

Main COVID-19 vaccines

Vaccine	Manufacturing	_	How it works	Method of	Eligibility
	name	combined		administration	
Nirmatrelvir and Ritonavir	Paxlovid	Combined	Nirmarelvir stops virus from growing and spreading.	Tablets	Within 5 days of
			Ritonavir helps nirmatrelvir from being broken to perform function (National Health Service, 2023),		starting symptoms.
			In depth explanation:		18 years
			Paxlovid is a protease enzyme inhibitor. It prevents the virus making copies within the human cell (viral replication).		and above.
					Amongst
			Normally, proteases cleave sites in the viral polyprotein where a		the highest
			chemical called pyrrolidone is replaced by an amino acid glutamine. This helps divide the viral polyprotein.		risk group.
					Nirmatrelvir
			Nirmatrelvir binds directly to the amino acid cysteine (Cys145)		is
			present in the catalytic binary site in M pro protease 306. This site		prescribed
			also contains the amino acid histidine.		at the first symptom,
			The binding of Nirmatrelvir to cysteine forms sulfides. Nirmatrelvir		and patients
			provides a g-lactam ring that uses glutamine amino acid residue in		do not
			the multiprotein site (Marzi et al., 2022).		require to
					be
			Ritonavir decreases metabolism of nirmatrelvir to maintain		hospitalized
			concentrations of Nirmatrelvir in the plasma via the CYP3A		
			cytochrome enzymes (liver enzymes) to inactivate the virus (Marzi <i>et al.</i> , 2022).		

			Ritonavir is also a HIV-1 protease inhibitor. This helps prevent resistance against HIV retrovirus.		
Sotrovimab	(Xevudy)	Single	A monoclonal antibody which is a type of synthetic protein that function similar to human antibodies. The aim is to stick the antibody to spike protein of virus and neutralise it to avoid it entering the lungs which may lead to infection (National Health Service, 2023) <i>In depth explanation.</i>	It is given intravenously via drip commonly the arm at the hospital or local medical centre for over	This is offered if Paxlovid does not work nor suitable.
			It is produced in Chinese Hamster Ovary (CHO) cells by recombinant DNA technology.	30 minutes. One dose required.	reinfection from COVID-19
			The drug targets the spike protein that the virus uses to attach to cells. This is especially at amino acid in the spike protein position 337, 340 and 356. This blocks the virus from entering, replication and producing new viruses in the body. This helps to overcome infection (European Medicines Agency, n.d.)	Dose: 500 mg	Patients aged 12 and above and weighing at least 40 kg
Molnupiravir	Lagevrio	Single	It stops COVID-19 from growing and spreading and prevent more severe symptoms (National Health Service, 2023) In depth explanation: Molnupiravir is a small molecular prodrug of the circulating metabolite and nucleoside derivative <i>N</i> -hydroxycytidine (NHC).	Taken orally Capsule Twice a day for 5 days.	To start within five days of symptoms starting. Highest risk
			NHC is commonly removed from the body via the kidneys by metabolising pyrimidines: cytidine and uridine. This mixes with nucleoiside.		Must have attempted with

NHC is taken up inside cells and phosphorylated (addition of	Paxlovid or
phosphate) to form NHC-triphosphate (NHC-TP).	Remdesivir
	before
NHC-TP competes with other to join and	offered this
attacks the enzyme RNA viral RNA polymerase. This helps prevent	treatment.
replication of SARS-CoV-2 (Marzi et al., 2022; Maas et al., 2024)	
Molnupiravir then enters the elongating RNA strand. It can substitute for cytidine or uridine. It can then complementarily pair with guanosine if cytidine or uridine if adenosine in the RNA template.	
Instead of terminating, Molnupiravir elongates/lengthens and produces more RNA templates that contain Molnupiravir. The RNA polymerase then uses the Molnupiravir-containing RNA templates for replication.	
This increases the number of errors across viral genome in RNA synthesis, it can negatively affect viral replication of and non-infectious viruses. This is called viral error induction (Maas <i>et al.</i> , 2024).	
More errors observed at 800 mg dose which is why it is chosen dose (Maas <i>et al.</i> 2024)	