracorporeal membrane oxygenation in comparison to non-pregnant women of reproductive age (Allotey *et al.*, 2021).

UK studies have suggested a higher than background rate of stillbirth in infected women (Allotey *et al.*, 2020, Gurol-Urganci *et al.*, 2021). The risk of preterm birth is also increased in women with symptomatic COVID-19 (Vousden *et al.*, 2021), usually as a result of a medical recommendation to deliver early to improve maternal oxygenation.¹

Pregnant women are more likely to have severe COVID-19 infection if they are overweight or obese, are of black and asian minority ethnic background, have co-morbidities such as diabetes, hypertension and asthma, or are 35 years old or older (Vousden *et al.*, 2021, Allotey *et al.*, 2020).

The risks to pregnant women and neonates following COVID-19 infection in the UK appear to have changed over the course of the pandemic. The proportion of pregnant women admitted to intensive care units, the maternal mortality ratio, the stillbirth rate and the number of neonatal deaths, increased between the wildtype SARS-CoV-2 dominant period to the Delta dominant period. (Vousden *et al.*, 2021a, Knight *et al.*, 2022).

In contrast, pregnant women infected with SARS-CoV-2 were substantially less likely to have a preterm birth or maternal critical care admission during the Omicron period than

1 NICE Guideline 25, 2019 <https://www.nice.org.uk/guidance/ng25>

during the Delta period; fewer stillbirths and no neonatal deaths were observed in the Omicron period (Stock *et al*, 2022). Despite this, even in the Omicron era, severity of COVID-19 was higher in unvaccinated than vaccinated women (Engjom *et al*, 2022).

COVID-19 vaccines

The recognition of the pandemic accelerated the development and testing of several vaccines using platforms investigated during previous emergencies such as the SARS pandemic (Amanat *et al*, 2020) and Ebola in West Africa. Candidate vaccines included nucleic acid vaccines, inactivated virus vaccines, live attenuated vaccines, protein or peptide subunit vaccines, and viral-vectored vaccines.

Most vaccine candidates focussed on immunisation against the spike (S) protein, which is the main target for neutralising antibodies. Neutralising antibodies that block viral entry into host cells through preventing the interaction between the spike protein Receptor Binding Motif (RBM) and the host cell Angiotensin-converting enzyme 2 (ACE2) were expected to be protective (Addetia *et al*, 2020, Thompson *et al*, 2020).

All vaccines now authorised for primary vaccination in the UK target the S protein of the original SARS-CoV-2 strain; two use an mRNA platform (Pfizer BioNTech COVID-19 BNT162b2 vaccine (Comirnaty® and Moderna mRNA-1273 COVID-19 vaccine/Spikevax®), two use an adenovirus vector (AstraZeneca COVID-19 ChAdOx1-S vaccine/ Vaxzevria® and COVID-19 vaccine Janssen Ad26.COVS-2 [recombinant]) and one uses a recombinant S protein (grown in baculovirus infected insect cells) as the antigen with the Matrix-MTM adjuvant (Novavax Nuvaxovid®). The latter adjuvant includes two saponins derived from tree bark. A recently approved booster vaccine (Sanofi Pasteur, VidPrevtyn Beta®) also uses a recombinant S protein but is targeted against the Beta variant and uses a different adjuvant (see the section on variant vaccines).

AstraZeneca COVID-19 ChAdOx1-S vaccine/ Vaxzevria® was used extensively during the primary vaccination campaign but was not routinely used as a booster and is no longer available in the UK. COVID-19 Vaccine Janssen has never been supplied in the UK and a small supply of Nuvaxovid® is currently only available at a limited number of sites.

The Pfizer BioNTech and Moderna COVID-19 vaccines that have been used for the bulk of the UK programme, are nucleoside-modified messenger RNA (mRNA) vaccines. mRNA vaccines use the pathogen's genetic code as the vaccine; this then exploits the host cells to translate the code and then make the target spike protein. The protein then acts as an intracellular antigen to stimulate the immune response (Amanat *et al*, 2020). mRNA is then normally degraded within a few days. Both the Moderna mRNA-1273 and the Pfizer BioNTech COVID-19 BNT162b2 vaccines have been generated entirely in vitro and are formulated in lipid nanoparticles which are taken up by the host cells (Vogel *et al*, 2020, Jackson *et al*, 2020). The Pfizer vaccine was tested in healthy adults between the ages of 18-55 and 65-85 years in phase 1 studies and the BNT162b2 vaccine product at a 30 microgram dose was chosen by Pfizer as the lead candidate in phase 2/3 trials (Walsh *et al*, 2020). The Moderna mRNA-1273 vaccine was tested at three dose levels in those aged 18-55 years and the 100 microgram dose chosen for phase 3 study (Jackson *et al*, 2020).

AstraZeneca COVID-19 vaccine uses a replication deficient chimpanzee adenovirus (ChAd) as a vector to deliver the full-length SARS-CoV-2 spike protein genetic sequence into the host cell (Van Doremalen *et al*, 2020). The adenovirus vector is grown in a human cell-line

(HEK293) ([see chapter 1](#)). ChAd is a non-enveloped virus; the glycoprotein antigen is not present in the vector, but is only expressed once the genetic code within the vector enters the target cells. The vector genes are also modified to render the virus replication incompetent, and to enhance immunogenicity (Garafalo *et al*, 2020). Once the vector is in the nucleus, mRNA encoding the spike protein is produced that then enters the cytoplasm. This then leads to translation of the target protein which acts as an intracellular antigen.

Vaccine effectiveness

Pfizer BioNTech COVID-19 BNT162b2 vaccine (Comirnaty®)

In phase 1/2 human trials, after prime and boost vaccination, neutralising antibodies were comparable or higher than in convalescent patients. Neutralising antibody responses were generally higher in the 18 to 55 year age group compared to the 65 to 85 year age group, but responses were comparable to levels in convalescent patients in both age groups. (Vogel *et al*, 2020).

A phase 3 study was conducted in around 44,000 individuals aged 12 years and above with a second dose delivered between 19 and 42 days. Initial analysis conducted as part of a phase 3 study demonstrated a two-dose vaccine efficacy of 95% in those aged 16 years and over and against symptomatic disease with the wild-type virus. Efficacy was consistent across age, gender, and ethnicity, and in the presence of co-morbidities (including asthma, obesity, diabetes, hypertension and lung disease). In naïve participants aged between 65 and 75 years, and in those aged 75 years and over, the efficacy was 94.7% (95% CI 66.7-99.9%) and 100% (95% CI 13.1-100%) respectively. Efficacy remained high when the analysis included those with evidence of prior immunity. Published efficacy between dose 1 and 2 of the Pfizer BioNTech vaccine was 52.4% (95% CI 29.5-68.4%). Based on the timing of cases accrued in the phase 3 study, most vaccine failures in the period between doses occurred shortly after vaccination, suggesting that short term protection from dose 1 is very high from day 10 after vaccination (Polack *et al*, 2020). Using data for those cases observed between day 15 and 21, efficacy against symptomatic COVID-19 after the first dose was estimated at 89% (95% CI 52-97%). (<https://www.fda.gov/media/144246/download>)

The Pfizer BioNTech COVID-19 vaccine BNT162b2 received approval to supply in the UK from the Medicine and Healthcare products Regulatory Agency (MHRA) on 2 December 2020.

Following a study in over 2000 children aged 12-15 years, which generated additional safety and efficacy data, the approval of a 30 microgram dose was extended to those in this age group in June 2021.

In September 2021, the MHRA approved the use of a 30 microgram dose of Pfizer BioNTech vaccine as a third or reinforcing dose, at least eight weeks after completion of a primary course of either an mRNA or adenovirus vectored vaccine.

Trials in children aged 5-11 years, using a 10 microgram dose of the vaccine formulated for children shown equivalent antibody response and slightly lower reactogenicity than the full adult/adolescent dose (30 micrograms) in those aged 16-25 years. In December 2021, MHRA approved the paediatric formulation of the 10 microgram dose for primary vaccination of children aged 5-11 years.

Studies using a 3 microgram dose in those aged 6 months to 4 years have been undertaken and suggested that, in naive participants, three doses are required to provide similar levels of

antibody to the original strain to that observed after a two dose course in adults. This was associated with a vaccine efficacy (against mainly mild disease) of over 70% during a predominant Omicron era. In December 2022, MHRA approved the infant formulation of the 3 microgram dose for primary vaccination of children aged 6 months to 4 years.

AstraZeneca COVID-19 vaccine (Vaxzevria®)

In phase 1/2 human trials AstraZeneca COVID-19 vaccine was compared with a meningococcal conjugate vaccine (MenACWY) control in healthy adults aged between 18-55 years (Folegatti *et al*, 2020). Preliminary findings showed that neutralising antibodies were induced at day 14 and 28 after the first vaccination and titres increased after a second dose. Specific T cell responses were also induced after a single immunisation and were maintained after the second dose. Final data showed that IgG spike antibody responses and neutralising antibody 28 days after the second dose were similar across the three age cohorts (18–55 years, 56–69 years, and ≥70 years). More than 99% (208/209) of the participants had neutralising antibody responses two weeks after the second dose. Peak T-cell responses were seen 14 days after the first dose and were broadly equivalent in the three age groups (Ramasamy *et al*, 2020). In analysis of over 11,000 patients in the phase 3 study, overall vaccine efficacy against symptomatic disease was 70.4% (95% CI: 54.8–80.6%) (Voysey *et al*, 2020). There were ten cases hospitalised for COVID-19, of which two were severe, all in the control group, suggesting very high protection against severe disease. High protection against hospitalisation was seen from 21 days after dose 1 until two weeks after the second dose, suggesting that a single dose will provide high short term protection against severe disease (Voysey *et al*, 2020). An exploratory analysis of participants who had received one standard dose of the vaccine suggested that efficacy against symptomatic COVID-19 was 73.0% (95% CI: 48.79-85.76%).

The AstraZeneca COVID-19 vaccine received approval to supply in the UK from the MHRA on 30 December 2020.

In September 2021, the MHRA approved the use of AstraZeneca vaccine as a third or reinforcing dose, at least eight weeks after completion of a primary course of AstraZeneca vaccine.

Moderna COVID-19 vaccine (Spikevax®)

In phase 1 testing of the Moderna mRNA-1273 vaccine, all patients seroconverted to IgG by Enzyme-Linked Immunosorbent Assay (ELISA) after the first dose of vaccine. Pseudo-neutralisation and wild virus neutralisation responses were detected in all participants after two 100 microgram doses of the Moderna mRNA-1273. Phase 3 placebo controlled testing in over 30,000 volunteers, showed a vaccine efficacy of 94.1% against symptomatic illness due to wild-type virus. Efficacy was similar in those over 65 years. Vaccine efficacy against severe COVID-19 was 100% (95% CI: 87.0-100%) (Baden *et al*, 2020).

The cumulative case numbers in the phase 3 study showed a clear divergence between the vaccine and placebo groups from about 14 days after the first dose. Re-analysis of the phase 3 data from 15 days after the first dose to the time of the second dose, suggested that efficacy of a single dose was 92.1% against symptomatic illness.

The Moderna vaccine (Spikevax®) was approved for use in the UK in January 2021. Following further studies of safety and efficacy in children, approval was extended to those aged 12-17 years August 2021. In 2022, a half dose (50 micrograms) of Moderna COVID-19 vaccine (Spikevax®) was then approved for those aged 6 to 11 years.

Novavax COVID-19 vaccine (Nuvaxovid®)

In a phase 2 study, a dose of 5 micrograms of the recombinant S protein combined with 50 micrograms of Matrix-M™ adjuvant were chosen (Mallory *et al*, 2021). Large vaccine efficacy studies in the UK (Heath *et al*, 2021) and the USA (Dunkle *et al*, 2021) showed an efficacy of 90% against symptomatic infection with 100% against severe disease. An efficacy of 49% was also shown in a South African trial, during a time when the Beta variant was circulating (Shinde *et al*, 2021).

Novavax vaccine was approved for primary vaccination in February 2022. The vaccine was approved as a heterologous booster in November 2022 (see below).

Reinforcing immunisation

Studies of boosting in the UK have shown that a third adult dose of AstraZeneca, Novavax, Moderna and Pfizer BioNTech vaccines successfully boosted individuals who had been primed with two doses of Pfizer BioNTech or AstraZeneca vaccine around 3 months earlier (Munro *et al*, 2021). Levels of IgG and neutralising antibody, including against Delta variant, were generally higher where an mRNA vaccine was used as either a heterologous or homologous boost, or where AstraZeneca was used as a heterologous boost after a primary course of Pfizer BioNTech. Although levels of antibody were lowest after an AstraZeneca boost in those primed with the same vaccine, levels were as good or better than those seen after the second dose; these antibody levels correlate with high levels of protection against severe disease and death. This finding was confirmed in a study where a third dose of AstraZeneca was given more than six months after the second dose. (Flaxman *et al*, 2021). A separate study using a half dose of Moderna (50 micrograms) in those who had received a primary course of Moderna (100 micrograms) showed good immunogenicity and a rate of reactions similar to the second dose of Moderna. (Choi *et al*, 2021).

Variant vaccines

Following the recognition of the Omicron variant becoming the dominant global circulating strain during 2022, many vaccine manufacturers rapidly developed second generation vaccines that may have broader coverage against SARS-CoV-2 variants. Those approved or approaching licensure have been developed as boosters and have either replaced the spike protein from the original vaccine strain with another strain, or developed a bivalent formulation containing spike protein sequences from both the ancestral strain and a newer variant. Those which use a well established format, such as mRNA vaccines, have been licensed on the basis of immunobridging - i.e. by showing non-inferiority of the neutralising antibody response to the ancestral strain, with potentially higher neutralising antibody response to the variant strain. Bivalent original and Omicron BA.1 mRNA vaccines were approved and became available for booster vaccination in the UK during the autumn of 2022. So far, the emergence of new variants has been too rapid to enable incorporation of a new strain in time to pre-empt an increase in disease. In late 2022, incidence was largely driven by infection with Omicron BA.4 and BA.5. Following the clinical data generated for BA.1 containing vaccines, an mRNA vaccine targeted against the BA.4/5 strains was approved, based on data from animal studies, in the UK in November 2022

Moderna bivalent (Spikevax® bivalent Original/Omicron vaccine) was approved by MHRA for use as a booster in August 2022. This vaccine contains 25 micrograms of mRNA directed against the ancestral strain and 25 micrograms of mRNA against Omicron BA.1. A similar

formulation manufactured by Pfizer BioNTech (Original/Omicron BA.1 Comirnaty®) containing 15 micrograms of mRNA directed against the ancestral strain and 15 micrograms of mRNA against Omicron BA.1 was approved by the MHRA in September 2022.

Around one month after a booster of the Moderna bivalent vaccine, neutralising antibody against the Omicron (BA.1) strain was around 1.6 times the level in those who received a 50 microgram dose of the original Moderna vaccine. Neutralising antibody levels against the ancestral strain were also slightly higher with the bivalent vaccine compared to the original vaccine (geometric mean ratios 1.22 in naive participants), as were the levels of binding antibodies against the alpha, beta, delta and gamma variants (geometric mean ratios between 1.10 and 1.17). The bivalent vaccine did effectively boost neutralising antibody to BA.4/BA.5, but geometric mean levels were lower than against Omicron BA.1 (<https://www.fda.gov/media/159492/download>). Reactogenicity was similar between the original and bivalent vaccine. A similar bivalent vaccine, targeting the BA.4/5 strain instead of the BA.1 Omicron strain, was approved in the UK in February 2023. Clinical studies have shown that neutralising antibody against the Omicron BA.4/5 strain are higher than after a booster of the original vaccine (Chalkias *et al*, 2022).

For Pfizer BioNTech, a booster of the bivalent vaccine produced neutralising antibody response against the Omicron (BA.1) strain around 1.56 times the level in those who received a 30 microgram dose of the original Pfizer BioNTech vaccine. Neutralising antibody levels against the ancestral strain were similar to those after the original vaccine. Neutralising antibody against BA.4/BA.5 was detected after the bivalent vaccine although geometric mean levels were around 3 fold lower than against Omicron BA.1. (<https://www.fda.gov/media/159496/download>). Reactogenicity was similar between the original and bivalent vaccine. This product has a similar reactogenicity profile to the initial vaccine and subsequent human trials has demonstrated superior neutralising response to BA.4/5 when compared to the original vaccine (Zou *et al*, 2022).

The adult/adolescent bivalent BA.4/5 mRNA products are expected to be available in the UK by spring 2023. A paediatric formulation of the Pfizer BioNTech bivalent vaccine, containing 5 micrograms of mRNA targeting the original strain and 5 micrograms of mRNA targeting the BA.4/5 strains was approved in the UK in February 2023. This product will be suitable for children aged 5 to 11 years and may become available later in 2023.

Sanofi Pasteur COVID-19 vaccine (VidPrevtyn Beta®)

In December 2022, a vaccine based on a recombinant spike protein from the Beta variant was approved in the UK. The vaccine (VidPrevtyn Beta®) is manufactured by Sanofi Pasteur and uses the AS03 adjuvant system. This adjuvant was developed for use during pandemics and was employed as part of an H1N1v influenza vaccine in 2010-2011. It is similar to the MF59 used in adjuvanted influenza vaccine, in that it contains squalene, but it also contains DL- α -tocopherol - a form of vitamin E which helps to modulate the innate immune system and therefore further enhance the immune response. VidPrevtyn Beta® had shown efficacy as a primary vaccine and was then studied as a booster in adults who had received primary vaccination with either mRNA or adenovirus vaccines. Although targeted against the Beta variant, 91 days after vaccination, a booster dose of Sanofi Pasteur COVID-19 vaccine (VidPrevtyn Beta®) achieved similar levels of pseudo neutralising antibody against the original, Beta, Delta, BA.1 and BA.4/5 strains as those seen after the Moderna and Pfizer bivalent vaccines targeting the BA.1 strain (Spikevax® bivalent Original/Omicron and Comirnaty® Original/Omicron BA.1). <https://www.fda.gov/media/164809/download>

Real world effectiveness

Vaccine effectiveness data from the UK has now been generated with successive SARS-CoV-2 variants. A single adult dose of either the Pfizer BioNTech or the AstraZeneca vaccines were shown to provide modest protection against symptomatic disease due to Alpha variant; with single vaccinated cases around 40% less likely to require hospital admission or to die (Lopez Bernal *et al*, 2021a). This was consistent with protection of around 80% against hospitalisation as seen in local studies (Vasileiou *et al*, 2021, AvonCAP, 2021). Protection against infection was also seen in healthcare workers, where a single dose of Pfizer BioNTech vaccine provided more than 70% protection against both symptomatic and asymptomatic infection (Hall *et al*, 2021a), and in care home residents where a single dose of either Pfizer BioNTech or AstraZeneca vaccines reduced the risk of infection by around 60% (Shrotri *et al*, 2021). The observed reduction in both symptomatic and asymptomatic infections suggested that vaccination had potential to reduce transmission; this was supported by a Scottish study during the pre-Delta era that showed a 30% reduction in risk of infection in the household members of vaccinated compared to unvaccinated healthcare workers after a single dose of the Pfizer BioNTech vaccine. (Shah *et al*, 2021).

Higher levels of protection against symptomatic disease due to Alpha variant were observed after the second dose for both Pfizer BioNTech (Lopez Bernal *et al*, 2021b) and AstraZeneca vaccines.

Following the introduction of the Delta variant to the UK in April 2021, further updates to the analysis of real world effectiveness have been undertaken (Lopez Bernal *et al*, 2021b). Protection against symptomatic infection with the Delta variant was slightly lower than against Alpha, particularly after a single dose. Protection against hospitalisation, however, was maintained with two doses of the AstraZeneca and Pfizer BioNTech vaccines providing over 90% short term protection against this outcome. (Stowe *et al*, 2021).

Since the emergence of the Omicron variant, vaccine effectiveness data confirms that protection against symptomatic disease from current vaccines is lower than for Delta. (Andrews *et al*, 2022a). Vaccination does provide higher levels of protection against hospitalisation due to Omicron. A summary of the most recent data on real world effectiveness for each variant is now published and updated regularly. <https://www.gov.uk/government/publications/covid-19-vaccine-surveillance-report>.

Duration of protection

Israel was the first country to demonstrate waning protection from Pfizer BioNTech vaccine showing a decline in protection, even against severe disease, at around 6 months (Goldberg *et al*, 2021). In the USA, protection against hospitalisation for Pfizer BioNTech and Moderna vaccines remained high (around 84%) between 3 and 6 months (Tenforde *et al*, 2021).

Updated UK analysis to late August 2021 suggests that protection against symptomatic infection due to the Delta variant appears to decline after the second dose, although remains above 50% overall after 5 months. (Andrews *et al*, 2022b) Levels of protection from AstraZeneca are lower than that seen after Pfizer BioNTech and remain around 20% lower after 5 months. In contrast, protection against hospitalisation and death from Delta variant appears to be well sustained, remaining around 85% at six months after primary vaccination with both AstraZeneca and Pfizer BioNTech vaccines. The decline in protection appears to be mainly driven by older people (over 65 years) and those with

clinical risk factors (including immunosuppression). For Omicron, protection from primary vaccination appears to decline to very low levels by six months after all three vaccines used in the UK. Waning of protection after booster doses is discussed in the section on reinforcing immunisation.

Reinforcing doses

In Israel, administration of a booster dose of Pfizer BioNTech to adults who had received a primary course of the same vaccine, has been associated with a major reduction in the risk of both confirmed and severe disease due to COVID-19. (Bar-On *et al*, 2021).

In the UK, early data showed a major increase in levels of protection after the first booster dose against both symptomatic disease and hospitalisation due to the Delta variant (Andrews *et al*, 2022c). Vaccine effectiveness data for Omicron confirms that protection against symptomatic disease soon after an mRNA booster dose increases to around 70-75% regardless of the primary vaccine series. Levels of protection after an AstraZeneca booster in those who received the same vaccine as a primary course were only slightly lower than those seen after the mRNA boosters. (Andrews *et al*, 2022b). More recent analysis confirms that protection against symptomatic disease after an mRNA booster declines substantially by three months after the dose was given. For the small number of individuals who received AstraZeneca as a booster, levels of protection against symptomatic Omicron infection appeared to be similar or slightly lower than those after an mRNA booster.

Protection against hospitalisation after an mRNA booster reaches over 90% in the two weeks after vaccination and then declines towards a stable plateau of around 60% by six months. Data real world effectiveness for more severe outcomes is published and updated regularly. <https://www.gov.uk/government/publications/covid-19-vaccine-surveillance-report>.

Variant vaccines

Although current vaccines offer lower levels of protection against mild disease caused by variants with mutations on the spike protein, protection against more severe COVID-19 appears to be relatively less affected. Higher levels of antibody against the original spike protein do appear to provide higher levels of protection against symptomatic infection due to distant variants.

The bivalent and monovalent variant vaccines were initially approved on the basis of neutralising antibody compared to the original vaccines. Overall the improvements in neutralising antibody levels are modest and likely to translate to small improvements in protection against new variants. Early evidence on the real world effectiveness of the bivalent BA.1 vaccines against hospitalisation for the current circulating Omicron strains (mainly due to BA.4/5) suggests that a booster dose increases the level of protection in the short term by an additional 60% - similar to what was observed after previous booster doses.

<https://www.gov.uk/government/publications/covid-19-vaccine-weekly-surveillance-reports>

Safety

Pfizer BioNTech COVID-19 BNT162b2 vaccine (Comirnaty®)

Local reactions at the injection site are fairly common after Pfizer BioNTech COVID-19 vaccine, primarily pain at the injection site, usually without redness and swelling. Systemic events reported were generally mild and short lived (Walsh *et al*, 2020). In the final safety analysis of over 21,000 participants 16 years and older, the most common events were injection site pain (>80%), fatigue (>60%), and headache (>50%). Myalgia, arthralgia and chills were also common with fever in 10-20%, mainly after the second dose. Most were classified as mild or moderate. Lymphadenopathy in the axillary, supraclavicular or cervical nodes on the same side as the injection was reported in less than 1% (Polack *et al*, 2020).

Side effects were less common in those aged over 55 than those aged 16 to 55 years. Severe systemic effects, defined as those that interfere with daily activity, included fatigue in 4% and headache in 2%. There was no signal to suggest that prior vaccination led to enhanced disease with only 1 case of severe COVID-19 in the 8 vaccine failures (Polack *et al*, 2020).

During post marketing surveillance, a number of cases of myocarditis and pericarditis have been reported after Pfizer BioNTech vaccine. The reported rate appears to be highest in those under 25 years of age and in males, and after the second dose. Onset is within a few days of vaccination and most cases are mild and have recovered without any sequelae. The MHRA has advised the benefits of vaccination still outweigh any risk in most individuals.

Since the widespread use of the vaccine, a number of other conditions have been reported after vaccination and have been or are about to be added to the Summary of Product Characteristics (SmPC). This includes reports of heavy menstrual bleeding (in most cases temporary and non-serious) and extensive swelling of the vaccinated limb. (<https://www.gov.uk/government/publications/regulatory-approval-of-pfizer-biontech-vaccine-for-covid-19>) A very small number of cases of Guillain-Barre Syndrome (GBS) have been reported after Pfizer-BioNTech vaccination but these reports have not reached the number expected to occur by chance in the immunised population.

Safety data reported from other countries after routine use of the paediatric dose of Pfizer BioNTech vaccine confirms the finding of lower rates of all reactions when compared to a full dose in older children and young people.

Moderna COVID-19 vaccine (Spikevax®)

A high proportion (more than 75%) of vaccine recipients had localised pain at the injection site after both dose 1 and dose 2 of the Moderna mRNA-1273 vaccine. Redness and swelling were also seen after the second dose and local pain tended to last longer (around 3 days). Mild systemic effects were also common, including headache, fatigue, joint and muscle aches and chills. Systemic events were more severe after dose 2 and fever was only seen after dose two. Both local and systemic reactions were less common in older participants (Baden *et al*, 2020). Adverse events were less common in those with pre-existing SARS-CoV-2 antibody. Axillary lymphadenopathy on the same side as the injection site was detected in more than one in ten recipients.

There were no cases of severe COVID-19 disease in the vaccine group, and thus no signal for enhanced disease (Baden *et al*, 2020).

During post-marketing surveillance, a number of cases of myocarditis and pericarditis have been reported after Moderna vaccine. The reported rate appears to be highest in those under 25 years of age and in males, and after the second dose. Onset is within a few days of vaccination and most cases are mild and have recovered without any sequelae. The MHRA has advised the benefits of vaccination still outweigh any risk in most individuals.

Since the widespread use of the vaccine, a number of other conditions have been reported after vaccination and have been recently added to the Summary of Product Characteristics (SmPC) this includes heavy menstrual bleeding (in most cases temporary and non-serious), capillary leak syndrome, extensive swelling of the vaccinated limb and urticaria. (<https://www.gov.uk/government/publications/regulatory-approval-of-covid-19-vaccine-moderna>) A very small number of cases of Guillain-Barre Syndrome (GBS) have been reported after Moderna vaccination but these reports have not reached the number expected to occur by chance in the immunised population.

AstraZeneca COVID-19 vaccine (Vaxzevria®)

From early phase trials, mild pain and tenderness at the injection site was common with AstraZeneca COVID-19 vaccine occurring in 88% of 18-55 year olds, 73% of 56-69 year olds and 61% of people aged 70 years or over; similar levels were reported after each dose. Short lived systemic symptoms including fatigue and headache were also common but decreased with age, being reported in 86%, 77%, and 65% of those aged 18-55, 56-69 and 70 years or over respectively; most of these were classified as mild or moderate. These reactions were unusual after the second dose (Ramasamy *et al*, 2020). Mild fever (>38°C) was recorded in the first 48 hours for around a quarter of younger participants but was not reported in those over 55 years of age or in any age group after the second dose (Ramasamy *et al*, 2020). Fever is modified by the prophylactic use of paracetamol, which does not affect the immune response to this vaccine (Folegatti *et al*, 2020). In the phase 3 study, injection site reactions, mild fever, headache, myalgia and arthralgia occurred in more than 10% of vaccinees. Less than 1% reported lymphadenopathy or an itchy rash. Only one serious adverse event was reported as possibly linked to the vaccine; this was a case of transverse myelitis which occurred 14 days after dose 2. There was no signal to suggest that prior vaccination led to enhanced disease (Voysey *et al*, 2020).

During post-marketing surveillance, a very rare condition involving serious thromboembolic events accompanied by thrombocytopenia, has been reported after AstraZeneca vaccination. The condition presents with unusual venous thrombosis, including cerebral venous sinus thrombosis, portal vein thrombosis, and sometimes arterial thrombosis, with low platelet count and high D-dimer measurements. The condition has similarities to heparin-induced thrombocytopenia and thrombosis (HITT or HIT type 2) and patients usually have positive antibody to platelet factor 4. The majority of the events occurred between 5 and 16 days following vaccination (Greinacher *et al*, 2021).

The current reported rate of this event in the UK is around 15 cases per million after the first dose, although a higher incidence is seen in younger individuals. After the second dose the reported rate is much lower, particularly in younger individuals. Overall, the Joint Committee on Vaccination and Immunisation (JCVI), MHRA and the WHO concluded that the benefits of vaccination outweighed this small risk for adults aged 40 years and over, and those at higher clinical risk.

GBS has been reported very rarely within six weeks of AstraZeneca vaccination, and rates appear to be higher than the background rates. This risk would equate to about 5.8 extra cases of GBS per million doses in the six weeks following the first dose of AstraZeneca vaccine, based on a unpublished UK study (<https://www.ucl.ac.uk/ion/news/2022/may/rise-guillain-barre-syndrome-following-astrazeneca-vaccine>). There was no evidence of a higher rate of reporting in individuals who had had a previous episode of GBS.

A small number of cases of capillary leak syndrome have been reported across Europe within 4 days of AstraZeneca vaccination. Around half of those affected had a history of capillary leak syndrome.

Cases of thrombocytopenia (without accompanying thrombosis) have been reported rarely in the first four weeks after receiving AstraZeneca vaccination. Some of these cases have occurred in individuals with a history of immune thrombocytopenia (ITP)

Novavax COVID-19 vaccine (Nuvaxovid®)

Side effects after the vaccine are similar to other COVID-19 vaccines, with slightly lower rates of local reactions and systemic effects when compared to mRNA vaccines. Around 50% of dose 1 and 70% of dose 2 recipients reporting pain and /or tenderness at the injection site and around 40-50% report systemic symptoms including fatigue, malaise, headache and muscle pain, with rates of fever below 10%. Overall, there was a higher incidence of adverse reactions in younger age group (18-64 years).

Small numbers of cases of myocarditis or pericarditis were reported across the trials and in post-marketing follow up. Myocarditis and pericarditis have now been added to the list of possible side effects in the SmPC.

Sanofi Pasteur COVID-19 vaccine (VidPrevtyn Beta®)

When given as a first booster in individuals previously vaccinated with mRNA or adenovirus vector vaccines the most common side effects were injection site pain (76.2%), headache (41.4%), myalgia (37.8%), malaise (33.0%), arthralgia (28.7%), and chills (19.9%). Most side effects were short lived (less than 3 days) and of mild to moderate severity.

Reinforcing immunisation

In the UK study, all boosters led to short term local and systemic reactions, similar to those seen after the primary course, including local pain, fatigue, headache and muscle pain. Rates of reactions were higher with heterologous than homologous boosters and in those aged under 70 years when compared to older recipients. Rates of local and systemic symptoms were higher where a full dose of Moderna was used to boost those who had received either AstraZeneca or Pfizer BioNTech for the primary course and when AstraZeneca was used to boost those who had Pfizer BioNTech as a primary course, when compared to Novavax or Pfizer BioNTech after either primary vaccination. Using a half dose (50 micrograms) of Moderna is expected to have a lower rate of side effects (including myocarditis) than a full dose.

Following implementation of booster doses, the nature of adverse events reported has been similar to that reported after the first two doses of the COVID-19 vaccines. Reports of suspected adverse events following COVID-19 boosters given at the same time as seasonal flu vaccines are also similar to that when the vaccines are given individually. There have been a small number of reports of suspected myocarditis and pericarditis following booster doses with Pfizer/BioNTech Moderna COVID-19 vaccines.

Storage

Pfizer BioNTech adult/adolescent formulation (Comirnaty®) is supplied frozen in packs of 195 vials. Frozen vials should be transferred to thaw at 2°C to 8°C; a 195 vial pack may take 3 hours to thaw. Alternatively, frozen vials may also be thawed for 30 minutes at temperatures up to 25°C for immediate use. After thawing, undiluted vaccine can be stored for up to 31 days at 2°C to 8°C.

Pfizer BioNTech paediatric formulations (Comirnaty®10 and Comirnaty® Original/Omicron BA.4/5 5+5) and (Comirnaty® 3) are supplied frozen in 10-vial packs. The vaccines can be thawed at 2°C to 8°C for 4 hours or individual vials can be thawed at room temperature (up to 30°C) for 30 minutes. Unopened vials can be stored for up to 10 weeks at 2°C to 8°C. Prior to use, the unopened vial can be stored for up to 12 hours at temperatures up to 30°C.

Pfizer BioNTech bivalent booster formulations (Comirnaty® Original/Omicron BA.4/5 or Comirnaty® Original/Omicron BA.1) are supplied frozen in 10 vial packs and should be stored - 90°C to -60°C. Packs can be thawed at 2 °C to 8 °C for 6 hours or individual vials can be thawed at room temperature for 30 minutes. Once thawed, the vaccine should not be re-frozen, but can be stored and transported at 2 °C to 8 °C for 10 weeks (within the overall shelf life).

The AstraZeneca vaccine (Vaxzevria®) should be stored at 2°C to 8°C. The vaccine does not contain any preservative. After first opening the vial, it should be used within 6 hours. The vaccine may be stored between 2°C and 25°C during this period. After this time, the vial must be discarded.

Moderna COVID-19 vaccine (Spikevax®) vials should be stored frozen (between -25° to -15°C for the original vaccine and -50° to -15° C for the bivalent vaccine) Once thawed, the vaccine may be stored refrigerated at 2°C to 8°C protected from light for up to 30 days if not punctured. The unopened vials are stable for 24 hours at 8° to 25°C.

Novavax COVID-19 vaccine (Nuvaxovid®) should be stored at 2°C to 8°C and protected from light. The vaccine is stable up to 12 hours at 25°C.

The Sanofi Pasteur COVID-19 vaccine (VidPrevtyn Beta®) should be stored in a refrigerator at 2°C to 8°C. After mixing the SmPC advises returning the product to the fridge, protecting it from light and then discarding after six hours. MHRA review of quality data have shown that the mixed antigen/adjuvant for VidPrevtyn Beta is stable at 23-27°C for several hours. As any risk of microbial growth would be minimal within a few hours of mixing, UKHSA advises that, in the clinic setting, the 10 doses of VidPrevtyn Beta should be used without returning to the fridge as soon as practicably possible, ideally within one clinic session (2-3 hours). In other settings, such as domiciliary vaccination, the product may be returned to the fridge or cool box between each vaccination, but must be discarded after 6 hours.

Presentation

A pack of the Pfizer BioNTech adult/adolescent formulation (Comirnaty®) contains 195 vials with a purple cap; each vial contains a minimum of 6 doses per vial (1170 doses per pack). A pack of the Pfizer BioNTech paediatric vaccine (Comirnaty®10 and Comirnaty® Original/Omicron BA.4/5 5+5) contains 10 vials with an orange cap, each containing sufficient for 10 doses. The Pfizer BioNTech infant and pre-school vaccine (Comirnaty® 3) contains 10 vials

with a maroon cap, each containing sufficient for 10 doses. All three vaccines are supplied with 0.9% sodium chloride diluent for injection in plastic ampoules. After dilution, the vaccines should be used as soon as practically possible, and within the maximum time as outlined in the SpC. The adult/ adolescent vaccine should be kept at 2°C to 25°C and used within 6 hours; the paediatric vaccines should be kept at 2°C to 30°C and used within 12 hours. Any unused vaccine should be discarded.

The Pfizer BioNTech bivalent boosters (Comirnaty® Original/Omicron BA.4/5 or Comirnaty® Original/Omicron BA.1) come in packs of 10 vials with a grey cap. The vaccine does not require dilution and each vial contains 6 doses. After the first puncture, the vaccine should be used as soon as practically possible, stored at 2°C to 30°C and used within 12 hours.

The AstraZeneca vaccine (Vaxzevria®) is supplied in packs of 10 vials. Each vial contains 8 or 10 doses of vaccine, and is a colourless to slightly brown, clear to slightly opaque liquid.

Moderna COVID-19 vaccine (Spikevax®) is supplied in multidose vials with a red cap and containing 10 primary doses of 0.5ml or 20 booster doses of 0.25ml. The Moderna bivalent boosters (Spikevax® bivalent Original/Omicron BA.4/5 or Spikevax® bivalent Original/Omicron) are supplied in multidose vials containing either 5 or 10 doses of 0.5ml and have a blue cap.

Novavax COVID-19 vaccine (Nuvaxovid®) is supplied in a multidose vial which contains 10 doses of 0.5 ml. Each vial contains a colourless to slightly yellow, clear to mildly opalescent fluid. The multidose vial should be gently swirled and inspected for particles, and used within 6 hours of first puncture.

The Sanofi Pasteur COVID-19 vaccine (VidPrevtyn Beta®) is supplied as two multi-dose vials, (containing 2.5ml of adjuvant and antigen each) that must be mixed before injection. The antigen solution is a colourless, clear liquid and the adjuvant emulsion is a whitish milky liquid. After mixing the vial contains 10 doses of 0.5ml.

Dosing and schedule

All COVID-19 vaccines

For both adenovirus vector and mRNA vaccines, there is evidence of better immune response and/or protection where longer intervals are used between doses in the primary schedule. (Amirthalingam *et al*, 2021, Payne *et al*, 2021, Voysey *et al*, 2021).

Based on this evidence, longer intervals are likely to provide more durable protection. For primary vaccination, JCVI is currently recommending a minimum interval of eight weeks between the first two doses for all of the approved COVID-19 vaccines. This interval should be followed in all eligible adults and children in clinical risk groups. Operationally, using the same minimum interval for all products will simplify supply and booking, and will help to ensure a good balance between achieving rapid and long-lasting protection.

The infant and pre-school formulation of the Pfizer-BioNTech vaccine (Comirnaty® 3) is approved as a three dose schedule. Most UK children have already been exposed to SARS-CoV-2 infection, and separating the first two doses is likely to make a better immune response than when delivered 3 weeks apart. Therefore JCVI has advised that eligible children aged 6 months to 4 years should be offered two doses at an interval of eight weeks. Those with severe immunosuppression may be considered for additional doses, as outlined in the section on additional doses for those with severe immunosuppression. Other clinical risk groups are expected to become eligible for further doses during future seasonal campaigns.

For those under 18 years who are not in a high risk group a 12 week interval was recommended by JCVI. This was based on precautionary advice from the JCVI based on emerging evidence of a lower rate of myocarditis in countries that use schedules of 8 to 12 weeks.

The main reason to vaccinate at a shorter interval (that is below eight weeks) could be for those about to commence immunosuppressive treatment. In these individuals, timing of administration should take account of the ability of the immune system to respond, and to accommodate this the interval may be reduced to the minimum approved as outlined below. JCVI has also recommended additional doses of vaccination for individuals who were severely immunosuppressed when they received their COVID-19 vaccination (see the section on additional doses for those with severe immunosuppression).

Pfizer BioNTech COVID-19 vaccines (Comirnaty[®], and Comirnaty[®] Original/ Omicron BA.1 and Comirnaty[®] Original/ Omicron BA.4/5)

For those aged 12 years and above, the dose of the bivalent vaccines is 0.3 ml of the undiluted (ready to use) Pfizer BioNTech COVID-19 vaccine containing 15 plus 15 micrograms. A 0.3 ml dose of the diluted original Pfizer BioNTech COVID-19 vaccine, containing 30 micrograms is an acceptable alternative.

The primary course is approved to be administered in two doses, a minimum of 21 days apart, but current JCVI advice is that the doses should be given at an eight week minimum interval.

A booster dose of Pfizer BioNTech COVID-19 vaccine is 30 micrograms contained in 0.3 ml of the diluted vaccine.

Paediatric Pfizer BioNTech COVID-19 vaccine (Comirnaty[®]10 and Comirnaty[®] Original/ Omicron BA.4/5 5+5)

For children aged 5-11 years, the dose of Pfizer BioNTech COVID-19 vaccine is 10 micrograms. The paediatric formulations, Comirnaty[®]10 and Comirnaty[®] Original/Omicron BA.4/5 (5 + 5 micrograms) are supplied in a multidose vial, with each vial containing 10 doses of 0.2 mL (after dilution with 1.3ml of saline). The paediatric formulation should be used in this age group, although 0.1ml of the adult/adolescent vaccines may be used in exceptional circumstances only.

The primary course is approved to be administered in two doses, a minimum of 21 days apart, but current JCVI advice is that the doses should be given at an eight week minimum interval.

Infant Pfizer BioNTech COVID-19 vaccine (Comirnaty[®]3)

For children aged 6 months to 4 years, the dose of Pfizer BioNTech COVID-19 vaccine is 3 micrograms. The infant formulation Comirnaty[®]3 is supplied in a multidose vial with a maroon cap, with each vial containing 10 doses of 0.2 ml injection volume (after dilution with 2.2 mL of saline).

The primary course is approved to be administered in three doses, with the second dose a minimum of 21 days after the first and the third dose after a further eight weeks (see above). Current JCVI advice is that two doses should be given eight weeks apart.

Novavax COVID-19 vaccine (Nuvaxovid®)

The dose of Novavax COVID-19 vaccine is 0.5 ml. The primary course is two doses, a minimum of 3 weeks apart, but current JCVI advice is that the doses should be given at an eight week minimum interval.

Moderna COVID-19 vaccines (Spikevax®, Spikevax® bivalent Original/Omicron and Spikevax® bivalent Original/Omicron® BA.4/5)

For primary vaccination, including third doses for those with severe immunosuppression, JCVI is now recommending using a 0.5ml reinforcing dose of the bivalent Moderna COVID-19 vaccine containing 25 + 25 microgrammes. A 0.5ml dose of the original Moderna COVID-19 vaccine, containing 100 micrograms is an acceptable alternative.

The primary course is approved to be administered in two doses, a minimum of 28 days apart, but current JCVI advice is that the doses should be given at an eight week minimum interval.

Reinforcing doses

For those eligible for vaccination in both the spring and the autumn campaigns, a single booster dose is usually scheduled at around six months from the previous dose. To facilitate operational delivery, boosting can take place, however, from three months after the previous dose. This interval applies to any booster and regardless of the product given for the previous doses.

Pfizer BioNTech bivalent vaccines (Comirnaty® Original/Omicron BA.4/5 or Comirnaty® - Original/Omicron BA.1)

A booster dose of the bivalent Pfizer BioNTech vaccine contains 15 plus 15 micrograms in 0.3ml of the undiluted (ready to use) vaccine.

Moderna bivalent COVID-19 vaccines (Spikevax® bivalent Original/Omicron BA.4/5 or Spikevax® bivalent Original/Omicron)

A single 0.5ml booster dose of the bivalent vaccines (containing 25 plus 25 micrograms) is recommended.

Novavax COVID-19 vaccine (Nuvaxovid®)

The dose of Novavax COVID-19 vaccine is 0.5ml, administered as a single booster.

Sanofi Pasteur COVID-19 vaccine (VidPrevtyn Beta®)

The dose of Sanofi Pasteur COVID-19 vaccine is 0.5ml, administered as a single booster.

Administration

Vaccines are routinely given intramuscularly into the upper arm or anterolateral thigh. This is to reduce the risk of localised reactions, which are more common when vaccines are given subcutaneously (Mark *et al*, 1999; Zuckerman, 2000; Diggle and Deeks, 2000).

Pfizer BioNTech COVID-19 vaccine (Comirnaty®) and the equivalent bivalent booster vaccines (Comirnaty® Original/Omicron BA.4/5 or Comirnaty® Original/Omicron BA.1), should be administered as an intramuscular injection into the deltoid muscle. A 1ml syringe with a 23g x 25mm needle will be provided for administration. A separate needle and syringe should be used for each individual. The vial should be discarded if the solution is discoloured or visible particles are observed.

AstraZeneca COVID-19 vaccine (Vaxzevria®) is administered as a single dose of 0.5ml intramuscular injection into the deltoid muscle. A 1ml syringe with a 23g/25g x 25mm needle will be provided for administration. The vaccine should be inspected visually for particulate matter and discolouration prior to administration. The vial should be discarded if the solution is discoloured or visible particles are observed. The vial should not be shaken. A separate needle and syringe should be used for each individual. It is normal for liquid to remain in the vial after withdrawing the final dose.

Moderna COVID-19 vaccine (Spikevax®) and the equivalent bivalent booster vaccines (Spikevax® bivalent Original/Omicron BA.4/5 or Spikevax® bivalent Original/Omicron) should be administered by intramuscular injection into the deltoid muscle. A 1ml syringe with a 23g x 25mm needle will be provided for administration. A separate needle and syringe should be used for each individual. It is normal for liquid to remain in the vial after withdrawing the final dose.

Novavax COVID-19 vaccine (Nuvaxovid®) should be administered by intramuscular injection, preferably in the deltoid muscle of the upper arm. A separate needle and syringe should be used for each individual. A 1ml syringe with a 23g/25g x 25mm needle will be provided for administration.

Sanofi Pasteur COVID-19 vaccine (VidPrevtyn Beta®) is a single dose of 0.5 ml and should be administered by intramuscular injection, preferably in the deltoid muscle of the upper arm. A 21g safety needle and 3ml syringe will be provided for preparation of the product. A 23g x 25mm needle and 1ml syringe will be provided for administration. A separate needle and syringe should be used for each individual.

Individuals with bleeding disorders may be vaccinated intramuscularly if, in the opinion of a doctor familiar with the individual's bleeding risk, vaccines or similar small volume intramuscular injections can be administered with reasonable safety by this route. If the individual receives medication/treatment to reduce bleeding, for example treatment for haemophilia, intramuscular vaccination can be scheduled shortly after such medication/treatment is administered. Individuals on stable anticoagulation therapy, including individuals on warfarin who are up-to-date with their scheduled INR testing and whose latest INR is below the upper level of the therapeutic range, can receive intramuscular vaccination. A fine needle (23 or 25 gauge) should be used for the vaccination, followed by firm pressure applied to the site without rubbing for at least 2 minutes (Advisory Committee on Immunization Practices 2019). The individual/parent/carer should be informed about the risk of haematoma from the injection.

Disposal

Equipment used for vaccination, including used vials, ampoules or syringes, should be disposed of by placing them in a proper, puncture-resistant 'sharps box' according to local authority regulations and guidance in Health Technical Memorandum 07-01: Safe management of healthcare waste (Department of Health, 2013).

AstraZeneca COVID-19 vaccine (Vaxzevria®) contains genetically modified organisms (GMOs). Sharps waste and empty vials should be placed into yellow lidded waste bins and sent for incineration; there is no need for specific designation as GMO waste. An appropriate virucidal disinfectant should be available for managing spills in all settings where vaccination is administered. Potentially contaminated gloves and aprons can be disposed in yellow/black striped bags for offensive waste.

The COVID-19 pandemic immunisation programme

The main objective of the COVID-19 pandemic immunisation programme was to protect those who are at highest risk from serious illness or death. The JCVI ranked the eligible groups according to risk. For the first phase this ranking was based on the risk of COVID-19 specific mortality, with the second phase concerned with prevention of hospitalisation.

Phase 1 recommendations for primary vaccination

Evidence from the UK indicates that the risk of poorer outcomes from COVID-19 infection increases dramatically with age in both healthy adults and in adults with underlying health conditions. Those over the age of 65 years have by far the highest risk, and the risk increases steeply with age. Residents in care homes for older adults have been disproportionately affected by the COVID-19 pandemic. Table 2 sets out the initial JCVI advice on priority groups for primary COVID-19 vaccination.

Table 2 – Phase 1 priority groups for primary vaccination advised by the Joint Committee on Vaccination and Immunisation

Priority group	Risk group
1	Residents in a care home for older adults Staff working in care homes for older adults
2	All those 80 years of age and over Frontline health and social care workers
3	All those 75 years of age and over
4	All those 70 years of age and over Individuals aged 16 to 69 in a high risk group ¹
5	All those 65 years of age and over
6	Adults aged 16 to 65 years in an at-risk group (Table 3)
7	All those 60 years of age and over
8	All those 55 years of age and over
9	All those 50 years of age and over

Definitions of individuals aged 16 years and over at clinical high risk (priority groups 4 and 6)

People previously defined as clinically extremely vulnerable (CEV) were considered to be at high risk of severe illness from COVID-19 (<https://www.gov.uk/government/publications/guidance-on-shielding-and-protecting-extremely-vulnerable-persons-from-covid-19/guidance-on-shielding-and-protecting-extremely-vulnerable-persons-from-covid-19#cev>) and these patients were flagged on the GP system. A hospital clinician or GP was able to add a patient to the list, based on their clinical judgement, because they considered them to be at very high risk of serious illness from COVID-19. All patients who were on the CEV list also fell into group 6, which included a broader range of disease categories that JCVI advised would constitute a higher clinical risk for COVID-19 vaccination (tables 3 and 4). When the shielding programme ended groups 4 and 6 were formally merged.

¹ Previously known as clinically extremely vulnerable

The examples in tables 3 and 4 are not exhaustive, and, within these broad groups, the prescriber may need to apply clinical judgment to take into account the risk of COVID-19 exacerbating any underlying disease that a patient may have, as well as the risk of serious illness from COVID-19 itself. A more comprehensive list of eligible diagnoses, and the appropriate clinical codes, can be found in the link at the end of the chapter.

In December 2021, following the recognition of pregnancy as a risk factor for severe COVID-19 infection and poor pregnancy outcomes during the Delta wave, pregnancy was added to the the clinical risk groups (see table 3).

Definitions of front line staff aged 16 years and over

Vaccination in phase one was also recommended for certain staff groups (see definitions below). The objective of occupational immunisation of health and social care staff was to protect those workers at high risk of exposure who may also expose vulnerable individuals whilst providing care. There is limited evidence that vaccination leads to a reduction in transmission, although a small effect may have major additional benefit for staff who may expose multiple vulnerable patients and other staff members.

Staff involved in direct patient care

This includes staff who have frequent face-to-face clinical contact with patients and who are directly involved in patient care in either secondary or primary care/community settings. This includes doctors, dentists, midwives and nurses, paramedics and ambulance staff, pharmacists, optometrists, occupational therapists, physiotherapists and radiographers. It should also include those working in independent, voluntary and non-standard healthcare settings such as hospices, and community-based mental health or addiction services. Staff working on the COVID-19 vaccination programme, temporary staff, students, trainees and volunteers who are working with patients are also be included.

Non-clinical staff in secondary or primary care/community healthcare settings

This includes non-clinical ancillary staff who may have social contact with patients but are not directly involved in patient care. This group includes receptionists, ward clerks, porters and cleaners.

Laboratory and pathology staff

Hospital-based laboratory and mortuary staff who frequently handle SARS-CoV-2 or collect or handle potentially infected specimens, including respiratory, gastrointestinal and blood specimens should be eligible as they may also have social contact with patients. This may also include cleaners, porters, secretaries and receptionists in laboratories. Frontline funeral operatives and mortuary technicians/embalmers are both at risk of exposure and likely to spend a considerable amount of time in care homes and hospital settings where they may also expose multiple patients.

Staff working in non-hospital-based laboratories and those academic or commercial research laboratories who handle clinical specimens or potentially infected samples will be able to use effective protective equipment in their work and should be at low risk of exposure, and of exposing vulnerable patients.

Frontline social care workers (priority groups 1 and 2) aged 16 years and over

This includes those working in long-stay residential and nursing care homes or other long-stay care facilities where rapid spread is likely to follow introduction of infection and cause high morbidity and mortality.

It also includes other front-line social care workers who regularly provide close personal care to those who are clinically vulnerable.

Those clinically vulnerable to COVID-19 are defined by the JCVI priority groups:

- a) children of any age with severe neuro-disability, severe or profound and multiple learning disabilities (including Down's syndrome and those on the learning disability register) or immunosuppression (as defined in table 4),
- b) adults who have underlying health conditions leading to greater risk of disease or mortality as defined in table 3,
- c) those of advanced age.

Definitions of other high-risk groups**Household contacts**

Individuals who expect to share living accommodation on most days (and therefore continuing close contact is unavoidable) with people who are immunosuppressed (defined as immunosuppressed in tables 3 or 4).

Carers

Those who are eligible for a carer's allowance, or those aged 16 years and over who are the sole or primary carer of an elderly or disabled person who is at increased risk of COVID-19 mortality and therefore clinically vulnerable.

Those clinically vulnerable to COVID-19 are defined by the following JCVI priority groups:

- a) children of any age with severe neuro-disability, severe or profound and multiple learning disabilities (including Down's syndrome and those on the learning disability register) or immunosuppression (as defined in table 4),
- b) adults who have underlying health conditions leading to greater risk of disease or mortality as defined in table 3,
- c) those of advanced age.

Phase 2 recommendations for primary immunisation**Adults aged 16 to 50 years not in high risk groups**

The objectives of the second phase of the COVID-19 immunisation programme were to protect those who are at risk from serious illness or death, and to protect the NHS by reducing the risks of hospitalisation and critical care admission. Phase 2 of the programme was accompanied by continued efforts to maximise coverage amongst those prioritised in Phase 1 but who remained unvaccinated, and to complete delivery of second doses to all those given first doses in Phase 1.

There is good evidence that the risks of hospitalisation and critical care admission from COVID-19 increases with age. JCVI therefore advised that the offer of vaccination during Phase 2 was offered in the following order:

- All those aged 40-49 years
- All those aged 30-39 years
- All those aged 18-29 years
- All those aged 16-17 years

Children and young people aged under 16 years at higher risk

In 2021 and 2022, primary vaccination was extended to children and young people aged 5 to 15 years at higher risk from the consequences of COVID-19, including:

- those aged 5 to 15 years in recognised clinical groups at higher risk of severe COVID-19 (see table 4)
- those aged 5 to 15 years (later restricted to those aged 12 to 15 years) who expect to share living accommodation on most days (and therefore those for whom continuing close contact is unavoidable) with individuals of any age who are immunosuppressed (defined as immunosuppressed in tables 3 or 4)

Initial JCVI advice on the paediatric clinical groups at higher risk of severe COVID was based on clinical reviews and analysis of primary care data (Williamson *et al*, 2021). Further analysis under an expert group commissioned by the Deputy Chief Medical Officer then identified a wider number of diagnostic groups with a high absolute risk (greater than 100/million) of paediatric intensive care admission over the 2020-21 period (Harwood *et al*, 2021, Smith *et al*, 2021, Ward *et al*, 2021). In addition to these distinct diagnoses, the analysis suggested that the admission rate was high in a pooled group of children with chronic conditions, based on those codes used for a Royal College of Paediatrics and Child Health review of mortality in 2013.¹ JCVI therefore decided that a set of underlying health conditions - similar to those prioritised for adult vaccination with the exception of obesity and mental illness - could reasonably also apply to children and young people (summarised in Table 4). The rate of admission for children with asthma was only slightly raised above the rate in healthy children, suggesting that, in line with the evidence from adults, only poorly controlled asthma constituted a clinical risk for the complications of COVID-19 infection.

In December 2021, following the recognition of pregnancy as a risk factor for severe COVID-19 infection and poor pregnancy outcomes during the Delta wave, pregnancy was added to the the clinical risk groups for adults and young people aged under 16 years (table 4).

In early 2023, JCVI recommended that primary vaccination could be extended to children aged 6 months to 4 years in recognised clinical risk groups (table 4). The programme will be implemented during spring and summer of 2023 in all UK countries.

Children and young people aged under 16 years and not in high risk groups

Over 2021 and 2022, the offer of primary vaccination was also extended to all children and young people aged 5 to 17 years. Because of the lower risk of the complications from COVID-19 infection in children and young people, at each stage the JCVI carefully considered the emerging evidence around the risks and benefits of the vaccination to younger people. For those aged 12 to 15 years, the committee took a precautionary

1 <https://www.rcpch.ac.uk/sites/default/files/CHR-UK - Retrospective Epidemiological Review of All-cause Mortality in CYP.pdf>

approach to mitigate against the rare risk of myocarditis, advising that the second primary dose should be given at an interval of 12 weeks. The longer interval in this age group reflected the evidence of high short term protection against severe disease from the first dose and early evidence from countries with a longer schedule (eight to twelve weeks) suggesting a lower rate of myocarditis after the second dose (Buchan *et al*, 2022). The committee also recommended that parents and young people should be fully informed about the benefits and risks of the vaccination.

Following the emergence of Omicron in late 2021, a one-off programme was also offered to those aged 5 to 11 years. As Omicron infection is particularly mild, vaccine induced protection against mild Omicron infection is short lived, and almost all children in this age group have been infected with COVID-19, JCVI also recommended that delivery of paediatric non-COVID-19 immunisation programmes should be a higher priority. Coverage in these other programmes fell behind during the pandemic, and this may have increased health inequalities.

Table 3: Clinical risk groups for individuals aged 16 years and over.

Clinical risk groups	
Chronic respiratory disease	Individuals with a severe lung condition, including those with poorly controlled asthma ¹ and chronic obstructive pulmonary disease (COPD) including chronic bronchitis and emphysema; bronchiectasis, cystic fibrosis, interstitial lung fibrosis, pneumoconiosis and bronchopulmonary dysplasia (BPD).
Chronic heart disease and vascular disease	Congenital heart disease, hypertension with cardiac complications, chronic heart failure, individuals requiring regular medication and/or follow-up for ischaemic heart disease. This includes individuals with atrial fibrillation, peripheral vascular disease or a history of venous thromboembolism.
Chronic kidney disease	Chronic kidney disease at stage 3, 4 or 5, chronic kidney failure, nephrotic syndrome, kidney transplantation.
Chronic liver disease	Cirrhosis, biliary atresia, chronic hepatitis.
Chronic neurological disease	Stroke, transient ischaemic attack (TIA). Conditions in which respiratory function may be compromised due to neurological or neuromuscular disease (e.g. polio syndrome sufferers). This group also includes individuals with cerebral palsy, severe or profound and multiple learning disabilities (PMLD) including all those on the learning disability register, Down's syndrome, multiple sclerosis, epilepsy, dementia, Parkinson's disease, motor neurone disease and related or similar conditions; or hereditary and degenerative disease of the nervous system or muscles; or severe neurological disability.
Diabetes mellitus and other endocrine disorders	Any diabetes, including diet-controlled diabetes, current gestational diabetes, and Addison's disease.
Immunosuppression	<p>Immunosuppression due to disease or treatment, including patients undergoing chemotherapy leading to immunosuppression, patients undergoing radical radiotherapy, solid organ transplant recipients, bone marrow or stem cell transplant recipients, HIV infection at all stages, multiple myeloma or genetic disorders affecting the immune system (e.g. IRAK-4, NEMO, complement disorder, SCID).</p> <p>Individuals who are receiving immunosuppressive or immunomodulating biological therapy including, but not limited to, anti-TNF, alemtuzumab, ofatumumab, rituximab, patients receiving protein kinase inhibitors or PARP inhibitors, and individuals treated with steroid sparing agents such as cyclophosphamide and mycophenolate mofetil.</p> <p>Individuals treated with or likely to be treated with systemic steroids for more than a month at a dose equivalent to prednisolone at 20mg or more per day for adults.</p> <p>Anyone with a history of haematological malignancy, including leukaemia, lymphoma, and myeloma.</p> <p>Those who require long term immunosuppressive treatment for conditions including, but not limited to, systemic lupus erythematosus, rheumatoid arthritis, inflammatory bowel disease, scleroderma and psoriasis.</p>

1 Poorly controlled asthma is defined as:

- ≥2 courses of oral corticosteroids in the preceding 24 months OR
- on maintenance oral corticosteroids OR
- ≥1 hospital admission for asthma in the preceding 24 months

<https://www.brit-thoracic.org.uk/covid-19/covid-19-information-for-the-respiratory-community/#jcvi-advice-on-covid-19-booster-vaccination-for-adults-in-clinical-at-risk-groups-and-adults-with-asthma>

	Some immunosuppressed patients may have a suboptimal immunological response to the vaccine (see Immunosuppression and HIV).
Asplenia or dysfunction of the spleen	This also includes conditions that may lead to splenic dysfunction, such as homozygous sickle cell disease, thalassemia major and coeliac syndrome.
Morbid obesity	Adults with a Body Mass Index (BMI) ≥ 40 kg/m ² .
Severe mental illness	Individuals with schizophrenia or bipolar disorder, or any mental illness that causes severe functional impairment.
Younger adults in long-stay nursing and residential care settings	Many younger adults in residential care settings will be eligible for vaccination because they fall into one of the clinical risk groups above (for example learning disabilities). Given the likely high risk of exposure in these settings, where a high proportion of the population would be considered eligible, vaccination of the whole resident population is recommended. Younger residents in care homes for the elderly will be at high risk of exposure, and although they may be at lower risk of mortality than older residents should not be excluded from vaccination programmes (see priority 1 above).
Pregnancy	All stages (first, second and third trimesters)

1 Those clinically vulnerable to COVID-19 are defined by the JCVI priority groups: a) children of any age with severe neuro-disability, severe or profound and multiple learning disabilities (including Down's syndrome and those on the learning disability register) or immunosuppression (as defined in table 4), b) adults who have underlying health conditions leading to greater risk of disease or mortality as defined in table 3, c) those of advanced age.

Table 4: Clinical risk groups for individuals aged under 16 years

Chronic respiratory disease	Including those with poorly controlled asthma ¹ that requires continuous or repeated use of systemic steroids or with previous exacerbations requiring hospital admission, cystic fibrosis, ciliary dyskinesias and bronchopulmonary dysplasia
Chronic heart conditions	Haemodynamically significant congenital and acquired heart disease, or less severe heart disease with other co-morbidity. This includes: <ul style="list-style-type: none"> • single ventricle patients or those palliated with a Fontan (Total Cavopulmonary Connection) circulation • those with chronic cyanosis (oxygen saturations <85% persistently) • patients with cardiomyopathy requiring medication • patients with congenital heart disease on medication to improve heart function • patients with pulmonary hypertension (high blood pressure in the lungs) requiring medication
Chronic conditions of the kidney, liver or digestive system	Including those associated with congenital malformations of the organs, metabolic disorders and neoplasms, and conditions such as severe gastro-oesophageal reflux that may predispose to respiratory infection
Chronic neurological disease	This includes those with <ul style="list-style-type: none"> • neuro-disability and/or neuromuscular disease that may occur as a result of conditions such as cerebral palsy, autism, epilepsy and muscular dystrophy • hereditary and degenerative disease of the nervous system or muscles, other conditions associated with hypoventilation • severe or profound and multiple learning disabilities (PMLD), Down's syndrome, including all those on the learning disability register • neoplasm of the brain
Endocrine disorders	Including diabetes mellitus, Addison's and hypopituitary syndrome
Immunosuppression	Immunosuppression due to disease or treatment, including: <ul style="list-style-type: none"> • those undergoing chemotherapy or radiotherapy, solid organ transplant recipients, bone marrow or stem cell transplant recipients • genetic disorders affecting the immune system (e.g. deficiencies of IRAK-4 or NEMO, complement disorder, SCID) • those with haematological malignancy, including leukaemia and lymphoma • those receiving immunosuppressive or immunomodulating biological therapy • those treated with or likely to be treated with high or moderate dose corticosteroids • those receiving any dose of non-biological oral immune modulating drugs e.g. methotrexate, azathioprine, 6-mercaptopurine or mycophenolate • those with auto-immune diseases who may require long term immunosuppressive treatments <p>Children who are about to receive planned immunosuppressive therapy should be considered for vaccination prior to commencing therapy.</p>
Asplenia or dysfunction of the spleen	Including hereditary spherocytosis, homozygous sickle cell disease and thalassemia major
Serious genetic abnormalities that affect a number of systems	Including mitochondrial disease and chromosomal abnormalities
Pregnancy	All stages (first, second and third trimesters)

1 Poorly controlled asthma is defined as:

- ≥2 courses of oral corticosteroids in the preceding 24 months OR
- on maintenance oral corticosteroids OR
- ≥1 hospital admission for asthma in the preceding 24 months

<https://www.brit-thoracic.org.uk/covid-19/covid-19-information-for-the-respiratory-community/#jcvi-advice-on-covid-19-vaccination-for-children-aged-12-15-years-in-clinical-at-risk-groups>

Reinforcing immunisation advice for 2021 and 2022

Initial recommendations for reinforcing immunisation

In September 2021, JCVI advised that the following groups should be offered a COVID-19 booster vaccine.

This included:

- those living in residential care homes for older adults
- all adults aged 50 years or over
- frontline health and social care workers
- all those aged 16 to 49 years with underlying health conditions that put them at higher risk of severe COVID-19 (table 3)
- all carers aged 16 years and above
- all those aged 16 years and above who are household contacts of immunosuppressed individuals (defined as immunosuppressed in tables 3 or 4) of any age

The first groups to receive boosters were therefore those prioritised in phase 1 of the COVID-19 programme (table 2, groups 1-9), with the booster offered six months from the completion of the primary course. JCVI also advised that operational flexibility should permit boosting of all adults, regardless of age, in certain closed settings or in populations such as those experiencing homelessness.

Additional surge recommendations following the emergence of the Omicron variant

Following the emergence of the Omicron variant in November 2021, JCVI advised accelerating the booster deployment in order of age and risk status. Individuals aged over 50 years and in risk groups were offered booster vaccination first, followed by those aged 18-49 years who were not in high risk groups, in descending age order. Reinforcing doses should not be given within three months of completion of the primary course.

In December 2021, JCVI advised that those aged 16-17 years, children and young people aged 12-15 who are at higher risk from COVID-19 (table 4) or those aged 12-15 years who are household contacts of immunosuppressed individuals of any age (defined as immunosuppressed in tables 3 or 4) should also be offered a booster. These children were also eligible for boosting in the autumn 2022 campaign (see below).

Additional considerations around reinforcing doses

In December 2021, following the recognition of pregnancy as a risk factor for severe COVID-19 outcomes and poor pregnancy in the Delta wave, pregnancy was added to the the clinical risk groups.

JCVI advised that those aged 5 years and over with severe immunosuppression (Boxes 1 and 2) who had not yet received their third dose should be given a third dose during the booster campaign to avoid further delay. Subsequent boosters were scheduled for at least three months after that dose, in line with the clinical advice on optimal timing in immunosuppressed individuals. See the later section on additional dose for those with severe immunosuppression.

The spring booster campaign 2022

In February 2022, recognising the small decline in observed vaccine effectiveness against hospitalisation for COVID-19 after the booster dose, JCVI recommended a spring booster

campaign for individuals at higher risk of severe COVID-19. Many of the oldest adults had received their booster vaccine dose in September or October 2021, and protection against severe disease was expected to continue to wane gradually by autumn 2022. As a precautionary strategy, an extra spring dose was advised, to sustain protection until the booster programme in autumn 2022.

The committee recommended that a booster dose should be given around 6 months after the last vaccine dose to:

- adults aged 75 years and over
- residents in a care home for older adults, and
- individuals aged 12 years and over who are immunosuppressed (defined as immunosuppressed in tables 3 or 4).

The vast majority of people aged over 75 reached an interval of around six months from their previous dose between March and June 2022. Operational flexibility was permitted, however, for individuals in care homes and for housebound patients, providing there was at least three months from the previous dose. Immunosuppressed individuals who had received an additional primary dose more recently, were also offered the booster during the spring campaign providing there was at least three months from the previous dose.

Someone in an eligible group who had received a full course of primary vaccination (two or three doses) but had not received their first booster by March 2022, was eligible for the spring booster in the campaign provided there was at least three months from the previous dose. An additional dose was not then recommended before the autumn.

The autumn booster campaign 2022

Following on from the spring campaign, the JCVI recommended a move to regular, planned and targeted boosting as the most important strategy to control COVID-19. For the 2022 autumn booster programme, the primary objective was to augment immunity in those at higher risk from COVID-19 and thereby optimise protection against severe COVID-19, specifically hospitalisation and death, over winter 2022/23.

The following groups were offered a COVID-19 booster vaccine in the autumn of 2022:

- residents in a care home for older adults and staff working in care homes for older adults
- frontline health and social care workers
- all adults aged 50 years and over
- persons aged 5 to 49 years in a clinical risk group, as set out in Tables 3 and 4
- persons aged 5 to 49 years who are household contacts of people with immunosuppression (as defined in Tables 3 and 4)
- persons aged 16 to 49 years who are carers (as defined in Table 3)

The booster was offered from September, allowing a minimum of three months from the previous dose. The programme prioritised delivery to those aged over 75 years and in care homes for older adults but recognised the need for operational flexibility based on the likely delivery models. The aim was to complete the campaign before December to provide additional protection in time for the expected winter peak of other seasonal viruses, but with mop-up opportunities during January.

Someone in the eligible groups above who had received a full course of primary vaccination (two or three doses) but had not received a booster before September 2022, were eligible for the autumn booster in the campaign provided there was at least three months from the previous dose. Additional doses are not then required. Children in high risk groups who turned five years of age after August 2022 became eligible for primary vaccination and may have been offered a booster during the autumn programme, provided there was at least three months since their second (or third) primary dose.

Living with COVID-19 vaccination programme

The UK COVID-19 pandemic vaccine programme was initiated in December 2020 with the primary objective to prevent severe disease, hospitalisations, and deaths. Now that the vast majority of the UK adult population have been vaccinated and seroprevalence studies indicate that most of the adult and childhood population have been naturally infected, the UK COVID-19 vaccination programme is expected to transition during 2023 towards a longer-term more sustainable programme.

Evidence is becoming clear that all the current vaccines provide only modest and short-term protection against infection and therefore against transmission. Protection against mild symptomatic disease is moderate but also only sustained over the short-term. With the newly emerged variants lower levels of protection against mild disease have been seen, declining to negligible levels within four to six months of primary vaccination and three to four months after booster doses. Protection against more severe forms of disease and death appears to be higher and maintained over the medium term.

Aim of the longer term COVID-19 vaccination programme

With current vaccines, the programme cannot be effectively used to interrupt transmission or to markedly impact on short term illness. The aim of the programme will therefore be to reduce severe disease (hospitalisation and mortality) and thus also to protect NHS capacity.

The risk of hospitalisation for COVID-19 continues to be disproportionately greater in those from older age groups, residents in care homes for older adults, and persons with certain underlying health conditions. Due to the high transmissibility of the Omicron variant, together with infection that can be asymptomatic or only mildly symptomatic, many persons who require hospital care for non-COVID-19 reasons may be coincidentally infected with SARS-CoV-2. Such hospitalisations are not sustainably prevented through COVID-19 vaccination, so future programmes need to be proportionate in focus.

Limitation of the pandemic COVID-19 vaccination offer

JCVI advises that with the close of the autumn 2022 vaccination campaign, the offer of a pandemic booster dose (in place since 2021) for persons aged 16 to 49 years who are not in a clinical or other high risk group should close. From the end of the spring 2023 campaign, primary course COVID-19 vaccination will become a targeted offer only to those at higher risk of severe COVID-19. After that, the offer is expected to be limited to older adults and those in a clinical risk group (tables 3 and 4). The primary offer will only be available to eligible individuals during the planned seasonal booster campaigns.

Otherwise healthy persons who develop a new health condition that places them in a clinical risk group would normally become eligible for primary vaccination when they would also become eligible for booster vaccination during a subsequent seasonal campaign or any surge response.

Individuals who develop severe immunosuppression (boxes 1 and 2) may be at high risk of severe COVID-19 and less able to sustain any protection from previous vaccination or exposure. Such individuals should be considered for catch-up primary vaccination or additional dose(s) of vaccination, before the next seasonal campaign, based on clinical judgement. The advice around such doses is outlined in the next section on severe immunosuppression, including advice about the optimal timing.

JCVI advice for 2023

Spring 2023

The committee recommended that a booster dose should be given to:

- adults aged 75 years and over
- residents in a care home for older adults, and
- individuals aged 5 years and over who are immunosuppressed (defined as immunosuppressed in tables 3 or 4).

The vast majority of people aged over 75 years should be reaching an interval of around six months from their last dose between late March and June 2023. Operational flexibility is permitted to offer the booster to eligible individuals expected to reach the target age during the spring campaign. Boosters should be offered around six months from the previous dose, but can be given three months from the previous dose; this may be particularly important to facilitate delivery of the programme to residents in care homes and the housebound. For individuals who may have received a second or third primary dose more recently, a booster can be offered during the spring campaign provided there is at least three months from the previous dose - additional doses are not then generally required until they become eligible during the next seasonal campaign.

As primary vaccination of children aged 6 months to 4 years at high clinical risk was only advised in early 2023, severely immunosuppressed children under five years of age will not be eligible for a booster in spring 2023. These children may be considered for additional doses at a later time point (see section on additional doses for individuals with severe immunosuppression).

Autumn 2023

The autumn 2023 programme is expected to target only those at higher risk of severe disease; and therefore be similar in scope to the autumn 2022 programme.

Surge response

Current Omicron variants have lower disease severity compared to infection due to previous SARS-CoV-2 variants. The emergence of a novel, more virulent, variant of concern, may require an emergency surge vaccine response. As such a variant is only likely to emerge because it escapes existing population immunity, the value of a booster programme with current vaccines may be limited. A targeted vaccine response offering readily available vaccine to those at higher risk may be required to boost background immunity, whilst waiting for availability of a more closely matched vaccine.

Additional doses for individuals with severe immunosuppression aged 6 months and above (see also later section on specific population groups)

Some individuals who are immunosuppressed due to underlying health conditions or medical treatment may not mount a full immune response to primary COVID-19 vaccination. Preliminary overall results from UK studies of real-world vaccine effectiveness (VE) in persons who are immunosuppressed suggest only a modest reduction in VE against symptomatic COVID-19 (Whitaker *et al*, 2022). Immunogenicity studies measuring binding or neutralising antibody and/or cellular responses have suggested that, amongst the immunosuppressed group, some individuals with more severe forms of immunosuppression make low or no detectable responses. A few published studies describing the effect of a third dose of mRNA vaccine in persons who are immunosuppressed report increased immune responses in varying proportions of persons. (Hall *et al*, 2021b, Kamar *et al*, 2021, Werbel *et al*, 2021).

Although there are no known correlates of protection against infection, symptomatic disease and severe COVID-19 (hospitalisation and deaths), JCVI considers that a small group of more severely immunosuppressed individuals should be offered additional doses of vaccination.

Primary vaccination

JCVI has advised that a third vaccine dose be offered to individuals aged 6 months and above who had severe immunosuppression in proximity to their first or second COVID-19 doses in the primary schedule (Boxes 1 and 2). Most individuals whose immunosuppression commenced at least two weeks after the second dose of vaccination do not require an additional primary vaccination at this stage. Individuals who had received brief immunosuppression (≤ 40 mg prednisolone per day or equivalent for children) for an acute episode (for example, asthma / COPD / COVID-19) and individuals on replacement corticosteroids for adrenal insufficiency are not considered severely immunosuppressed sufficient to have prevented response to the primary vaccination.

The third dose should be given at least eight weeks from their second primary dose.

From the end of the spring 2023 campaign, the primary course of COVID-19 vaccine becomes a targeted offer to those at higher risk and only during seasonal campaigns. The main exception to this would be unvaccinated individuals aged five years and above who become or have recently become severely immunosuppressed. These individuals should be considered for primary vaccination, regardless of the time of year. Clinical judgement should be used to decide on the best timing to commence vaccination (see section on timing).

Booster doses

Vaccinated individuals aged 6 months and above in this group will also require booster doses, in line with the rest of the eligible population, to extend protection from their primary course. Following the recognition of the Omicron variant in 2021, JCVI advised that the first reinforcing dose should be offered from three months after the third primary dose. Individuals aged over 12 years with severe immunosuppression were also eligible for the spring 2022 booster dose (see section on the spring booster campaign 2022) and for the autumn 2022 booster.

Severely immunosuppressed individuals aged 5-11 years became eligible for primary vaccination in early 2022, so were not eligible for boosting in the spring 2022 campaign, but were eligible for boosting in the autumn 2022 campaign (see below).

Severely immunosuppressed individuals aged 6 months to 4 years will be eligible for primary vaccination in 2023 and are likely to be eligible for boosters in future campaigns.

From the end of the autumn 2022 campaign, vaccinated individuals aged 6 months and above who become or have recently become severely immunosuppressed (i.e. those commencing immunosuppressive therapy or those who have developed an immunosuppressive condition) should be considered for an additional dose of COVID-19 vaccine, regardless of the time of year. This would represent either a third primary dose or an additional booster. Clinical judgement should be used to decide which individuals should be given an additional booster dose soon after their diagnosis rather than waiting for the next campaign and thus getting extra protection during the season and at the same time as other high risk groups. The optimal timing should also take account of the degree of immune suppression (see section on timing) and can be offered from three months after the second or subsequent dose.

For those that receive this additional dose, a further booster dose would normally be recommended during the next seasonal campaign, provided there is at least three months from the previous dose.

In contrast to other eligible risk groups, those who are eligible for a booster due to severe immunosuppression but miss vaccination during the campaign period, may be considered for a booster at a later date based on individual clinical judgement, balancing their immediate level of risk against the advantages of waiting till the next seasonal campaign.

Timing of additional doses

In general, vaccines administered during periods of minimum immunosuppression are more likely to generate better immune responses. Therefore, any additional doses should ideally be given with special attention paid to current or planned immunosuppressive therapies.

For example:

- the small number of patients who are about to receive planned immunosuppressive therapy should be considered for primary or booster vaccination prior to commencing therapy (ideally at least two weeks before), when their immune system is better able to make a response
- where possible, third primary or additional booster doses should be delayed until two weeks after the period of immunosuppression, in addition to the time period for clearance of the therapeutic agent
- alternatively, consideration should be given to vaccination during a treatment 'holiday' or when the degree of immunosuppression is at a minimum

Any decision to defer immunosuppressive therapy or to delay possible benefit from vaccination until after immunosuppressive therapy should only be taken after due consideration of the risks of exacerbating their underlying condition, as well as the risks from COVID-19.

Specific advice for patients on chemotherapy is available at <https://www.ukchemotherapyboard.org/publications>.

Box 1: Criteria for a third primary dose of COVID-19 vaccine in those aged 12 years and above**Individuals with primary or acquired immunodeficiency states at the time of vaccination due to conditions including:**

- acute and chronic leukaemias, and clinically aggressive lymphomas (including Hodgkin's lymphoma) who were under treatment or within 12 months of achieving cure at the time of vaccination
- individuals under follow up for a chronic lymphoproliferative disorders including haematological malignancies such as indolent lymphoma, chronic lymphoid leukaemia, myeloma, Waldenstrom's macroglobulinemia and other plasma cell dyscrasias (Note: this list is not exhaustive)
- adults and children aged 12 years and over with immunosuppression due to HIV/AIDS with a current CD4 count of <200 cells/ μ l
- Primary or acquired cellular and combined immune deficiencies – those with lymphopaenia ($<1,000$ lymphocytes/ μ l) or with a functional lymphocyte disorder
- those who had received a stem cell transplant or chimaeric antigen receptor (CAR)-T cell therapy in the 24 months before vaccination
- those who had received a stem cell transplant more than 24 months before vaccination but had ongoing immunosuppression or graft versus host disease (GVHD)
- persistent agammaglobulinaemia (IgG < 3 g/L) due to primary immunodeficiency (e.g. common variable immunodeficiency) or secondary to disease / therapy

Individuals on immunosuppressive or immunomodulating therapy at the time of vaccination including:

- those who were receiving immunosuppressive therapy for a solid organ transplant at the time of vaccination
- those who were receiving or had received in the previous 3 months targeted therapy for autoimmune disease, such as JAK inhibitors or biologic immune modulators including B-cell targeted therapies (including rituximab but in this case the recipient would be considered immunosuppressed for a 6 month period), T-cell co-stimulation modulators, monoclonal tumour necrosis factor inhibitors (TNFi), soluble TNF receptors, interleukin (IL)-6 receptor inhibitors., IL-17 inhibitors, IL 12/23 inhibitors, IL 23 inhibitors. (Note: this list is not exhaustive)
- those who were receiving or had received immunosuppressive chemotherapy or radiotherapy for any indication in the 6 months before vaccination

Individuals with chronic immune-mediated inflammatory disease who were receiving or had received immunosuppressive therapy prior to vaccination including:

- high dose corticosteroids (equivalent to ≥ 20 mg prednisolone per day) for more than 10 days in the month before vaccination
- long term moderate dose corticosteroids (equivalent to ≥ 10 mg prednisolone per day for more than 4 weeks) in the 3 months before vaccination
- non-biological oral immune modulating drugs, such as methotrexate >20 mg per week (oral and subcutaneous), azathioprine >3.0 mg/kg/day; 6-mercaptopurine >1.5 mg/kg/day, mycophenolate >1 g/day) in the 3 months before vaccination
- certain combination therapies at individual doses lower than above, including those on ≥ 7.5 mg prednisolone per day in combination with other immunosuppressants (other than hydroxychloroquine or sulfasalazine) and those receiving methotrexate (any dose) with leflunomide in the 3 months before vaccination

Individuals who had received high dose steroids (equivalent to >40 mg prednisolone per day for more than a week) for any reason in the month before vaccination

Box 2: Criteria for a third primary dose of COVID-19 vaccine in children aged 6 months to 11 years

Individuals with primary or acquired immunodeficiency states at the time of vaccination due to conditions including:

- acute and chronic leukaemias, and clinically aggressive lymphomas (including Hodgkin's lymphoma) who were under treatment or within 12 months of achieving cure at the time of vaccination
- individuals under follow up for a chronic lymphoproliferative disorders including haematological malignancies
- children with immunosuppression due to HIV/AIDS (children with a current CD4 count of <500 cells/ μ l in those aged 5 years and <200 cells/ μ l in those aged 6-11 years)
- Primary or acquired cellular and combined immune deficiencies – those with lymphopaenia (<1,000 lymphocytes/ μ) or with a functional lymphocyte disorder
- those who had received a stem cell transplant or chimaeric antigen receptor (CAR)-T cell therapy in the 24 months before vaccination
- those who had received a stem cell transplant more than 24 months before vaccination but had ongoing immunosuppression or graft versus host disease (GVHD)
- persistent agammaglobulinaemia (IgG < 3g/L) due to primary immunodeficiency (e.g. common variable immunodeficiency) or secondary to disease / therapy

Individuals on immunosuppressive or immunomodulating therapy at the time of vaccination including:

- those who were receiving immunosuppressive therapy for a solid organ transplant at the time of vaccination
- those who were receiving or had received in the previous 3 months targeted therapy for autoimmune disease, such as JAK inhibitors or biologic immune modulators including B-cell targeted therapies (including rituximab but in this case the recipient would be considered immunosuppressed for a 6 month period), T-cell co-stimulation modulators, monoclonal tumour necrosis factor inhibitors (TNFi), soluble TNF receptors, interleukin (IL)-6 receptor inhibitors., IL-17 inhibitors, IL 12/23 inhibitors, IL 23 inhibitors. (Note: this list is not exhaustive)
- those who were receiving or had received immunosuppressive chemotherapy or radiotherapy for any indication in the 6 months before vaccination

Individuals with chronic immune-mediated inflammatory disease who were receiving or had received immunosuppressive therapy prior to vaccination including:

- high dose corticosteroids (equivalent to \geq 1mg prednisolone per kg per day) for more than 10 days in the month before vaccination
- long term moderate dose corticosteroids (equivalent to \geq 0.5 mg prednisolone per kg per day for more than 4 weeks) in the 3 months before vaccination
- any dose of non-biological oral immune modulating drugs (with the exception of hydroxychloroquine and sulfasalazine), such as methotrexate, azathioprine, 6-mercaptopurine or mycophenolate in the 3 months before vaccination. (Note: this list is not exhaustive)

Individuals who had received high dose steroids (equivalent to >2mg prednisolone per kg per day for more than a week) for any reason in the month before vaccination

Age specific recommendations on vaccine type

Primary vaccination

The age appropriate advice for primary vaccination is outlined below (see the section on dosing and schedule).

Children aged 6 months to 4 years

For children aged 6 months to 4 years, an infant dose of Pfizer BioNTech (Comirnaty® 3 micrograms) is recommended.

Children below six years of age, including those who commenced immunisation with the 3 microgram infant dose before turning five, may commence and complete primary vaccination with the 3 microgram infant dose of Pfizer BioNTech if that is the only vaccine readily available in the clinic.

Children aged 5-11 years (including some who have turned 12 years)

Children in this age group should receive the paediatric formulation of Pfizer BioNTech (Comirnaty®10). Based on the JCVI advice, Comirnaty® Original/Omicron BA.4/5 (5/5 micrograms), the bivalent vaccine targeting the latest variant may be used when this vaccine is deployed. The paediatric formulation should be used in this age group, although 10 micrograms (0.1ml) of the diluted adult/adolescent vaccine may be used in exceptional circumstances only.

Children aged 5-11 years who have commenced immunisation with the paediatric dose of Pfizer BioNTech and then turn 12 years of age should also complete vaccination with the paediatric dose although the adult/adolescent dose is an acceptable alternative if this is the only supply available. Children aged 5-11 years who were given a fractional dose of the adult preparation may complete with the paediatric formulation.

Children and young people aged 12-17 years

The adult/adolescent (30 micrograms) dose of the Pfizer BioNTech vaccine (Comirnaty®) has approval for use from 12 years old and currently has the most extensive safety data in those aged 12-17 years. Pfizer BioNTech vaccines are therefore the preferred mRNA products for children. Based on current supply, the full booster dose of Pfizer BioNTech bivalent vaccine (0.3ml or 15 + 15 micrograms of Comirnaty® Original/Omicron) should be offered when someone attends for primary vaccination. Based on the JCVI advice, the preference is to use the bivalent mRNA vaccines containing the latest variant (currently BA.4/5), however using a bivalent with a previous variant (such as BA.1) or full dose mRNA vaccine (Pfizer BioNTech 30 micrograms) may be used if there would otherwise be a delay in vaccination. Although Moderna products are approved in this age group, Pfizer is currently preferred due to a lower reported rate of myocarditis.¹ Children who turned age 12 years and commenced vaccination with the 10 microgram dose may complete with the 10 microgram dose (see above).

AstraZeneca vaccine is no longer being supplied for routine use in the UK. When mRNA vaccines are not considered clinically suitable, Novavax COVID-19 vaccine may be used for primary vaccination of those aged 12-17 years.

1 <https://www.gov.uk/government/publications/coronavirus-covid-19-vaccine-adverse-reactions/coronavirus-vaccine-summary-of-yellow-card-reporting>

Adults aged 18 years and over

Evidence suggests that the risk of serious COVID-19 disease is strongly related to age, and the risk of COVID-19 mortality, hospitalisation and ICU admission is lower in younger adults. Based on current supply, the full booster dose of the bivalent Moderna COVID-19 vaccine (0.5ml or 25 + 25 micrograms of Spikevax® bivalent Original/Omicron) or the Pfizer BioNTech vaccine (0.3ml or 15 + 15 micrograms of Comirnaty® Original/Omicron) should be offered when someone attends for primary vaccination. Based on the JCVI advice, the preference is to use the bivalent mRNA vaccines containing the latest variant (currently BA.4/5), however a bivalent with a previous variant, a full dose mRNA vaccine, or, in those aged 65 years and over, Sanofi Pasteur COVID-19 vaccine (VidPrevtyn Beta®) may be used if there would otherwise be a delay in vaccination.

AstraZeneca vaccine is no longer being supplied for routine use in the UK. When mRNA vaccines are not considered clinically suitable, Novavax vaccine may be used for primary vaccination of adults, with Sanofi Pasteur as an alternate in those over 65 years.

Pregnant women in eligible groups

Pfizer BioNTech and Moderna bivalent vaccines are the preferred vaccines for primary and reinforcing vaccination of eligible women (for those under 18 years, Pfizer BioNTech vaccine (Comirnaty®) is preferred), because of more extensive experience of their use in pregnancy. When mRNA vaccines are not considered clinically suitable, Novavax COVID-19 vaccine may be used for vaccination of pregnant women, including to complete a course or as a booster, although experience in pregnancy is relatively limited.

Third primary doses for those aged six months and over with severe immunosuppression

Those who were severely immunosuppressed in proximity to primary vaccination should now receive additional doses using the age appropriate vaccine type as recommended for reinforcing doses (see next section).

When mRNA vaccines are not considered clinically suitable, Novavax COVID-19 vaccine may be used for a third dose primary vaccination of those aged 12 years and over.

Reinforcing vaccination

For mRNA vaccines, JCVI considered evidence around the differences in neutralising antibody after the bivalent vaccines compared to the original vaccine when used as a booster. The committee considered that the improvement in neutralising antibody levels were modest and likely to translate to only small improvements in protection against the BA.1 strain, and potentially against other Omicron strains, with no clear advantage against other variants. This marginally improved protection against one variant was considered insufficient to justify any substantial delay in offering boosters to those at highest risk. On this basis the committee concluded that Moderna or Pfizer BioNTech bivalent vaccine should be offered in the autumn 2022 booster programme to adults, but only where supply is sufficient to avoid delays in a planned implementation timetable. For spring 2023, the committee expect mRNA bivalent vaccines targeting BA.4/5 strains to be approved and available. These bivalent vaccines appear to be more immunogenic against the matched strains, and although BA.4/5 strains have been less prevalent in 2023, the committee agreed that the latest mRNA bivalent product should be used where authorised, available and deployed for that age group.

Based on evidence of similar immune responses as the bivalent mRNA vaccines three months after vaccination and against a range of variants, JCVI have advised that Sanofi Pasteur COVID-19 vaccine (VidPrevtyn Beta®) may also be used as a booster for older individuals in the spring 2023 campaign. The committee considered that an adjuvanted vaccine may offer some advantage for this age group and the product offers more flexible storage conditions for use in outreach setting. Data in immunosuppressed individuals is limited and so mRNA remains the preferred vaccine format in younger immunosuppressed people.

Individuals, including children at higher risk, should be clearly advised that boosting is required to ensure timely protection and therefore to accept whichever booster vaccine they were offered.

For all reinforcing doses, including additional doses for those with severe immunosuppression, the age appropriate advice below should be followed, regardless of the vaccine received as a primary dose or for previous boosters.

Eligible adults aged 75 years or over (including residents aged over 65 years in care homes for the elderly)

- A full 0.3ml dose of bivalent (15/15 micrograms) Pfizer BioNTech vaccine (currently Comirnaty® Original/Omicron BA.4/5)
- A full 0.5ml booster dose of the bivalent (25/25 micrograms) of the bivalent Moderna COVID-19 vaccine (currently Spikevax® bivalent Original/Omicron BA.4/5)
- A full 0.5ml dose of the Sanofi Pasteur COVID-19 vaccine (VidPrevtyn Beta®)

If the bivalent mRNA vaccine containing the latest variant (currently BA.4/5) is not available, a bivalent with a previous variant (such as Comirnaty® Original/Omicron BA.1 or Spikevax® bivalent Original/Omicron) or a full dose mRNA vaccine (Pfizer BioNTech 30 micrograms or Moderna 100 micrograms) may be used if there would otherwise be a delay in vaccination. When mRNA vaccines are not considered clinically suitable, either Sanofi Pasteur or Novavax COVID-19 vaccine may be used for boosting.

Eligible adults aged 18 - 74 years (including pregnant women)

- A full 0.3ml dose of bivalent (15/15 micrograms) Pfizer BioNTech vaccine (currently Comirnaty® Original/Omicron BA.4/5)
- A full 0.5ml booster dose of the bivalent (25/25 micrograms) of the bivalent Moderna COVID-19 vaccine (currently Spikevax® bivalent Original/Omicron BA.4/5)

As part of operational flexibility, eligible individuals aged 65 to 74 years may also receive Sanofi Pasteur vaccine where it would simplify delivery in that setting. This includes vaccination in domiciliary settings. When mRNA vaccines are not considered clinically suitable, Novavax COVID-19 vaccine may be used for boosting across this age group. For those aged 65-74 years Sanofi Pasteur vaccine is also a suitable alternative.

If the bivalent mRNA vaccine containing the latest variant (currently BA.4/5) is not available, a bivalent with a previous variant (such as Comirnaty® Original/Omicron BA.1 or Spikevax® bivalent Original/Omicron) or a full dose mRNA vaccine (Pfizer BioNTech 30 micrograms or Moderna 100 micrograms) may be used if there would otherwise be a delay in vaccination.

Eligible children and young adults aged 12-17 years (including pregnant women)

- A full 0.3ml dose of bivalent (15/15 micrograms) Pfizer BioNTech vaccine (currently Comirnaty® Original/Omicron BA.4/5)

When mRNA vaccines are not considered clinically suitable, Novavax COVID-19 vaccine may be used for boosting across this age group.

If the bivalent mRNA vaccine containing the latest variant (currently BA.4/5) is not available, a bivalent with a previous variant (such as Comirnaty® Original/Omicron BA.1) or a full dose mRNA vaccine (Pfizer BioNTech 30 micrograms) may be used if there would otherwise be a delay in vaccination.

Eligible children aged 5-11 years

- A full 0.2ml dose (10 micrograms) of paediatric Pfizer-BioNTech vaccine (Comirnaty®10)
- A full 0.2ml dose (5 + 5 micrograms) of paediatric Pfizer-BioNTech vaccine Comirnaty® Original/Omicron BA.4/5 (if available and deployed at that stage)

Vaccination of those aged 5 to 11 years in whom mRNA vaccines are unsuitable requires an individual clinical judgement.

Eligible children aged 6 month-4 years

- A full 0.2ml dose (3 micrograms) of paediatric Pfizer-BioNTech vaccine (Comirnaty®3).
- Vaccination of those aged 6 months to 4 years in whom mRNA vaccines are unsuitable requires an individual clinical judgement.

Other considerations

Previous incomplete vaccination

If the course is interrupted or delayed, it should be resumed using the same vaccine but any delayed doses should not be repeated. Evidence suggests that those who receive mixed schedules, including mRNA and adenovirus vectored vaccines, make a good immune response, (Liu *et al*, 2021) although rates of side effects with a heterologous second dose are higher. (Shaw *et al*, 2021). Accumulating evidence now supports the use of heterologous schedules for primary and reinforcing immunisation. For individuals who started the schedule and who attend for vaccination where the same vaccine is not available or suitable, or if products received previously are unknown or not available, a dose of the locally available product should be given. Individuals who experienced severe expected reactions after a previous dose of AstraZeneca or Pfizer BioNTech vaccines should be informed about the higher rate of such reactions when they receive a second dose of an alternate vaccine. (Powell *et al*, 2021)

Children aged 5-12 years who have commenced immunisation with the paediatric dose of Pfizer BioNTech should complete vaccination with the paediatric dose (although an adult/adolescent dose is an alternate in those who turn 12 years of age between doses). Those who present for the second dose over the age of 12 years should be given an adult/adolescent dose of vaccine.

Individuals vaccinated overseas or as part of clinical trials

Individuals who have been vaccinated abroad are likely to have received a vaccine based on the spike protein or an inactivated whole viral vaccine and are expected to be boosted by the vaccines currently used in the UK. Specific advice on vaccination of those who received COVID-19 vaccine overseas is available from UKHSA. <https://www.gov.uk/government/publications/covid-19-vaccination-programme-guidance-for-healthcare-practitioners>.

Individuals who have participated in a clinical trial of either primary or booster COVID-19 vaccination should be provided with written advice on whether and when they should be safely vaccinated in the routine programme. Trial participants who are eligible for boosters should be offered vaccination in line with the general population, at least three months after the dose considered as the final primary dose or the final revaccination (if the latter is required for certification purposes).

Co-administration with other vaccines

Initially data on co-administration of COVID-19 with other vaccines was limited. In the absence of such data first principles would suggest that interference between inactivated vaccines with different antigenic content is likely to be limited (see Chapter 11). Based on experience with other vaccines, any potential interference is most likely to result in a slightly attenuated immune response to one of the vaccines. There is no evidence of any safety concerns, although it may make the attribution of any adverse events more difficult. Similar considerations apply to co-administration of inactivated (or non-replicating) COVID-19 vaccines with live vaccines such as MMR. In particular, live vaccines which replicate in the mucosa, such as live attenuated influenza vaccine (LAIV) are unlikely to be seriously affected by concomitant COVID-19 vaccination.

A UK study of co-administration of AstraZeneca and Pfizer BioNTech COVID-19 vaccines with inactivated influenza vaccines confirmed acceptable immunogenicity and reactogenicity (Lazarus *et al*, 2021). Although a study of co-administration of Novavax COVID-19 vaccine with inactivated influenza did show some attenuation of the antibody response to COVID-19 (Toback *et al*, 2022), co-administration was still associated with high efficacy against COVID-19 in the phase 3 study (Heath *et al*, 2022). A recent study has shown an acceptable safety profile when COVID-19 is co-administered with inactivated shingles vaccine (Naficy *et al*, 2022).

Based on the evidence above, and, as COVID-19 vaccines are considered inactivated (including the non-replicating adenovirus vaccine), where individuals in an eligible cohort present having recently received one or more inactivated or another live vaccine, COVID-19 vaccination should still be given. The same applies for other live and inactivated vaccines where COVID-19 vaccination has been received first or where a patient presents requiring two or more vaccines. It is generally better for vaccination to proceed to avoid any further delay in protection and to avoid the risk of the patient not returning for a later appointment. This includes but is not limited to vaccines commonly administered around the same time or in the same settings (including pneumococcal polysaccharide vaccine and shingles vaccine in those aged over 65 years, pertussis-containing vaccines and influenza vaccines in pregnancy, and LAIV, HPV, MenACWY and Td-IPV vaccines in school age children).

Monoclonal antibodies to SARS-CoV-2 have recently been licensed for the treatment and prophylaxis of COVID-19 infection. Primate data suggests that administration of the AstraZeneca combination monoclonal antibody product did not interfere with the subsequent response to active vaccination. Based on this limited evidence, therefore, no specific interval is required between receipt of these products and COVID-19 vaccination, or vice versa. As the use of these products is likely to be prioritised to those who are less able to respond to vaccination, for example immunosuppressed individuals, additional doses of vaccine may be required as outlined above.

Specific population groups

Pregnancy

There is no known risk associated with giving inactivated, recombinant viral or bacterial vaccines or toxoids during pregnancy or whilst breastfeeding (Kroger *et al*, 2013). Since inactivated vaccines cannot replicate, they cannot cause infection in either the mother or the fetus. Although AstraZeneca COVID-19 vaccine contains a live adenovirus vector, this virus is not replicating so will not cause infection in the mother or the fetus. As with most pharmaceutical products, large clinical trials of COVID-19 vaccine in pregnancy have not been carried out.

Developmental and reproductivity testing of the Pfizer BioNTech, Moderna and AstraZeneca vaccines in animals have not raised any concerns. Adenovirus vectors, similar to those used in the AstraZeneca COVID-19 vaccine, have been widely used to vaccinate women against Ebola without raising any concern; formal trials of these vaccines in pregnancy are due to proceed.

A large amount of observational data from women vaccinated in pregnancy in the USA have not shown an increase in adverse pregnancy outcomes.¹ JCVI has therefore advised that women who are pregnant should be recommended to receive primary immunisation, and that pregnancy is considered a clinical risk group for the autumn booster programme. Although data systems and collection periods differ between the UK countries, around 150,000 women in England, 25,000 in Scotland and 4,500 in Wales have received at least one dose of COVID-19 vaccine whilst pregnant. Initial analysis of birth outcomes in women who had received at least one dose of the vaccine and delivered between January to November in England showed a similar or higher rate of good birth outcomes than in unvaccinated women. <https://www.gov.uk/government/publications/covid-19-vaccine-weekly-surveillance-reports>.

Routine questioning about last menstrual period and/or pregnancy testing is not required before offering the vaccine. Women who are planning pregnancy or in the immediate postpartum should be vaccinated with a suitable product for their age and risk status.

If a woman finds out she is pregnant after she has started a course of vaccine, she should complete vaccination at the recommended interval.

Termination of pregnancy following inadvertent immunisation should not be recommended. Surveillance of the inadvertent administration of COVID-19 vaccines in early pregnancy is no longer being conducted for the UK by the UK Health Security Agency.² This surveillance was being undertaken to document safety in women who unknowingly receive a vaccine in early pregnancy to inform women and health professionals. As above, any women who are inadvertently vaccinated in early pregnancy should complete vaccination at the recommended interval.

Breastfeeding

There is no known risk associated with being given a non-live vaccine whilst breastfeeding. JCVI advises that breastfeeding women should be offered any suitable COVID-19 vaccine. Emerging safety data is reassuring: mRNA was not detected in the breast milk of recently vaccinated women (Golan *et al*, 2021) and protective antibodies have been detected in breast milk (Gray *et al*, 2021).

The developmental and health benefits of breastfeeding are clear and should be discussed with the woman, along with her clinical need for immunisation against COVID-19.

1 <https://www.cdc.gov/coronavirus/2019-ncov/vaccines/safety/vsafepregnancyregistry.html>

2 <https://www.gov.uk/guidance/vaccination-in-pregnancy-vip>

Children

SARS-CoV-2 vaccine trials have been conducted in those aged 12-15 years using the adult/adolescent dose of both Pfizer BioNTech and Moderna vaccines. A 10 microgram dose of Pfizer BioNTech vaccines formulated for children, has been approved by the MHRA for use in children aged 5-11 years.

Children and young people have a very low risk of COVID-19, severe disease or death due to SARS-CoV-2 compared to adults. Vaccination of school children may offer short term protection against school absences.¹

Children under 16 years of age are at low risk of COVID-19 mortality, and risk of hospital admission is largely confined to those in clinical risk groups. Some children, such as those with profound and multiple LD (PMLD) and severe LD, and including children with Down's syndrome and cerebral palsy are at higher risk of hospitalisation for COVID-19 (Williamson *et al*, 2021). Analysis undertaken by an expert group recently concluded that rates of intensive care admission exceeded 100 per million over the first year of the pandemic in children with a broad range of underlying conditions. Eligible children aged 6 months and above with these conditions are therefore recommended to receive Pfizer BioNTech COVID-19 vaccine (see section on children and tables 3 and 4).

Immunosuppression and HIV (see also earlier section on additional doses for individuals with severe immunosuppression)

Individuals who have immunosuppression and HIV infection (regardless of CD4 count) should be given COVID-19 vaccine in accordance with the recommendations and contraindications above. Although AstraZeneca COVID-19 vaccine contains a live adenovirus vector, this virus is not replicating and is considered safe in immunosuppressed people. Other adenovirus vector vaccines have been trialled in populations with high prevalence of HIV and shown no serious adverse events (Kennedy *et al*, 2017). Many of the COVID-19 vaccines were studied in people living with HIV infection who are stable on treatment.

Individuals with immunosuppression may not make a full immune response to vaccination. As there is limited evidence on the response for each specific immunosuppressing therapy or condition then general principles are needed to inform the optimal timing of delivery. It is expected that vaccines administered during periods of minimum immunosuppression are more likely to generate better immune responses. Therefore, vaccination should be given ideally, with special attention paid to current or planned immunosuppressive therapies. Recent studies have suggested immune responses were better in patients with cancer who received their chemotherapy at least two weeks before vaccination (Monin-Aldama *et al*, 2021), or people on methotrexate when their therapy was temporarily stopped for two weeks after vaccination (Abhishek *et al*, 2022). Specialists may advise their patients based on their knowledge and understanding of their immune status and likely immune response to vaccination, but should also consider the risk from COVID-19 and the patient's likelihood of exposure. **Any decision to defer immunosuppressive therapy or to delay possible benefit from vaccination until after therapy should only be taken with due consideration of the risks of exacerbating their underlying condition, as well as the risks from COVID-19.**

Emerging evidence suggests that many patients with immunosuppression do derive

1 <https://www.gov.uk/government/publications/universal-vaccination-of-children-and-young-people-aged-12-to-15-years-against-covid-19>

protection after two doses of vaccination. (Whitaker *et al*, 2022). As for immunocompetent individuals, however, protection from primary vaccination does decline over time. As those with immunosuppression remain at higher risk of serious complications from COVID-19 infection, this group of individuals have been eligible for a booster in the spring and autumn 2022 campaigns and are expected to be eligible for future seasonal campaigns. To reduce the risk of exposure, household contacts of individuals with immunosuppression were eligible for primary vaccination and, in certain ages group, for boosting in seasonal campaigns.

Some individuals with more severe immunosuppression do not make a good immune response to a complete primary course of vaccine. This includes, but is not limited to, individuals on immunosuppression for solid organ transplants (Prendecki *et al*, 2021), those with haematological cancers who are within six months of completing curative therapy (Lim *et al*, 2021), and those on certain monoclonal antibody therapies (Mahil *et al*, 2021). JCVI has therefore recommended that some individuals require additional doses (see Boxes 1 and 2), and after the autumn 2022 campaign, individuals becoming severely immunosuppressed may be considered for an additional dose between campaigns (see section on additional doses).

Post-vaccination testing for spike antibody may be considered by specialists managing individuals with severe immunosuppression. Low levels of detectable antibody may indicate poor protection against mild infection, although protection against severe disease may still be present due to T and B cell immunological memory. Taking into account their underlying immune defect, specialists may use this information to advise individuals to take additional precautions to avoid exposure or to receive information about access to early antiviral therapy.

Individuals who receive bone marrow transplants, and many individuals who receive CAR-T therapy for certain conditions, may lose immunological memory from vaccination received prior to the treatment and the development of the underlying condition. After treatment and recovery, these individuals should be considered for a full course of revaccination for **all** vaccines used in the routine programme (see chapter 7). Specialist advice should be followed on which vaccines can be safely given and on the optimal timing for commencing revaccination.

Contraindications

There are very few individuals who cannot receive the COVID-19 vaccines approved in the UK. Where there is doubt, rather than withholding vaccination, appropriate advice should be sought from the relevant specialist, or from the local immunisation or health protection team.

The following are relative contra-indications to receiving a COVID-19 vaccine:

- individuals who have had a previous anaphylaxis reaction to a COVID-19 vaccine
 - individuals who received Astra-Zeneca, Sanofi Pasteur or Novavax vaccines may be given a different product to the one implicated in any setting, with observation for 30 minutes
 - some individuals with initial apparent allergic reaction to an mRNA vaccine can tolerate a second dose of the same vaccine. Where there were no objective signs of anaphylaxis and symptoms rapidly resolved (with no more than 1 dose of IM adrenaline), a further dose of the same vaccine can be given in any vaccination setting. If the reaction might have been anaphylaxis, obtain expert advice; if a decision is made to administer the same vaccine, then this should be done under medical supervision in the hospital setting. See flowchart for further information.

- individuals with a prior allergic reaction to any component (excipient) of the COVID-19 vaccine e.g. polyethylene glycol
 - published data now show that some individuals with prior allergic reaction to PEG-containing medicines (eg. PEG-asparaginase) can tolerate the Pfizer BioNTech vaccine (although the historical reaction may have been due to a non-PEG component) (Mark *et al*, 2021). Expert advice should be obtained and if a decision is made to administer an mRNA vaccine, then this should only be done in hospital under medical supervision

Precautions

Minor illnesses without fever or systemic upset are not valid reasons to postpone immunisation. If an individual is acutely unwell, immunisation may be postponed until they have fully recovered. This is to avoid confusing the differential diagnosis of any acute illness (including COVID-19) by wrongly attributing any signs or symptoms to the adverse effects of the vaccine.

Individuals with a history of COVID-19 infection

There are no safety concerns from vaccinating individuals with a past history of COVID-19 infection, or with detectable COVID-19 antibody.

Vaccination of individuals who may be infected or asymptomatic or incubating COVID-19 infection is unlikely to have a detrimental effect on the illness, although individuals with suspected COVID-19 infection should not attend vaccination sessions to avoid infecting others. As clinical deterioration can occur up to two weeks after infection, vaccination should ideally be deferred until clinical recovery. There is no longer any need to defer immunisation in individuals after recovery from a recent episode of compatible symptoms, whether or not they are tested for COVID-19. During care home outbreaks, vaccination of residents with confirmed COVID-19 may go ahead, provided the residents are clinically stable and infection control procedures can be maintained. These populations are likely to be highly vulnerable and this policy should help to maximise vaccination coverage without the need for multiple visits.

Having prolonged COVID-19 symptoms is not a contraindication to receiving COVID-19 vaccine but if the patient is seriously debilitated, still under active investigation, or has evidence of recent deterioration, deferral of vaccination may be considered to avoid incorrect attribution of any change in the person's underlying condition to the vaccine.

Individuals with a history of allergy

A very small number of individuals have experienced anaphylaxis when receiving a COVID-19 vaccine. Anyone with a history of allergic reaction to an excipient in the COVID-19 vaccine should not receive that vaccine (except with expert advice), but those with any other allergies (such as a food allergy) – including those with prior anaphylaxis – can have the vaccine.

The Pfizer BioNTech and Moderna mRNA vaccines contain polyethylene glycol (PEG). PEGs (also known as macrogols) are a group of known allergens commonly found in medicines, many household products and cosmetics. Medicines containing PEG include some tablets, laxatives, depot steroid injections, and some bowel preparations used for colonoscopy. Known allergy to PEG is rare. Evidence now shows that PEG allergy is implicated in only a minority of allergic reactions reported after COVID-19 vaccines.

The rate of anaphylaxis reported to date after the AstraZeneca vaccine is in line with the expected rate of anaphylaxis to non-COVID vaccines. The AstraZeneca, Novavax and Sanofi

Pasteur vaccines do not contain PEG but do contain a related compound called polysorbate 80. Rarely, people with PEG allergy may also be allergic to polysorbate 80. However, polysorbate 80 is widely used in medicines and foods, and is present in many medicines including monoclonal antibody preparations. Some injected influenza vaccines (including the main vaccine used in over 65 year olds) contain polysorbate 80. Individuals who have tolerated injections that contain polysorbate 80 (including the adjuvanted influenza vaccine, Fludax[®] and the GlaxoSmithKline vaccine Fluarix[®]) are likely to tolerate the AstraZeneca and Novavax vaccines.

The Sanofi Pasteur vaccine also contains PS80 at a higher level than these influenza vaccines, as well as small amounts of polysorbate 20 (a similar compound). Despite very limited experience with this vaccine, it is unlikely that individuals with an allergy to PEG would react to the Sanofi Pasteur vaccine, particularly if they have tolerated a previous influenza vaccine and/or an AstraZeneca or Novavax vaccine. Advice on the management of patients with allergy is summarised in table 5.

Following COVID-19 vaccine administration, individuals should be observed for any immediate reactions whilst they are receiving any verbal post vaccination information and exiting the centre. Facilities for management of anaphylaxis should be available at all vaccination sites (see chapter 8). Advice has also been issued by the Resuscitation Council.¹

According to the Summaries of Product Characteristics, it is recommended that all recipients of the Pfizer BioNTech, Moderna, Novavax and Sanofi Pasteur vaccines are kept for observation and monitored for a minimum of 15 minutes. In recognition of the need to accelerate delivery of the programme in response to the emergence of the Omicron variant, the UK Chief Medical Officers recommended suspension of this requirement for the two mRNA vaccines (Pfizer BioNTech and Moderna) in both children and adults. The suspension of the observation period in individuals without a history of allergy has since been agreed by the Commission on Human Medicines.² The advice to suspend the routine 15 minute observation period therefore applies to all currently available COVID-19 vaccines in all age groups, including the bivalent mRNA products and the both the Novavax and Sanofi Pasteur vaccines.

The MHRA will continue to closely monitor anaphylaxis post-COVID-19 vaccination; reporting of adverse events via the Yellow Card Scheme is strongly encouraged (www.mhra.gov.uk/yellowcard).

Vaccinated individuals should be informed about how to access immediate healthcare advice in the event of displaying any symptoms. In some settings, for example domiciliary vaccination, this may require a responsible adult to be present for at least 15 minutes after vaccination. Patients with a personal history of allergy should be managed as in table 5. No specific management is required for patients with a family history of allergies.

As fainting can occur following vaccination, all those vaccinated with any of the COVID-19 vaccines should not drive for 15 minutes after vaccination.

1 www.resus.org.uk/about-us/news-and-events/rcuk-publishes-anaphylaxis-guidance-vaccination-settings

2 <https://www.gov.uk/government/publications/coronavirus-covid-19-vaccine-adverse-reactions/coronavirus-vaccine-summary-of-yellow-card-reporting>

Table 5: Management of patients with a history of allergy

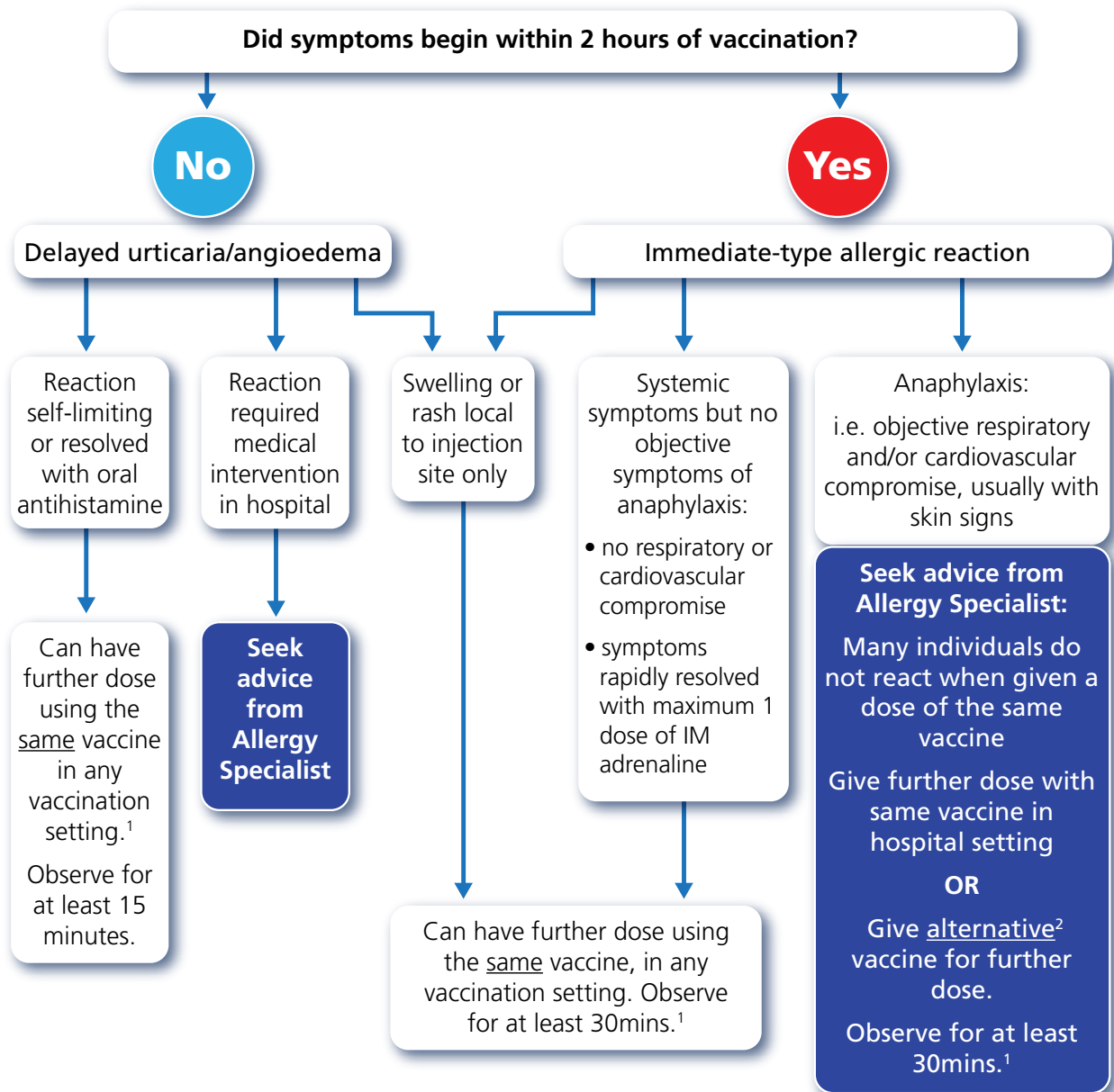
	Proceed with vaccination (no special precautions)	Special precautions	Vaccination contra-indicated
PATIENT CHARACTERISTICS	<ul style="list-style-type: none"> ● previous allergic reaction (including anaphylaxis) to a food, insect sting and most medicines (where trigger has been identified) ● previous non-systemic reaction to a vaccine ● hypersensitivity to non-steroidal anti-inflammatory drugs e.g. aspirin, ibuprofen ● mastocytosis 	<ul style="list-style-type: none"> ● prior non-anaphylaxis allergic reaction to COVID-19 vaccine ● history of immediate anaphylaxis to multiple, different drug classes, with the trigger unidentified (this may indicate PEG allergy) ● history of anaphylaxis to a vaccine, injected antibody preparation or a medicine likely to contain PEG (e.g. depot steroid injection, laxative) ● history of idiopathic anaphylaxis 	<ul style="list-style-type: none"> ● prior anaphylaxis reaction to COVID-19 vaccine ● prior systemic allergic reaction to a component of the vaccine <p>(for known PEG allergy see text above)</p>
ACTIONS	<ul style="list-style-type: none"> ● proceed with vaccination in any setting ● some individuals may be reassured by being observed for 15 minutes (may not be required if previously tolerated the same vaccine) ● some patients (e.g. those with mastocytosis) may benefit from pretreatment with anti-histamine to reduce allergic symptoms 	<ul style="list-style-type: none"> ● consider possibility of PEG allergy and seek allergy advice if needed ● a person has previously tolerated a dose of the same vaccine, it is safe to administer in any setting. <p>Otherwise</p> <ul style="list-style-type: none"> - consider giving vaccine and observe for 30 minutes 	<ul style="list-style-type: none"> ● refer to allergist or other appropriate specialist ● consider administration of the implicated mRNA vaccine under medical supervision in hospital, or, where reaction was to a non-mRNA vaccine give alternative vaccine in any setting ● consider observation for 30 minutes

Patients with undiagnosed PEG allergy often have a history of immediate onset-unexplained anaphylaxis or anaphylaxis to multiple classes of drugs. Such individuals should not be vaccinated with the Pfizer BioNTech or Moderna vaccines, except on the expert advice of an allergy specialist or where at least one dose of the same vaccine had been tolerated previously. A non-mRNA vaccine (such as Novavax or Sanofi Pasteur) can be used as an alternative (unless otherwise contraindicated), particularly if they previously tolerated the adjuvanted influenza vaccine. The vaccine should be administered in a setting with full resuscitation facilities (e.g. a hospital), and a 30 minute observation period is recommended. Advice for children with cancer who may be receiving PEG containing drugs is available at <https://www.cclg.org.uk/Coronavirus-advice>.

The British Society for Allergy and Clinical Immunology (BSACI) has advised that individuals who have a reaction to the first dose of a COVID-19 vaccine may be able to receive a second dose of vaccine, as in the flowchart below. Many individuals have tolerated subsequent doses of the same vaccine, and this is preferred as it avoids an individual being wrongly labelled as allergic for life.

Individuals with non-allergic reactions (vasovagal episodes, non-urticarial skin reaction or non-specific symptoms) to the first dose of a COVID-19 vaccine can receive the second dose of vaccine in any vaccination setting. Observation for 15 minutes is recommended.

Flowchart for managing patients who have allergic reactions to a previous dose of COVID-19 vaccine



COVID-19 - SARS-Cov-2

1 Consider pre-treatment with non-sedating antihistamine, at least 30mins prior to vaccination.

2 If reaction was to AstraZeneca, Novavax or Sanofi Pasteur vaccine, complete or boost with a different vaccine, which may include an mRNA vaccine. If reaction was to an mRNA vaccine, give any mRNA vaccine or Novavax or Sanofi Pasteur vaccine in a hospital setting.

Thrombosis and thrombocytopenia syndrome (TTS) occurring after COVID-19 vaccination

A condition involving serious thromboembolic events accompanied by thrombocytopenia, has been reported after AstraZeneca vaccination.

There is no evidence of any underlying risk factors in the individuals affected by this condition who have mainly been previously healthy. The condition is rare, tends to present with unusual forms of clotting and the mechanism is believed to be an idiosyncratic reaction related to an immune response to the AstraZeneca vaccine. Because of this likely immune

mechanism, there is no reason to believe that individuals with a past history of clots or of certain thrombophilic conditions would be at increased risk of this very rare condition. Similarly, although pregnancy increases the risk of clotting conditions, there is no evidence that pregnant women, those in the post-partum or women on the contraceptive pill are at higher risk of the specific condition of thrombosis in combination with thrombocytopenia after the AstraZeneca vaccine. There have been no confirmed cases reported in pregnant women to date. Caution should be used, however, when vaccinating individuals who have a history of a previous episode of heparin induced thrombocytopenia and thrombosis (HITT or HIT type 2).

Individuals who experience a clotting episode with concomitant thrombocytopenia following the first dose of AstraZeneca vaccine should be properly assessed. If they are considered to have the reported condition, further vaccination should be deferred until their clotting has completely stabilised. A number of patients with confirmed TTS (n=23) in the UK have now tolerated a second dose of Pfizer BioNTech, with a smaller number having tolerated Moderna (n=2) or AstraZeneca (n=1) vaccines (Lacy J *et al*, 2021). A recent study documented decline in anti-PF4 antibody by 12 weeks following the first dose of vaccine; follow up of 5 patients with confirmed TTS who received Pfizer BioNTech for the second dose at a minimum interval of 10 weeks had no further episodes (Schönborn *et al*, 2021). Current evidence would therefore support a decision to complete the primary course or boost patients with a history of TTS with an mRNA vaccine, provided at least 12 weeks has elapsed from the implicated dose.

Individuals who have received the first dose of AstraZeneca vaccine without developing this rare condition are advised to receive the second dose of the same vaccine at the currently recommended interval. To date, there is no signal of an increased risk of this condition after the second dose and the rate of other reactions is lower at the second dose than after the first dose of this vaccine.

Based on current evidence JCVI advised a preference for an alternative vaccine for healthy people under 40 years of age, including health and social care workers, unpaid carers and household contacts of immunosuppressed individuals.

Individuals with past clotting episodes and those diagnosed with thrombophilia, whether or not they are on long term anti-coagulation, remain at risk of COVID-19 disease. There is no evidence that those with a prior history of thrombosis or known risk factors for thrombosis are more at risk of developing this immune-mediated condition of thrombosis in combination with thrombocytopenia after the AstraZeneca vaccine. For most of these individuals, the risk of recurrent thrombosis due to COVID-19 infection, remains far greater than the risk of this syndrome. Therefore individuals aged 40 years and over with such a history should be vaccinated with any of the available vaccines (provided they are not otherwise contra-indicated). The same consideration applies to those who experience common clotting episodes after the first dose of AstraZeneca vaccine but without concomitant thrombocytopenia.

Other rare conditions

Myocarditis and pericarditis

Cases of myocarditis and pericarditis have been reported rarely after COVID-19 vaccination. If an individual develops myocarditis or pericarditis following the first COVID-19 vaccination they should be assessed by an appropriate clinician to determine whether it is likely to be vaccine related <https://www.gov.uk/government/publications/myocarditis-and-pericarditis->

[after-covid-19-vaccination/myocarditis-and-pericarditis-after-covid-19-vaccination-guidance-for-healthcare-professionals](#). As the mechanism of action and risk of recurrence of myocarditis and pericarditis are being investigated, the current advice is that an individual's second or subsequent doses should be deferred pending further investigation and careful consideration of the risks and benefits. For those that experience myocarditis or pericarditis within two weeks of the first dose of an mRNA vaccine, testing for N antibody may indicate prior exposure to COVID-19. These individuals are likely to be well protected and therefore the benefit from a second or subsequent dose is likely to be more limited. Where N antibody is negative or in other circumstances where a further dose is considered necessary, for example in those higher risk of the complications of COVID-19 infection, a second or booster dose of Pfizer BioNTech vaccine should be considered once the patient has fully recovered. Emerging evidence suggests that an interval of at least 12 weeks should be observed from the previous dose. Pfizer BioNTech is preferred over Moderna due to a slightly higher rate of myocarditis reported after the latter vaccine; AstraZeneca should not be offered to those who have previously received an mRNA vaccine given the more serious nature of thrombosis and thrombocytopenia syndrome. As cases of myocarditis and pericarditis have been reported after Novavax vaccine and there is less overall safety experience, this vaccine should only be used with caution in those who have had myocarditis after a previous dose of an mRNA vaccine.

Guillain-Barré syndrome

Very rare reports have been received of GBS following COVID-19 vaccination, so healthcare professionals should be alert to the signs and symptoms of GBS to ensure correct diagnosis and to rule out other causes, in order to initiate adequate supportive care and treatment.

Individuals who have a history of GBS should be vaccinated as recommended for their age and underlying risk status. Cases of GBS reported following vaccination may occur by chance (the background rate of GBS is 2 per 100,000 per year in the population) and no causal mechanism with COVID-19 vaccination has been proven. There is evidence to suggest that having had a prior diagnosis of GBS does not predispose an individual to further episodes of GBS when immunised with other vaccines (Baxter *et al*, 2012) and for the Pfizer BioNTech COVID-19 vaccine (Shapiro Ben David *et al*, 2021). In those who are diagnosed with GBS after the first dose of vaccine, the balance of risk benefit is in favour of completing a full COVID-19 vaccination schedule. On a precautionary basis, however, where GBS occurs within six weeks of an Astra Zeneca vaccine, for any future doses Pfizer or Moderna COVID-19 vaccines are preferred. Where GBS occurs following either of the mRNA vaccines, further vaccination can proceed as normal, once recovered.

Thrombocytopenia

Cases of thrombocytopenia (without accompanying thrombosis) have been reported rarely in the first four weeks after receiving AstraZeneca vaccination. Some of these cases have occurred in individuals with a history of immune thrombocytopenia (ITP). Previous ITP is not a contra-indication for vaccination but platelet monitoring is advised for patients with a history of ITP who receive AstraZeneca vaccine. Although evidence suggests a raised risk ITP after the AstraZeneca vaccine (Simpson *et al*, 2021), ITP has also been reported with other COVID-19 vaccines (Lee *et al*, 2021). Guidance produced by the UK ITP Forum Working Party therefore advises discussing the potential for a fall in platelet count in patients with a history of ITP receiving any COVID-19 vaccine and recommends a platelet count check 2-5 days after vaccination (<https://b-s-h.org.uk/about-us/news/covid-19-updates/>). Individuals who experience ITP in the four weeks after the first dose of AstraZeneca vaccine should be

assessed by a haematologist and the risk benefit of further vaccination and with which product should be considered on an individual basis. If receiving further vaccination, the platelet count should be monitored.

Capillary leak syndrome

Extremely rare reports of capillary leak syndrome have been reported after AstraZeneca and Moderna vaccines in individuals with a prior history of this condition. Individuals with a history of capillary leak syndrome, should be carefully counselled about the risks and benefits of vaccination and advice from a specialist should be sought.

Reporting anaphylaxis and other allergic reactions

Anaphylaxis is a very rare, recognised side effect of most vaccines and suspected cases should be reported via the Yellow Card Scheme (www.mhra.gov.uk/yellowcard). [Chapter 8](#) of the Green Book gives detailed guidance on distinguishing between faints, panic attacks and the signs and symptoms of anaphylaxis. If a case of suspected anaphylaxis meets the clinical features described in Chapter 8, this should be reported via the Yellow Card Scheme as a case of 'anaphylaxis'. Cases of less severe allergic reactions (i.e. not including the clinical features of anaphylaxis) should not be reported as anaphylaxis but as 'allergic reaction'.

As these vaccines are labelled with a black triangle, all adverse reactions occurring in individuals of any age after vaccination should be reported to the MHRA using the Yellow Card Scheme. Anyone can report a suspected adverse reaction to the Medical and Healthcare products Regulatory Agency (MHRA) using the Yellow Card reporting scheme (www.mhra.gov.uk/yellowcard). Any adverse reaction should also be documented in accordance with local procedures.

Management of suspected cases and contacts

There is currently limited evidence to support the use of COVID-19 vaccines as post-exposure prophylaxis or to interrupt transmission during outbreaks. The use of vaccine to provide direct protection to vulnerable individuals in prolonged community outbreaks should be discussed with the local health protection teams.

Current recommendations for testing and contact tracing and guidance on infection control is regularly updated and can be found in the following links:

<https://www.gov.uk/coronavirus>

<https://www.gov.scot/collections/coronavirus-covid-19-guidance/>

<https://www.hps.scot.nhs.uk/a-to-z-of-topics/covid-19/>

<https://phw.nhs.wales/topics/latest-information-on-novel-coronavirus-covid-19/>

<https://www.publichealth.hscni.net/covid-19-coronavirus/guidance-hsc-staff-healthcare-workers-and-care-providers>

Additional resources

Supplies

COVID-19 vaccines for those authorised by the NHS to deliver the programme will be made available for ordering on the ImmForm website <https://portal.immform.phe.gov.uk/> telephone 0207 183 8580 or through the Foundry ordering platform in England.

Arrangements in Scotland, Wales and Northern Ireland may be different, please contact Public Health Agencies in each respective administration for local details.

Key links

The full specification for those diagnoses, and associated clinical codes, eligible for COVID-19 vaccination has been developed and is available on the PRIMIS website <https://www.nottingham.ac.uk/primis/covid-19/covid-19.aspx>. Access to the link is available to NHS professionals and requires online registration.

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