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COVID-19 - SARS-CoV-2

NOTIFIABLE

The virus

COVID-19 disease first emerged as a presentation of severe respiratory infection in Wuhan, China in late 2019 (WHO, 2020). By January 2020, lower respiratory samples taken from affected patients were sequenced and demonstrated a novel coronavirus (SARS-CoV-2) (Huang *et al*, 2020). The first two cases in the UK were seen in late January (Lillie *et al*, 2020). In March 2020, the WHO declared a SARS-CoV-2 pandemic (WHO Director-General, 2020).

SARS-CoV-2 is a member of the family of Coronaviridae and genus Betacoronavirus (Zhu *et al*, 2020). Phylogenetic analysis of SARS-CoV-2 has shown that it is genetically distinct from the SARS coronavirus (Dhama, *et al*. 2020), but appears to share strong sequence similarity to bat coronaviruses in China (Lam *et al*, 2020).

As with other coronaviruses, SARS-CoV-2 is an RNA virus which 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 and consists of an S1 and S2 subunit (Kaur *et al* 2020). The S1 subunit has two functional domains: the N terminal domain (NTD) and receptor binding domain (RBD) which contains the receptor binding motif (RBM) (Kaur *et al*, 2020). The RBM binds to angiotensin converting enzyme 2 (ACE2) on host cells and is endocytosed with subsequent release of the viral genome into the cytoplasm (Amanat *et al*, 2020).

SARS-CoV-2 is primarily transmitted by person to person spread through respiratory aerosols, direct human contact and fomites (Kaur *et al*, 2020). Estimates of the basic reproduction number [R] were initially between 2 and 3 although a recent estimate was as high as 5.7 (Sanche *et al*, 2020). This high transmissibility indicates that stringent control measures, such as active surveillance, physical distancing, early quarantine and contact tracing, are needed in order to control viral spread. Perinatal transmission has been reported although the exact transmission route has not been elucidated (ECDCa, 2020).

After the initial exposure, patients typically develop symptoms within 5-6 days (incubation period) although about 20% of patients remain asymptomatic throughout infection (Cevik *et al*, 2020). Polymerase chain reaction (PCR) tests can detect viral SARS-CoV-2 RNA in the upper respiratory tract for a mean of 17 days, although transmission is maximal in the first week of illness. Symptomatic and pre-symptomatic transmission (1-2 days before symptom onset), is thought to play a greater role in the spread of SARS-CoV-2 than asymptomatic transmission.

The disease

In adults, the clinical picture varies widely. A significant proportion of individuals are likely to have mild symptoms and may be asymptomatic at the time of diagnosis.

Symptoms are commonly reported as a new onset of cough and fever (Grant *et al*, 2020), but may include headache, loss of smell, nasal obstruction, lethargy, myalgia (aching muscles), rhinorrhea (runny nose), taste dysfunction, sore throat, diarrhoea, vomiting and confusion; fever may not be reported in all symptomatic individuals. Patients may also be asymptomatic (He *et al*, 2020).

Progression of disease, multiple organ failure and death will occur in some individuals (Pachetti *et al*, 2020).

Currently available data suggest that increasing age and male gender are significant risk factors for severe infection. However, there are also groups of patients with underlying comorbidities, where infection may result in increased risk of serious disease (Docherty *et al*, 2020). In a large review of primary care records pseudonymously linked with SARS-CoV-2 status, comorbidities including diabetes, cancer and severe asthma were associated with increased risk of death (Williamson *et al*, 2020).

Infection fatality ratios (IFR) for COVID-19, derived from combining mortality data with infection rates in seroprevalence studies, show a marked increase in IFR in the oldest age groups (Table 1) (Ward *et al*, 2020).

Table 1: Infection fatality ratio and estimated total numbers of deaths (February to July 2020)

Category	Population Size	SARS-CoV-2 antibody prevalence% (95% CI) ¹	Confirmed COVID-19 deaths ²	Infection fatality ratio % (95% CI) ²	Estimated number of infections (95% CI)
Total	56,286,961	6.0 (5.7, 6.8)	30180	0.9 (0.9, 0.9)	3,362,037 (3,216,816; 3,507,258)
Sex					
Male	27,827,831	6.5 (5.8, 6.6)	18575	1.1 (1.0, 1.2)	1,729,675 (1,614,585; 1,844,766)
Female	28,459,130	5.8 (5.4, 6.1)	11600	0.7 (0.7, 0.8)	1,633,785 1,539,821; 1,727,749)
Age (years)					
15-44	21,335,397	7.2 (6.7, 7.7)	524	0.0 (0.0, 0.0)	1,535,884 (1,436,941; 1,634,826)
45-64	14,405,759	6.2 (5.8, 6.6)	4657	0.5 (0.5, 0.5)	895,238 (837,231; 953,244)
65-74	5,576,066	3.2 (2.7, 3.7)	5663	3.1 (2.6, 3.6)	181,044 (153,426; 208,661)
75+	4,777,650	3.3 (2.5, 4.1)	19330	11.6 (9.2, 14.1)	166,077 (131,059; 200,646)

COVID-19 -
SARS-Cov-2

- 1 All estimates of prevalence adjusted for imperfect test sensitivity and specificity (see text for details). Responses have been re-weighted to account for differential sampling (geographic) and for variation in response rate (age, gender, ethnicity and deprivation) in final column to be representative of the England population (18+).
- 2 Infection fatality ratios were calculated excluding care home residents. Confirmed COVID-19 death counts were obtained from <https://fingertips.phe.org.uk/static-reports/mortality-surveillance/excess-mortality-in-England-week-ending-17-jul-2020.html>. Deaths in care homes by age on 12 June 2020 were obtained from www.ons.gov.uk. Total deaths in care residents up to 17 July 2020 were obtained from www.ons.gov.uk. The age stratified estimates of COVID-19 deaths were then estimated using the total deaths from 17 July and the age distribution from 12 June. We assumed that age distribution of deaths did not change between 12 June and 17 July 2020.

In Europe and the UK, deaths attributed to SARS-CoV-2 have been reported disproportionately from residential care homes (ECDCb, 2020, Graham *et al* 2020). Other notable risk groups include healthcare workers (Nguyen *et al*, 2020) who may acquire infection both in the hospital or within the community setting (Bielicki *et al*, 2020). Current evidence suggests that deprivation and being from black and asian minority ethnic groups results in a higher risk for death from SARS-CoV-2 infection (Williamson *et al*, 2020), although the factors that contribute to this are not yet clear.

Children

Fewer than 5% of COVID-19 cases are amongst children and in general they appear to exhibit mild disease. Although cough and fever are the main symptoms in children (Ladhani *et al*, 2020), a UK study tracking children of healthcare workers has recently shown that of those who were seropositive, gastrointestinal symptoms were also commonplace (Waterfield *et al*, 2020). Preliminary evidence suggested that not only do children have a lower susceptibility to SARS-CoV-2 infection, but they are also unlikely to be key drivers of transmission at a population level (Viner *et al*, 2020). However, a recent prospective study found higher secondary attack rates where the household index case was a child (Lopez-Bernal *et al*, 2020).

A spectrum of multi system inflammatory disease similar to Kawasaki disease (KD) was recently described in children admitted during the SARS-CoV-2 pandemic, temporally associated with severe acute respiratory syndrome attributed to SARS-CoV-2 (Paediatric multisystem inflammatory syndrome temporally associated with SARS-CoV-2 infection (PIMS-TS)) (Whittaker *et al*, 2020). This severe presentation in children is extremely rare, but appears to encompass a wide range of features, including fever, gastrointestinal symptoms, rash, myocardial injury and shock (Swann *et al*, 2020).

Pregnant women and neonates

The risk to pregnant women and neonates following COVID-19 infection is generally low: more than half of pregnant women who test positive for SARS-CoV-2 are asymptomatic, and there is no increase in stillbirth or neonatal death rates associated with COVID-19 in pregnancy [Allotey *et al*, 2020]. It is still unclear whether SARS-CoV-2 can be transmitted vertically, and only about 2% of neonates born to COVID-positive mothers in the UK test positive for SARS-CoV-2 in the first 12 hours of life (Vousden *et al*, 2021). However, the risk of preterm birth is increased two to threefold for women with symptomatic COVID-19 (Vousden *et al*, 2021), usually as a result of a medical recommendation to deliver early to improve maternal oxygenation [NICE Guideline 25, 2019 <https://www.nice.org.uk/guidance/ng25>]. Furthermore, a small proportion of pregnant women can have severe or fatal COVID-19. The largest global systematic review indicates that pregnant women are more likely to be admitted to the intensive care unit (ICU) with COVID-19 than age-matched non-pregnant women, (Mullins *et al*, 2021) and there is a signal that this is true in the UK as well (<https://www.icnarc.org/Our-Audit/Audits/Cmp/Reports>).

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).

COVID-19 vaccines

The recognition of the pandemic has 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 include nucleic acid vaccines, inactivated virus vaccines, live attenuated vaccines, protein or peptide subunit vaccines, and viral-vectored vaccines.

Most vaccine candidates focus on immunisation with 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) are expected to be protective (Addetia *et al*, 2020, Thompson *et al*, 2020).

In the UK, three vaccines targeting the S protein have been authorised for supply; two use an mRNA platform (Pfizer BioNTech COVID-19 mRNA vaccine BNT162b2 and Moderna mRNA-1273 COVID-19 vaccine) and the third uses an adenovirus vector (AstraZeneca COVID-19 vaccine).

The Pfizer BioNTech and Moderna COVID-19 vaccines 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 µg 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 100µg 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-CoV2 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 mRNA vaccine BNT162b2

Two doses of Pfizer BioNTech COVID-19 mRNA vaccine BNT162b2 successfully reduced the levels of detectable viral RNA in Rhesus macaques when followed by intra-nasal and intra-tracheal challenge with SARS-CoV-2 (Vogel *et al*, 2020). 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.

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% (with credibility intervals from 90.3% to 97.6%) in those aged 16 years and above. 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 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 mRNA vaccine BNT162b2 received approval to supply in the UK from the MHRA on 2 December 2020.

AstraZeneca COVID-19 vaccine

AstraZeneca COVID-19 vaccine elicited increased neutralisation antibodies in Rhesus macaques as well as a reduction in detectable virus in the lower respiratory tract following challenge with SARS-CoV-2 (Van Doremalen *et al*, 2020). 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 73.00% (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.

Moderna COVID-19 vaccine

In phase 1 testing of the Moderna mRNA-1273 vaccine, all patients seroconverted to IgG by 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 100µg doses of the Moderna mRNA-1273. Phase 3 placebo controlled testing in over 30,000 volunteers, showed a vaccine efficacy of 94.1% (95% CI: 89.3-96.8%). 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% (95% CI 68.8%-99.1%).

The Moderna vaccine was approved for use in the UK in January 2021.

Real world effectiveness

Vaccine effectiveness data from the UK is now emerging. A single dose of either the Pfizer or the AstraZeneca vaccines has been shown to provide around 60% protection against symptomatic disease; vaccinated cases are also around 40% less likely to require hospital admission or to die (LopezBernal *et al*, 2021). This is consistent with protection of around 80% against hospitalisation as seen in local studies (Vasileiou *et al* 2021, AvonCAP, 2021).

Protection against infection has also been seen in healthcare workers, where a single dose of Pfizer vaccine provided more than 70% protection against both symptomatic and asymptomatic infection (Hall *et al* 2021), and in care home residents where a single dose of either Pfizer or AstraZeneca vaccines reduced the risk of infection by around 60% (Shrotri *et al*, 2021). This suggests that vaccination has potential to reduce transmission; this was supported by a Scottish study that showed a 30% reduction in risk of infection in the household members of vaccinated compared to unvaccinated healthcare workers (Shah *et al*, 2021).

Safety

Pfizer BioNTech COVID-19 mRNA vaccine BNT162b2

Local reactions at the injection site are fairly common after Pfizer BioNTech COVID-19 mRNA vaccine BNT162b2, 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). Four cases of Bell's palsy were reported in vaccine recipients in the trial. Although within the expected background rate, this will be monitored closely post-implementation.

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).

Moderna COVID-19 vaccine

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 2, and 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.

Bell's palsy was reported by three participants in the vaccine group and one participant in the placebo group. As for the Pfizer vaccine, this will be monitored closely post-implementation. There were no cases of severe COVID-19 disease in the vaccine group, and thus no signal for enhanced disease (Baden *et al*, 2020).

AstraZeneca COVID-19 vaccine

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 ($>38^{\circ}\text{C}$) 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 can be 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).

Recently, a 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 5 per million, although a higher incidence appears to be seen in younger individuals. Overall, the JCVI, MHRA and the WHO remain clear that the benefits of vaccination outweigh this small risk for adults aged 30 years and over, adults who are clinically extremely vulnerable and those with underlying clinical risks as defined in table 3.

Storage

The Pfizer BioNTech COVID-19 mRNA vaccine BNT162b2 should be stored in a freezer at -80°C to -60°C (-90°C to -60°C in the thermal container). Shelf life is 6 months at -80°C to -60°C . Frozen vials should be transferred to 2°C to 8°C to thaw; 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, stability data have demonstrated that undiluted vaccine can be stored for up to 5 days at 2°C to 8°C. Once thawed, the vaccine cannot be re-frozen.

The AstraZeneca vaccine should be stored at 2°C to 8°C and has a shelf life of 6 months. 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 vials should be stored frozen between -25° to -15°C and has a shelf life of 7 months at these temperatures. 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 vial is stable for 12 hours at 8° to 25°C.

Presentation

Each pack of the Pfizer BioNTech COVID-19 mRNA vaccine BNT162b2 contains 195 vials with a minimum of 5 doses per vial (975 doses per pack). It is supplied with 0.9% sodium chloride diluent for injection in plastic ampoules. After dilution, the vaccine should be kept at 2°C to 25°C and used within 6 hours. Any unused vaccine should be discarded.

The AstraZeneca vaccine 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 is supplied in multidose vials containing 10 doses of 0.5ml.

Dosing and schedule

Pfizer BioNTech COVID-19 mRNA vaccine BNT162b2

The dose of Pfizer BioNTech COVID-19 vaccine is 30µg contained in 0.3ml of the diluted vaccine. After dilution each multidose vial can be used to deliver five or six doses of 0.3ml.

The vaccine should be administered in two doses, a minimum of 21 days apart.

AstraZeneca COVID-19 vaccine

The dose of AstraZeneca COVID-19 vaccine is 0.5ml.

Based on good evidence of higher clinical protection, JCVI currently recommend that, ideally, an eight week minimum interval should be observed for this vaccine. An interval of 28 days may be observed when rapid protection is required (for example for those about to receive immunosuppressive treatment).

Moderna COVID-19 vaccine

The dose of Moderna COVID-19 vaccine is 0.5ml.

The vaccine should be administered in two doses, a minimum of 28 days apart.

All COVID-19 vaccines

Operationally, it is recommended that a consistent interval should be used for all vaccines to avoid confusion and simplify booking. Currently, a schedule of around 12 weeks is being followed to allow more people to benefit from the protection provided from the first dose during the roll out phase. Longer term protection will then be provided by the second dose.

If an interval longer than the recommended interval is left between doses, the second dose should still be given (preferably using the same vaccine as was given for the first dose if possible). The course does not need to be restarted.

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 mRNA vaccine BNT162b2 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. The vial should be discarded if the solution is discoloured or visible particles are observed.

AstraZeneca COVID-19 vaccine 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 should be administered as an intramuscular injection. 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.

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 (ACIP 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 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 offensive waste bags.

The COVID-19 immunisation programme

Phase 1 recommendations for the use of the vaccine

The objectives of first phase of the COVID-19 immunisation programme is to protect those who are at highest risk from serious illness or death. The Joint Committee of Vaccination and Immunisation (JCVI) has set out a prioritisation for persons at risk. JCVI ranked the eligible groups according to risk, largely based on prevention of COVID-19-specific mortality.

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 with age. Residents in care homes for older adults have been disproportionately affected by the COVID-19 pandemic. Table 2 sets out JCVI advice on priority groups for COVID-19 vaccination. Table 3 sets out JCVI advice on clinical and other risk groups for COVID-19 vaccination.

Table 2 – Priority groups for 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 Clinically extremely vulnerable individuals (not including those under 16 years of age)
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

Clinically extremely vulnerable (group 4) and adults in priority group 6

People who are defined as clinically extremely vulnerable (CEV) are 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>); these patients should be flagged on the GP system. A hospital clinician or GP can also add a patient to the list, based on their clinical judgement, because they consider them to be at very high risk of serious illness from COVID-19.

Many individuals considered CEV are in the oldest age groups and will be among the first to receive vaccine. Given the level of risk seen in this group as a whole, the remainder of the CEV group should be offered vaccine alongside those 70-74 years of age.

Children who are on the CEV list are not generally recommended for vaccination unless they have serious neurodisabilities (see section on children).

All patients on the CEV list will also fall into the broader disease categories outlined in table 3, but are in priority group 4 because of more recent treatment, more advanced condition or co-morbidities. Other patients in the same clinical risk group, but not on the CEV list at the time group 4 is called, should be called in priority group 6, or with their appropriate age cohort.

Table 3 Clinical and other risk groups 16 years of age and over who should receive COVID-19 immunisation.

Clinical risk groups	
Chronic respiratory disease	Individuals with a severe lung condition, including those with asthma that requires continuous or repeated use of systemic steroids or with previous exacerbations requiring hospital admission, 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 disease (e.g. polio syndrome sufferers). This includes individuals with cerebral palsy, severe or profound learning disabilities, 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	Any diabetes, including diet-controlled diabetes.
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 and those with systemic lupus erythematosus and rheumatoid arthritis, and psoriasis who may require long term immunosuppressive treatments.</p>

	<p>Most of the more severely immunosuppressed individuals in this group should already be flagged as CEV. Individuals who are not yet on the CEV list but who are about to receive highly immunosuppressive interventions or those whose level of immunosuppression is about to increase may be therefore be offered vaccine alongside the CEV group, if therapy can be safely delayed or there is sufficient time (ideally two weeks) before therapy commences.</p> <p>Some immunosuppressed patients may have a suboptimal immunological response to the vaccine (see Immunosuppression and HIV).</p>
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	<p>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.</p> <p>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).</p> <p>For consideration of children under 16 see below.</p>
Other risk groups	
Adult household contacts of people with immunosuppression	Individuals who expect to share living accommodation on most days (and therefore for whom continuing close contact is unavoidable) with individuals who are immunosuppressed (defined as above).
Adult carers	Those who are eligible for a carer's allowance, or those 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. ¹

The examples above are not exhaustive, and, within these groups, the prescriber should 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 list of eligible diagnoses, and the appropriate clinical codes, can be found in the link at the end of the chapter on page 13.

Recommendations by staff groups

The objective of occupational immunisation of health and social care staff is to protect workers at high risk of exposure who may also expose vulnerable individuals whilst providing care. There is increasing evidence that vaccination will lead to a reduction in transmission,

¹ Those clinically vulnerable to COVID-19 include children with severe neuro-disabilities, those who are designated Clinically Extremely vulnerable (CEV), adults who have underlying health conditions (as defined in table 3), and those who need care because of advanced age. Eligible carers should be vaccinated in priority group 6.

and even a small effect will have major additional benefit for staff who could expose multiple vulnerable patients and other staff members. Potential exposure to COVID-19, and therefore the priority for vaccination, may vary from workplace to workplace. Therefore, it is recommended that these staff groups are vaccinated against COVID-19 as follows.

Frontline healthcare staff

This includes the following groups:

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 must 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

This would include:

- 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
- social care staff directly involved in the care of their patients or clients
- others involved directly in delivering social care such that they and vulnerable patients/clients are at increased risk of exposure

Young people age 16-18 years, who are employed in, studying or in training for health and social care work should be offered vaccination alongside their colleagues, if a suitable vaccine is available (see section on age specific recommendations). Younger people who are taking part in health and social care work as volunteers, interns or for the purposes of

work experience, should make all efforts to avoid exposure to infection; vaccination would not normally be required.

Phase 2 advice

The objectives of the second phase of the COVID-19 immunisation programme is to continue to protect those who are at highest risk from serious illness or death, but also to protect the NHS by reducing the risks of hospitalisation and critical care admission. Phase 2 of the programme should therefore be accompanied by continued efforts to maximise coverage amongst those prioritised in Phase 1 but who remain 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 increase with age. **JCVI has therefore advised that the offer of vaccination during Phase 2 is age-based starting with the oldest adults first** and proceeding in the following order:

All those aged 40-49 years

All those aged 30-39 years

All those aged 18-29 years

As there is an increased risk of hospitalisation in males, those from certain ethnic minority backgrounds, those who are obese or morbidly obese, and those from socio-economically deprived areas, JCVI advises that specific focus should be used to promote and deliver vaccination to these groups.

Age specific recommendations on vaccine type

Children under 16 and young adults age 16-18 years

The Pfizer BioNTech vaccine has approval for use from 16 years old and currently has the most extensive safety data in those aged 12-15 years. This vaccine is therefore the preferred vaccine in this age group. It is likely that Moderna will have similar safety profile and may therefore be used as an alternate, "off-licence". Young people who have had a first dose of AstraZeneca vaccine, however, should complete with the same vaccine (see contraindications and precautions).

Healthy adults aged 18-30 years

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 evidence JCVI are advising a preference for a vaccine other than AstraZeneca to be offered to healthy people under 30 years of age, including health and social care workers, unpaid carers and household contacts of immunosuppressed individuals. In the absence of a suitable alternative these individuals may defer or choose to receive the AstraZeneca vaccine provided they have been informed and understand the relative risks and benefits. They should be given the latest version of the COVID-19 vaccination and blood clotting leaflet (<https://www.gov.uk/government/publications/covid-19-vaccination-and-blood-clotting>). Those who have already received a dose of AstraZeneca vaccine should complete with the same vaccine (see contraindications and precautions).

Aged over 30 and in high risk groups

Individuals in aged over 30 years, those who are clinically extremely vulnerable and those clinical risk groups (see table 3) are at high risk of the complications of COVID-19. The JCVI, MHRA and WHO consider that any risk from the rare side effects seen after AstraZeneca vaccine are outweighed by the benefits of vaccination. These individuals can therefore be offered vaccination with any of the available products, unless otherwise contra-indicated.

Pregnant women of any age

Pfizer and Moderna vaccines are the preferred vaccines for pregnant women of any age, because of more extensive experience of their use in pregnancy. Pregnant women who commenced vaccination with AstraZeneca, however, are advised to complete with the same vaccine (see section on pregnancy).

Previous incomplete vaccination

If the course is interrupted or delayed, it should be resumed using the same vaccine but the first dose should not be repeated. There is no evidence on the interchangeability of the COVID-19 vaccines although studies are underway. Therefore, every effort should be made to determine which vaccine the individual received and to complete with the same vaccine. For individuals who started the schedule and who attend for vaccination at a site where the same vaccine is not available, or if the first product received is unknown, it is reasonable to offer one dose of the locally available product to complete the schedule. This option is preferred if the individual is likely to be at immediate high risk or is considered unlikely to attend again. In these circumstances, as the vaccines are based on the spike protein, it is likely the second dose will help to boost the response to the first dose. For this reason, until additional information becomes available, further doses would not then be required.

Individuals who have received a first dose of the AstraZeneca vaccine should complete the course with the same vaccine, with the exception of those who experience an episode of thrombosis combined with thrombocytopenia (see contraindications and precautions).

Individuals who are participating in a clinical trial of COVID-19 vaccines who present for vaccination should be referred back to the investigators. Eligible persons who are enrolled in vaccine trials should then be provided with written advice on whether and when they can be safely vaccinated in the routine programme.

Individuals who have been vaccinated abroad are likely to have received either an mRNA or vector vaccine based on the spike protein, or an inactivated whole viral vaccine. Specific advice on completing vaccination in these individuals is available from Public Health England. <https://www.gov.uk/government/publications/covid-19-vaccination-programme-guidance-for-healthcare-practitioners>

Reinforcing immunisation

Booster doses of COVID-19 vaccine are not yet recommended because the need for, and timing of, boosters has not yet been determined.

Co-administration with other vaccines

Although no data for co-administration of COVID-19 vaccine with other vaccines exists, 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.

Because of the absence of data on co-administration with COVID-19 vaccines, it should not be routine to offer appointments to give this vaccine at the same time as other vaccines. Based on current information about the first COVID-19 vaccines being deployed, scheduling should ideally be separated by an interval of at least 7 days to avoid incorrect attribution of potential adverse events.

As all of the early COVID-19 vaccines are considered inactivated (including the non-replicating adenovirus vaccine), where individuals in an eligible cohort present having received another inactivated or live vaccine, COVID-19 vaccination should still be considered. The same applies for other live and inactivated vaccines where COVID-19 vaccination has been received first or where a patient presents requiring two vaccines. In most cases, vaccination should proceed to avoid any further delay in protection and to avoid the risk of the patient not returning for a later appointment. In such circumstances, patients should be informed about the likely timing of potential adverse events relating to each vaccine.

Pregnancy

There is no known risk associated with giving inactivated, recombinant viral or bacterial vaccines or toxoids during pregnancy or whilst breast-feeding (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.

Although clinical trials on the use of COVID-19 vaccines during pregnancy are not advanced, the available data do not indicate any harm to pregnancy. JCVI has therefore advised that women who are pregnant should be offered vaccination at the same time as non-pregnant women, based on their age and clinical risk group. There is now extensive post-marketing experience of the use of the Pfizer BioNTech and Moderna vaccines in the USA with no safety signals so far (<https://www.cdc.gov/coronavirus/2019-ncov/vaccines/safety/vsafepregnancyregistry.html>). These vaccines are therefore the preferred vaccines to offer to pregnant women. Clinicians should discuss the risks and benefits of vaccination with the woman, who should be told about the limited evidence of safety for the vaccine in pregnancy.

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 can be vaccinated with a suitable product for their age and clinical risk group.

If a woman finds out she is pregnant after she has started a course of vaccine, she may complete vaccination during pregnancy using the same vaccine product (unless contra-indicated). Alternatively, vaccination should be offered as soon as possible after pregnancy.

Termination of pregnancy following inadvertent immunisation should not be recommended. Surveillance of the inadvertent administration of COVID-19 vaccines in early pregnancy is being conducted for the UK by the PHE Immunisation Department, to whom such cases

should be reported <https://www.gov.uk/guidance/vaccination-in-pregnancy-vip>. As above, women who are inadvertently vaccinated in early pregnancy should be offered the second dose of the same product.

Breastfeeding

There is no known risk associated with being given a non-live vaccine whilst breastfeeding. JCVI advises that breastfeeding women may be offered any suitable COVID-19 vaccine.

The developmental and health benefits of breastfeeding should be considered along with the mother's clinical need for immunisation against COVID-19; at the same time, women should be informed about the absence of full safety data for the vaccine in breastfeeding.

Children

SARS-CoV-2 vaccine trials have only just begun in children and there are, therefore, very limited data on safety and immunogenicity in this group. Children and young people have a very low risk of COVID-19, severe disease or death due to SARS-CoV-2 compared to adults and so COVID-19 vaccines are not routinely recommended for children and young people under 16 years of age.

Children under 16 year of age, even if they are CEV, are at low risk of serious morbidity and mortality, and, given the absence of safety and efficacy data on the vaccine, are not recommended for vaccination. Limited data however, suggest that children with neurological comorbidities may be at a greater risk of developing severe COVID-19.

Given the very high risk of exposure to infection and outbreaks in institutional settings, vaccination may be considered for children with severe neuro-disabilities who tend to get recurrent respiratory tract infections and who frequently spend time in specialised residential care settings for children with complex needs. As older children have higher risk of acquiring and becoming sick from infection and there are some safety data on the Pfizer BioNTech COVID-19 mRNA vaccine BNT162b2 in children aged 12 years and older, vaccination of older children in these settings should be considered with this vaccine. If the Pfizer vaccine is not available, Moderna mRNA vaccine may be used "off-license" as an alternate.

Recommendations on vaccinating children with other underlying conditions will be reviewed after the initial roll-out phase by which time additional data on use of the vaccines in adults should allow a better assessment of risks and benefits.

Immunosuppression and HIV

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). Although individuals with stable treated HIV infection were not excluded from the phase 3 trial of the Pfizer and Moderna mRNA vaccines, data on safety and effectiveness in this group have not been presented. A study of the AstraZeneca vaccines in people living with HIV infection is underway.

Individuals with immunosuppression may not make a full immune response to vaccination. As there is no evidence on response in immunosuppressed individuals there is also no evidence upon which to base advice on the optimal timing of delivery. A recent study

suggested immune responses were better, however, in patients with cancer who received their chemotherapy at least two weeks earlier (Monin-Aldama *et al*, 2021). 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. The small number of patients who are about to receive planned immunosuppressive therapy should be considered for vaccination prior to commencing therapy (ideally at least two weeks before), when their immune system is better able to make a response. Where possible, it would also be preferable for the 2-dose schedule to be completed prior to commencing immunosuppression. This would entail offering the second dose at the recommended minimum for that vaccine (three or four weeks from the first dose) to provide maximum benefit that may not be received if the second dose was given during the period of immunosuppression. Any decision to defer immunosuppressive therapy or to delay possible benefit from vaccination until after therapy should not be taken without due consideration of the risks from COVID-19 and from their underlying condition. Although the immune correlates of protection are currently unknown, post-vaccination testing for spike antibody may be considered. Until further information becomes available vaccinated patients with immunosuppression should continue to follow advice to reduce the chance of exposure, and their adult household contacts should also be offered vaccine.

Contraindications and precautions

There are very few individuals who cannot receive the Pfizer BioNTech, Moderna or AstraZeneca COVID-19 vaccines. 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 vaccine should not be given to those who have had a previous systemic allergic reaction (including immediate-onset anaphylaxis) to:

- a previous dose of the same COVID-19 vaccine
- any component (excipient) of the COVID-19 vaccine e.g. polyethylene glycol (PEG)

Further advice on anaphylaxis and on the thrombosis and thrombocytopaenia syndrome are outlined below.

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.

There is no evidence of any 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. Vaccination should be deferred in those with confirmed infection to avoid confusing the differential diagnosis. As clinical deterioration can occur up to two weeks after infection, ideally vaccination should be deferred until clinical recovery to around four weeks after onset of symptoms or four weeks from the first confirmed positive specimen in those who are asymptomatic.

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.

Anaphylaxis following COVID-19 vaccination

A very small number of individuals have experienced anaphylaxis when vaccinated with the Pfizer BioNTech vaccine. Following close national surveillance, the MHRA is **no longer** advising that individuals with a history of anaphylaxis to any vaccine, medicine or food do not get the vaccine. Anyone with a previous history of allergic reactions to the ingredients of the vaccine should not receive it, but those **with any other allergies** (such as a food or penicillin allergy) **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 but would contraindicate receipt of this vaccine (Sellaturay *et al*, 2020). It is unclear whether PEG is the only cause of allergic reactions in patients with systemic allergic symptoms after the first dose of Pfizer-BioNTech vaccine.

The rate of anaphylaxis reported to date to the AstraZeneca vaccine is in line with the expected rate of anaphylaxis to non-COVID vaccines. The AstraZeneca vaccine does not contain PEG but does contain a related compound called polysorbate 80. Some 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 (such as certain influenza vaccines) are likely to tolerate the AstraZeneca vaccine. This is summarised in table 4.

Table 4: Management of patients with a history of allergy

	Proceed with vaccination	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) family history of allergies 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"> 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 systemic allergic reaction to the COVID-19 vaccine for an mRNA-based COVID-19 vaccine, prior allergic reaction to another mRNA vaccine prior allergic reaction to a component of the vaccine, including PEG
ACTIONS	<ul style="list-style-type: none"> proceed with vaccination as normal, according to local guidelines 	<ul style="list-style-type: none"> discuss with allergy specialist and consider possibility of PEG-allergy consider observation for 30 minutes if vaccination proceeds (see precautions) some patients may benefit from pretreatment with antihistamine, however this may mask initial symptoms of a reaction 	<ul style="list-style-type: none"> do not give vaccine in question refer to allergist

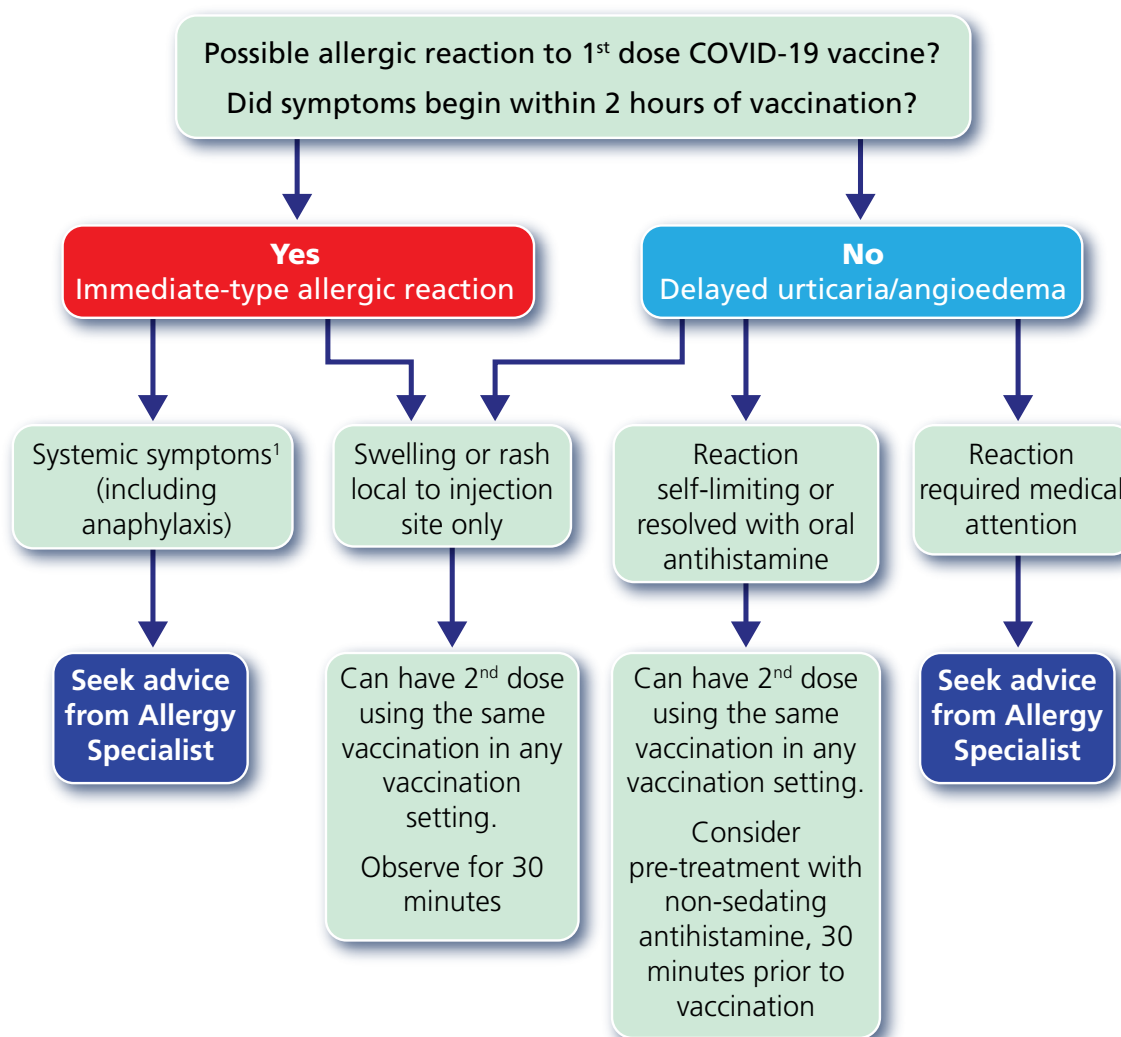
All recipients of the Pfizer BioNTech and Moderna vaccines should be kept for observation and monitored for a minimum of 15 minutes. Facilities for management of anaphylaxis should be available at all vaccination sites (see chapter 8). Advice has also been issued by the Resuscitation Council.¹

Patients with undiagnosed PEG allergy often have a history of immediate onset-unexplained anaphylaxis or anaphylaxis to multiple classes of drugs or an unexplained anaphylaxis. Such individuals should not be vaccinated with the Pfizer BioNTech or Moderna vaccines, except on the expert advice of an allergy specialist. The AstraZeneca vaccine can be used as an alternative (unless otherwise contraindicated), particularly if they previously tolerated an injected 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.

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 (figure).

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

Figure: Flowchart for managing patients who have allergic reactions to the first dose of COVID-19 vaccine



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.

Thrombosis and thrombocytopenia occurring after COVID-19 vaccination

A recently recognised condition involving serious thromboembolic events accompanied by thrombocytopaenia, 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. This may be related to the recipient's polymorphisms in genes encoding Fc receptors in the immune system and is an area of active research. 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 thrombocytopaenia after the AstraZeneca vaccine. There have been no confirmed cases reported in pregnant women to date.

The contra-indications to vaccination with the AstraZeneca COVID-19 vaccine have now been amended to include individuals who have a history of a previous episode of heparin-induced thrombocytopenia and thrombosis (HITT or HIT type 2). These individuals may be offered vaccination with an alternative COVID-19 vaccine.

Individuals who experience a clotting episode with concomitant thrombocytopaenia 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, and they should then be considered for a second dose of an alternative product. Based on the analogy of HITT, antibodies to platelet factor 4 may persist for around six months.

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 of around 12 weeks. To date, there have been no confirmed cases 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. Using an alternative product for the second dose may be more likely to lead to common side effects.

Based on current evidence JCVI is advising a preference for an alternative vaccine for healthy people under 30 years of age, including health and social care workers, unpaid carers and household contacts of immunosuppressed individuals.

Individuals over 30 years of age 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 thrombocytopaenia 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 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 thrombocytopaenia.

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 Coronavirus Yellow Card Scheme (www.coronavirus-yellowcard.mhra.gov.uk). [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.yellowcard.gov.uk). 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 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>

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.

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.

References

- Addetia A, Crawford KHD, Dingens A, *et al* (2020) Neutralizing antibodies correlate with protection from SARS-CoV-2 in humans during a 1 fishery vessel outbreak with high attack rate. *J Clin Microbiol* 58(11): e2107-20
- Advisory Committee on Immunization Practices (2019). General Best Practice Guidelines for Immunization: Best Practices Guidance of the Advisory Committee on Immunization Practices (ACIP). Special Situations <https://www.cdc.gov/vaccines/hcp/acip-recs/general-recs/special-situations.html>
- Allotey J, Bonet M, Kew T, *et al*. (2020) Clinical manifestations, risk factors, and maternal and perinatal outcomes of coronavirus disease 2019 in pregnancy: Living systematic review and meta-analysis. *Br Med J* 370:m3320.
- Amanat F, Krammer F. SARS-CoV-2 Vaccines: Status Report. *Immunity*. 2020 Apr 14; 52(4):583-589. doi: 10.1016/j.immuni.2020.03.007. Epub 2020 Apr 6.
- Baden LR, El Sahly HM, Essink B *et al*. Efficacy and safety of the mRNA-1273 SARS-CoV-2 vaccine. *NEJM* 2020. doi: 10.1056/NEJMoa2035389.
- Bielicki JA, Duval X, Gobat N, *et al*. Monitoring approaches for health-care workers during the COVID-19 pandemic. *Lancet Infect Dis*. 2020 Oct;20(10):e261-e267. doi: 10.1016/S1473-3099(20)30458-8.
- Cevik M, Kuppalli K, Kindrachuk J, Peiris M. Virology, transmission, and pathogenesis of SARS-CoV-2. *BMJ*. 2020 Oct 23;371:m3862. doi: 10.1136/bmj.m3862.
- Department of Health, 2013. Health Technical Memorandum 07-01 – Safe management of healthcare waste. <https://www.gov.uk/government/publications/guidance-on-the-safe-management-of-healthcare-waste>
- Dhama, K, Sharun K, Tiwari R, *et al*. Coronavirus Disease 2019 - COVID-19. *Clinical Microbiology Reviews* 2020, 33(4): DOI: 10.1128/CMR.00028-20
- Diggle L and Deeks J (2000). Effect of needle length on incidence of local reactions to routine immunisation in infants aged 4 months: randomised controlled trial. *BMJ* 321(7266): 931-3.
- Docherty A B, Harrison EM, Green CA, Hardwick HE, Pius R, Norman L *et al*. Features of 20 133 UK patients in hospital with covid-19 using the ISARIC WHO Clinical Characterisation Protocol: prospective observational cohort study *BMJ* 2020; 369 :m1985
- ECDCa. Risk factors and risk groups. <https://www.ecdc.europa.eu/en/covid-19/latest-evidence/epidemiology>. (Accessed: 3rd October 2020). <https://www.ecdc.europa.eu/en/covid-19/latest-evidence/epidemiology>
- ECDCb. Surveillance of COVID-19 at long term care facilities in the EU/EEA. <https://www.ecdc.europa.eu/en/publications-data/surveillance-COVID-19-long-term-care-facilities-EU-EEA>
- Elshafeey F, Magdi R, Hindi N *et al*. A systematic scoping review of COVID 19 during pregnancy and childbirth. *Int J Gynaecol Obstet*. 2020 Jul;150(1):47-52. doi: 10.1002/ijgo.13182.
- Folegatti, P. M. *et al*. Safety and immunogenicity of the ChAdOx1 nCoV-19 vaccine against SARS-CoV-2: a preliminary report of a phase 1/2 single-blind, randomised controlled trial. (2020) *Lancet* 396:467-8
- Garafalo M, Staniszewska M, Salmaso S, *et al*. Prospects of Replication-Deficient Adenovirus Based Vaccine Development against SARS- CoV-2. *Vaccines (Basel)*. 2020 Jun 10;8(2):293. doi: 10.3390/vaccines8020293.
- Graham NSN, Junghans C, Downes R, *et al*. SARS-CoV-2 infection, clinical features and outcome of COVID-19 in United Kingdom nursing homes. *J Infect*. 2020 Sep;81(3):411-419. doi: 10.1016/j.jinf.2020.05.073.
- Grant MC, Geoghegan L, Arbyn M, *et al*. The prevalence of symptoms in 24,410 adults infected by the novel coronavirus (SARS- CoV-2; COVID-19): A systematic review and meta-analysis of 148 studies from 9 countries. *PLoS One* 2020 Jun 23;15(6): e0234765. doi: 10.1371/journal.pone.0234765.
- Greinacher A, Thiele T, Warkentin TE *et al*. Thrombotic Thrombocytopenia after ChAdOx1 nCov-19 Vaccination. *NEJM*, 2021. DOI: 10.1056/NEJMoa2104840
- Hall VJ, Foulkes S, Saei A, Andrews N, Oguti B *et al*. Effectiveness of BNT162b2 mRNA Vaccine Against Infection and COVID-19 Vaccine Coverage in Healthcare Workers in England, Multicentre Prospective Cohort Study (the SIREN Study). Available at SSRN: <https://ssrn.com/abstract=3790399> or <http://dx.doi.org/10.2139/ssrn.3790399>
- He J, Guo Y, Mao R, Zhang J. Proportion of asymptomatic coronavirus disease 2019: A systematic review and meta-analysis. *J Med Virol*. 2020 Jul 21;10.1002/jmv.26326. doi: 10.1002/jmv.26326.
- Huang C, Wang Y, Li X, Ren L, Zhao *et al*. Clinical features of patients infected with 2019 novel coronavirus in Wuhan, China. *Lancet*. 2020 Feb 15;395 (10223):497-506. doi: 10.1016/S0140-6736(20) 30183-5.

- Hyams C, Marlow R, Maseko Z, King J and Ward, L. Assessing the Effectiveness of BNT162b2 and ChAdOx1nCoV-19 COVID-19 Vaccination in Prevention of Hospitalisations in Elderly and Frail Adults: A Single Centre Test Negative Case-Control Study. Available at SSRN: <https://ssrn.com/abstract=3796835> or <http://dx.doi.org/10.2139/ssrn.3796835>
- Jackson LA, Anderson EJ, Roupael NG *et al.* An mRNA vaccine against SARS-CoV-2 - preliminary report. *NEJM* 2020; 383: 1920-31.
- Karimi-Zarchi M, Neamatzadeh H, Dastgheib SA, *et al.* Vertical Transmission of Coronavirus Disease 19 (COVID-19) from Infected Pregnant Mothers to Neonates: A review. *Fetal Pediatr Pathol.* 2020 Jun;39(3): 246-250. doi: 10.1080/15513815.2020.
- Kaur SP, Gupta V. COVID-19 Vaccine: A comprehensive status report. *Virus Res.* 2020 Oct 15; 288:198114. Doi: 10.1016/j.viruses.2020.198114.
- Kennedy s, Bolay F, Keih M, *et al* (2017) Phase 2 Placebo-Controlled Trial of Two Vaccines to Prevent Ebola in Liberia. *N Eng J Med* 377: 1438-1447.
- Kroger AT, Atkinson WL, Pickering LA. General immunization Practices. in Plotkin SA, Orenstein WA, Offit PA. *Vaccines* (6th Edition). Elsevier Saunders 2013.
- Ladhani SN, Amin-Chowdhury Z, Davies HG, *et al.* COVID-19 in children: analysis of the first pandemic peak in England. *Arch Dis Child.* 2020 Dec;105(12): 1180-1185. doi: 10.1136/archdischild-2020-320042.
- Lam TT, Jia N, Zhang YW, Shum MH, *et al.* Identifying SARS-CoV-2- related coronaviruses in Malayan pangolins. *Nature.* 2020 Jul;583 (7815):282-285. doi: 10.1038/s41586-020-2169-0.
- Lillie PJ, Samson A, Li A, *et al.* Novel coronavirus disease (Covid-19): The first two patients in the UK with person to person transmission. *J Infect.* 2020 May;80(5):578-606. doi: 10.1016/j.jinf.2020.02.020. Epub 2020 Feb 28.
- Lopez Bernal, J. *et al.* Transmission dynamics of COVID-19 in household and community settings in the United Kingdom. (2020) <https://doi.org/10.1101/2020.08.19.20177188>.
- Lopez Bernal J, Andrews A, Gower C, *et al.* Early effectiveness of COVID-19 vaccination with BNT162b2 mRNA vaccine and ChAdOx1 adenovirus vector vaccine on symptomatic disease, hospitalisations and mortality in older adults in England <https://www.medrxiv.org/content/10.1101/2021.03.01.21252652v1>
- Mark A, Carlsson RM and Granstrom M (1999) Subcutaneous versus intramuscular injection for booster DT vaccination of adolescents. *Vaccine* 17(15-16): 2067-72.
- Mullins E, Hudak M, Banerjee J, *et al.* Pregnancy and neonatal outcomes of COVID-19 – coreporting of common outcomes from the PAN-COVID and AAP SONPM registry. <https://doi.org/10.1101/2021.01.06.21249325>
- Monin-Aldama L, Laing AG, Muñoz-Ruiz M. Interim results of the safety and immune-efficacy of 1 versus 2 doses of COVID-19 vaccine BNT162b2 for cancer patients in the context of the UK vaccine priority guidelines. <https://doi.org/10.1101/2021.03.17.21253131>
- Nguyen LH, Drew DA, Graham MS *et al.* Risk of COVID-19 among front-line health-care workers and the general community: a prospective cohort study. *Lancet Public Health.* 2020 Sep;5(9): e475-e483. doi: 10.1016/S2468-2667(20) 30164-X. Epub 2020 Jul 31.
- Pachetti M, Marini B, Giudici F, *et al.* Impact of lockdown on COVID-19 case fatality rate and viral mutations spread in 7 countries in Europe and North America. *J Transl Med.* 2020 Sep 2;18(1):338. doi: 10.1186/s12967-020-02501-x.
- Polack, FP, Thomas SJ, Kitchin N *et al.* Safety and Efficacy of the BNT162b2 mRNA Covid-19 Vaccine. *NEJM* 2020. DOI: 10.1056/NEJMoa2034577
- Ramasamy MN, Minassian AM, Ewer KJ, *et al.* Safety and immunogenicity of ChAdOx1 nCoV-19 vaccine administered in a prime-boost regimen in young and old adults (COV002): a single-blind, randomised, controlled, phase 2/3 trial. *Lancet* 2020 Nov 18: S0140-6736(20)32466-1. doi: 10.1016/S0140-6736(20)32466-1. [https://doi.org/10.1016/S0140-6736\(20\)32466-1](https://doi.org/10.1016/S0140-6736(20)32466-1).
- Sanche S, Lin YT, Xu C, Romero-Severson E, Hengartner N, Ke R. High Contagiousness and Rapid Spread of Severe Acute Respiratory Syndrome Coronavirus 2. *Emerg Infect Dis.* 2020 Jul;26(7):1470-1477. doi: 10.3201/eid2607.200282.
- Sellaturay P, Nasser S, Ewan P. Polyethylene Glycol-Induced Systemic Allergic Reactions (Anaphylaxis). *J Allergy Clin Immunol Pract.* 2020 Oct 1:S2213-2198(20)31007-2. doi: 10.1016/j.jaip.2020.09.029. Epub ahead of print. PMID: 33011299.
- Shah A, Gribben C, Bishop J *et al.* Effect of vaccination on transmission of COVID-19: an observational study in healthcare workers and their households View ORCID Profile doi: <https://doi.org/10.1101/2021.03.11.21253275>

- Shrotri M, Krutikov M, Palmer T *et al.* Vaccine effectiveness of the first dose of ChAdOx1 nCoV-19 and BNT162b2 against SARS-CoV-2 infection in residents of Long-Term Care Facilities (VIVALDI study). medRxiv [preprint], doi: 10.1101/2021.03.26.21254391
- Swann OV, Holden KA, Turtle L, Pollock L, Fairchild CJ, Drake TM *et al.* Clinical Characteristics of children and young people admitted to hospital with COVID-19 in United Kingdom: prospective multicentre observational cohort study BMJ 2020; 370:m3249
- Thompson CP, Grayson NE, Paton RS, *et al.* Detection of neutralising antibodies to SARS-CoV-2 to determine population exposure in Scottish blood donors between March and May 2020. Euro Surveill. 2020 Oct;25(42):2000685. doi: 10.2807/1560-7917.ES.2020.25.42.2000685
- van Doremalen N, Lambe T, Spencer A, *et al.* ChAdOx1 nCoV-19 vaccination prevents SARS-CoV-2 pneumonia in rhesus macaques. 2020 Oct;586 (7830):578. doi: 10.1038/s41586-020-2608-y.
- Vasileiou E, Simpson CR, Robertson C, *et al.* Effectiveness of First Dose of COVID-19 Vaccines against hospital admissions in Scotland: National prospective cohort study of 5.4 million people. Available at: https://papers.ssrn.com/sol3/papers.cfm?abstract_id=3789264 (pre-print)
- Viner RM, Mytton OT, Bonell C, *et al.* Susceptibility to SARS-CoV-2 Infection Among Children and Adolescents Compared with Adults: A Systematic Review and Meta-analysis. JAMA Pediatr. 2020 Sep 25:e204573. doi: 10.1001/jamapediatrics.2020.4573. Epub ahead of print. PMID:32975552; PMCID: PMC7519436.
- Vogel, A. *et al.* A prefusion SARS-CoV-2 spike RNA vaccine is highly immunogenic and prevents lung infection in non-human primates. (2020) <https://www.biorxiv.org/content/10.1101/2020.09.08.280818v1>
- Vousden N, Bunch K, Morris E MD. The incidence, characteristics and outcomes of pregnant women hospitalized with symptomatic and 2 asymptomatic SARS-CoV-2 infection in the UK from March to September 2020: a national cohort 3 study using the UK Obstetric Surveillance System (UKOSS). <https://www.medrxiv.org/content/10.1101/2021.01.04.21249195v1>
- Voysey M, Clemens S, Shabir AM *et al.* Safety and efficacy of the ChAdOx1 nCoV-19 vaccine (AZD1222) against SARS-CoV-2: an interim analysis of four randomised controlled trials in Brazil, South Africa, and the UK. Lancet 2020. [https://doi.org/10.1016/S0140-6736\(20\)32661-1](https://doi.org/10.1016/S0140-6736(20)32661-1)
- Walsh EE, Frenck RW Jr, Falsey AR, *et al.* Safety and Immunogenicity of Two RNA-Based COVID-19 Vaccine Candidates. N Engl J Med 2020 Oct 14;NEJMoa2027906. doi: 10.1056/NEJMoa2027906
- Ward H, Atchison CJ, Whitaker M, *et al.*, 2020. Antibody prevalence for SARS-CoV-2 in England following first peak of the pandemic: REACT2 study in 100,000 adults <https://www.medrxiv.org/content/10.1101/2020.08.12.20173690v2>
- Waterfield T, Watson C, Moore R, *et al.* Seroprevalence of SARS-CoV2 antibodies in children: a prospective multicentre cohort study. Arch Dis Child. 2020 Nov 10; archdischild-2020-320558. doi: 10.1136/archdischild-2020-320558.
- Whittaker E, Bamford A, Kenny J, *et al.* Clinical Characteristics of 58 Children With a Pediatric Inflammatory Multisystem Syndrome Temporally Associated with SARS-CoV-2. JAMA. 2020 Jul 21;324(3):259-269. doi: 10.1001/jama.2020.10369.
- WHO Director-General's opening remarks at the media briefing on COVID-19-11 March 2020. Available at: <https://www.who.int/director-general/speeches/detail/who-director-general-s-opening-remarks-at-the-media-briefing-on-covid-19---11-march-2020>. (Accessed: 1st October 2020)
- WHO I Novel Coronavirus – China. Available at: <https://www.who.int/csr/don/12-january-2020-novel-coronavirus-china/en/>. (Accessed 1 October 2020)
- Williamson EJ, Walker AJ, Bhaskaran K, *et al.* Factors associated with COVID-19-related death using OpenSAFELY. Nature. 2020 Aug;584(7821):430-436. doi: 10.1038/s41586-020-2521-4.
- Zhu N, Zhang D, Wang W, *et al.* A Novel Coronavirus from Patients with Pneumonia in China, 2019. N Engl J Med. 2020 Feb 20;382(8):727-733. doi: 10.1056/NEJMoa2001017.
- Zuckerman JN (2000) The importance of injecting vaccines into muscle. Different patients need different needle sizes. BMJ 321(7271): 1237-8.