COVID-19 Vaccine Summary Chart



Find the following information in this quick reference for pharmacy:

- Quick links and guidance
- Dosing and administration
- Storage

- Dose preparation
- Efficacy and safety information

- Clinical considerations
- Special populations
- Ingredients

Quick Links

- CDC: <u>Frequently Asked Questions about COVID-19</u> <u>Vaccination</u>
- CDC: <u>Understanding and Explaining Viral Vector</u> <u>COVID-19 Vaccines</u>
- FDA: <u>COVID-19 Vaccines</u>

- CDC: <u>V-safe After Vaccination Health Checker</u>
- CDC: <u>VaxTextsm COVID-19 Vaccination Second-Dose</u> <u>Reminder</u>
- USP: <u>COVID-19 Vaccine Handling: Operational</u> <u>Considerations for Healthcare Practitioners</u>

Vaccine	Pfizer-BioNTech (BNT162b2)	Moderna (mRNA-1273)	Janssen (Ad26.CoV2.S)
EUA	Issued December 11, 2020	Issued December 18, 2020	Issued February 27, 2021
Fact sheet	 <u>Health care providers</u> <u>Recipients/caregivers</u> 	 <u>Health care providers</u> <u>Recipients/caregivers</u> 	 <u>Health care providers</u> <u>Recipients/caregivers</u>
ACIP	Interim recommendation for use: Persons aged ≥ 16 years for prevention of COVID-19	Interim recommendation for use: Persons aged ≥ 18 years for prevention of COVID-19	Interim recommendation for use: Persons aged ≥ 18 years for prevention of COVID-19
CDC resources	Pfizer-BioNTech COVID-19 Vaccine	Moderna COVID-19 Vaccine	Janssen COVID-19 Vaccine
CDC clinical considerations		Interim Clinical Considerations	·



Vaccine	Pfizer-BioNTech (BNT162b2)	Moderna (mRNA-1273)	Janssen (Ad26.CoV2.S)	
Dosing and Administration				
Vaccine type	mRNA		Viral Vector	
Administer	Intramuscular (I.M.)			
Dose	30 mcg (0.3 mL each)	100 mcg (0.5 mL each)	5x10 ¹⁰ viral particles (0.5 mL each)	
Doses per vial	6	10	5	
Schedule	Two-dose series	Two-dose series	Single dose	
Recommended interval	21 days from first dose	28 days from first dose	N/A	
Earliest interval	17 days from first dose	24 days from first dose	N/A	
Latest interval	42 days from first dose		N/A	



Vaccine	Pfizer-BioNTech (BNT162b2)	Moderna (mRNA-1273)	Janssen (Ad26.CoV2.S)
Storage*	1		
How product arrives	Frozen liquid. No preservative.		Liquid suspension. No preservative.
Long-term storage	Ultra-low freezing until expiry date OR store frozen for up to 2 weeks	Store frozen until expiry date	Refrigerate until expiry date
Thawing	Thaw in refrigerator for at least 2-3 hours or at room temperature; must be at room temperature for at least 30 mins before dilution; do NOT refreeze	Thaw in refrigerator for at least 2-3 hours or at room temperature; must be at room temperature for at least 30 mins before administration; do NOT refreeze	Product is stored frozen by manufacturer until shipped at refrigerated temperatures; If vaccine is still frozen upon receipt, thaw at refrigerated temperature or if immediate use is required, thaw at room temperature; do NOT refreeze
Max time refrigerated unpunctured	5 days	30 days	Until expiry date
Max time at room temperature unpunctured	2 hours	12 hours	12 hours

*Temperature Key:

- Ultra-low Frozen Temperature: -80°C to -60°C (-112°F to 76°F)
- Frozen Temperature: -25°C to -15°C (-13°F to 5°F)
- Refrigerated Temperature: 2°C to 8°C (36°F to 46°F)
- Room Temperature: 9°C to 25°C (47°F to 77°F)



Vaccine	Pfizer-BioNTech (BNT162b2)	Moderna (mRNA-1273)	Janssen (Ad26.CoV2.S)	
Dose Preparation				
Dilution	Dilute with 1.8 mL of 0.9% sodium chloride (normal saline, preservative free).	Not diluted.		
Coloring	Off-white s	suspension	Colorless to slightly yellow, clear very opalescent suspension	
Handling	Do NOT shake; invert only	Do NOT shake; swirl before drawing up dose		
Max time refrigerated after first punctured	6 hours after dilution	6 hours	6 hours	
Max time at room temperature after first punctured	6 hours after dilution	6 hours	2 hours	
Efficacy and Safety Infor	mation			
Publications	Dagan, et al. NEJM. Feb 24, 2021	Baden, et al. NEJM. Feb 4, 2021	Sadoff, et al. NEJM. Jan 13, 2021	
	Polack, et al. <i>NEJM</i> . Dec 31, 2020	Anderson, et al. NEJM. Dec 17, 2020		
	Walsh, et al. NEJM. Dec 17, 2020	Jackson, et al. <i>NEJM</i> . Nov 12, 2020		
Overall efficacy;	95% beginning 7 days after second	94% beginning 14 days after second	67% beginning 14 days after single	
infection	trial data in 43,538 volunteers	trial data in >30,000 volunteers	trial data in >40,000 volunteers	
Prevention of severe COVID-19 infection	89%	100%	85%	
Prevention of asymptomatic COVID-19 infection	Under evaluation	Limited data suggest some degree of prevention	Data suggest a 60% reduction in asymptomatic infection from 29 days after dose	



Vaccine	Pfizer-BioNTech (BNT162b2)	Moderna (mRNA-1273)	Janssen (Ad26.CoV2.S)
Efficacy and Safety Information (continued)			
Study demographics	Diversity of volunteers: 81.9% White; 26.2% Hispanic/Latino; 9.8% African American; 4.4% Asian; <3% other races/ethnicities	Diversity of volunteers: 79.4% White; 20% Hispanic/Latino; 9.7% African American; 4.7% Asian; <3% other races/ethnicities	Diversity of volunteers: 59% White; 45% Hispanic/Latino ; 19% African American; 3% Asian ; 9% Native American
	Age and sex distribution: 50.6% male; 49.4% female; 21.4% 65 years and older	Age and sex distribution: 52.6% male; 47.4% female; 25.3% 65 years and older	Age and sex distribution: 55% male; 45% female; 34% 60 years and older
Postvaccination symptoms	 Injection site: pain, swelling, erythema at injection site, localized axillary lymphadenopathy (80%–89% of vaccinated persons*) Systemic: fever, fatigue, headache, chills, myalgia, arthralgia (55%–83% of vaccinated persons*; acetaminophen or ibuprofen may be used) These symptoms tend to be more common after the second dose and resolve 1–3 days after vaccination Anaphylaxis following vaccination is noted in US postmarket surveillance at a rate of 4.7 cases/million for Pfizer-BioNTech and at a rate of 2.5 cases/million for Moderna as of 1/18/21; unless contraindicated, benefit of vaccination outweighs risk of anaphylaxis; refer to CDC's guidance on Managing Anaphylaxis 		Injection site: pain, swelling, erythema
			 Systemic: headache, fatigue, muscle ache, nausea, fever Severe allergic reactions
			including anaphylaxis, were reported in clinical studies
			 Access a comprehensive summary of local reactions, systemic reactions, adverse events, and serious adverse events for the <u>Janssen</u> COVID-19 vaccine
	 Access a comprehensive summar reactions, adverse events, and se or <u>Moderna</u> COVID-19 vaccines 	rious adverse events for the <u>Pfizer</u>	
	* Depending on the vaccine, age gro	pup, and vaccine dose	



Vaccine	Pfizer-BioNTech (BNT162b2)	Moderna (mRNA-1273)	Janssen (Ad26.CoV2.S)			
Efficacy and Safety Info	Efficacy and Safety Information (continued)					
Contraindications	traindications • Severe allergic reaction (e.g., anaphylaxis) to any component of the vaccine					
	 Persons with a contraindication to mRNA COVID-19 vaccines (including due to a known allergy to polyethylene glycol [PEG]) have a precaution to Janssen COVID-19 vaccine, and vice versa 					
 Persons with a contraindication to Janssen COVID-19 vaccine (including due to a known polysorbate) have a precaution to mRNA COVID-19 vaccines 						
	revious dose or known (diagnosed)					
	• Persons with contraindication to one mRNA vaccine should not receive doses of either mRNA vaccine (Pfizer-BioNTech or Moderna)					
• If screen positive for a contraindication, do not vaccinate and consider referral to alle			er referral to allergist-immunologist			
Precautions	• Among persons without a contraindication, a history of any immediate (within 4 hours) allergic re to other vaccines or injectable therapies					
	 Persons with a contraindication to mRNA COVID-19 vaccines (Pfizer-BioNTech or Moderna) have a precaution to Janssen COVID-19 vaccine, and vice versa 					
	If screen positive for a precaution, complete a risk assessment, consider referral to allergist- immunologist, and observe for 30 minutes postvaccination					



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Clinical Considerations				
Interchangeability of COVID-19 vaccines	COVID-19 vaccines are not interchangeable; if the first dose of an mRNA COVID-19 vaccine was received, but the patient is unable to complete the series (e.g., contraindication), then the Janssen COVID-19 vaccine may be given at a minimum interval of 28 days from mRNA dose and the patient is considered to have received a valid, single-dose Janssen vaccination not a mixed vaccination series.			
Coadministration with other vaccines	Administer alone; separate COVID-19 vaccination a minimum of 14 days from other vaccines, unless the benefits of vaccination outweigh the risks of waiting to vaccinate			
Coadministration with antipyretic/analgesic	Prophylactic administration of antipyretic or analgesic medications for the prevention of postvaccination symptoms is NOT recommended; these medications <i>may be used if postvaccination symptoms occur, and patient need exists</i>			
Persons with a history of SARS-CoV-2 infection	Vaccination should be offered regardless of prior SARS-CoV-2 infection; while vaccine supplies remain limited, persons with a history of infection may choose to delay vaccination, if desired			
Persons treated with antibodies	Persons who received antibody therapy for COVID-19 should defer vaccination for 90 days			
Special Populations				
Immunocompromised persons	May be vaccinated; safety and efficacy data limited; counsel on the potential for a reduced immune response to the vaccine (efficacy) and the need to follow current guidance to protect themselves against COVID-19 (e.g., masks, social distancing)			
Persons with autoimmune disorder	May be vaccinated; no safety and efficacy data available, but persons with autoimmune disorders were included in clinical trials			
Pregnant/lactating women	May be vaccinated; pregnant or breastfeeding women were not included in the clinical trials; postauthorization <u>safety monitoring</u> of >30,000 women has not revealed a safety problem; mRNA and viral vector COVID-19 vaccines are not considered live virus vaccines and are not considered a risk to the breastfeeding infant			
Children and adolescents	Adolescents aged 16 to 17 years are eligible for vaccination	Not recommended to persons ≤18 years of age	Not recommended to persons ≤18 years of age	
Other populations	Persons with a history of Guillain-Barre syndrome or Bell's palsy may be vaccinated; persons with a history of dermal filler use may experience temporary swelling at or near the site of filler injection following vaccination and should follow up with their health care provider if this occurs			



Vaccine	Pfizer-BioNTech (BNT162b2)	Moderna (mRNA-1273)	Janssen (Ad26.CoV2.S)		
Ingredients	Ingredients				
	Nucleoside-modified mRNA encoding the viral spike (S) glycoprotein of SARS-CoV-2	 Nucleoside-modified mRNA encoding the viral spike (S) glycoprotein of SARS-CoV-2 	 5×10¹⁰ virus particles Citric acid 		
	• 2[(polyethylene glycol)*-2000]- N,N-ditetradecylacetamide	 Polyethylene glycol (PEG)* 2000 dimyristoyl glycerol 	Trisodium citrateEthanol		
	 1,2-distearoyl-sn-glycero-3- phosphocholine Cholesterol 	 (DMG) 1,2-distearoyl-sn-glycero-3- phosphocholine Cholesterol 	 2-hydroxypropyl-β-cyclodextrin Polysorbate-80* Sodium chloride 		
	• (4-hydroxybutyi)azanediyi) bis(hexane-6,1-diyl)bis(2- hexyldecanoate)	 SM-102 (proprietary to Moderna) 			
	Potassium chloride	Tromethamine			
	 Monobasic potassium phosphate 	Tromethamine hydrochlorideAcetic acid			
	Sodium chloride	Sodium acetate			
	 Dibasic sodium phosphate dihydrate 	Sucrose			
	Sucrose				

*As of March 1, 2021, mRNA COVID-19 vaccines are the only vaccines in the United States that contain PEG, though several vaccines contain polysorbate (more information can be found in CDC's vaccine excipient summary).

Disclaimer: Information related to the COVID-19 pandemic is changing rapidly and continuously. The material and information contained in this publication is believed to be current as of the date included on this document. The American Pharmacists Association assumes no responsibility for the accuracy, timeliness, errors or omission contained herein. Links to any sources do not constitute any endorsement of, validity, or warranty of the information contained on any site. The user of these materials should not under any circumstances solely rely on, or act based on this publication. Pharmacy professionals retain the responsibility for using their own professional judgment and practicing in accordance with all rules, regulations, and laws governing the pharmacy practice within their jurisdiction.

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