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	1		
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			
How product arrives	Frozen liquid. No preservative.		Liquid suspension. No preservative.
Long-term storage	Ultra-low freezing between -80°C to -60°C (-112°F to 76°F) until expiry date OR freeze at -25°C to -15°C (-13°F to 5°F) for up to 2 weeks	Freeze at -25°C to -15°C (-13°F to 5°F) until expiry date	Refrigerate at 2°C to 8°C (36°F to 46°F) for up to 3 months or 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	Thaw at room temperature for 1-2 hours if frozen upon receipt and immediate use is required. Product is stored frozen by manufacturer until shipped at refrigerated temperatures; refrigerate to thaw if immediate use is not required; do NOT refreeze
Max time refrigerated unpunctured	5 days	30 days	3 months
Max time at room temperature unpunctured	2 hours	12 hours	12 hours



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	suspension Colorless to slightly yellow, overy opalescent suspension	
Handling	Do NOT shake; invert only	Do NOT shake; swirl b	efore 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. <i>NEJM</i> . Feb 24, 2021 <u>Polack, et al. <i>NEJM</i>. Dec 31, 2020</u> Walsh, et al. <i>NEJM</i> . Dec 17, 2020	Baden, et al. <i>NEJM</i> . Feb 4, 2021 Anderson, et al. <i>NEJM</i> . Dec 17, 2020 Jackson, et al. <i>NEJM</i> . Nov 12, 2020	Sadoff, et al. NEJM. Jan 13, 2021
Overall efficacy; prevention of COVID-19 infection	95% beginning 7 days after second dose: primary analysis of Phase III trial data in 43,538 volunteers	94% beginning 14 days after second dose: <u>primary analysis</u> of Phase III trial data in >30,000 volunteers	67% beginning 14 days after single dose: <u>primary analysis</u> of Phase III 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 Info	ormation (continued)		
Study demographics	Diversity of volunteers: 81.9% White; 26.2% Hispanic/Latino; 9.8% African American; 4.4% Asian; <3% other races/ethnicities Age and sex distribution: 50.6% male; 49.4% female; 21.4% 65 years and older	Diversity of volunteers: 79.4% White; 20% Hispanic/Latino; 9.7% African American; 4.7% Asian; <3% other races/ethnicities Age and sex distribution: 52.6% male; 47.4% female; 25.3% 65 years and older	Diversity of volunteers: 59% White; 45% Hispanic/Latino ; 19% African American; 3% Asian ; 9% Native American Age and sex distribution: 55% male; 45% female; 34% 60 years and older
Postvaccination symptoms			• Injection site: pain, swelling, erythema
	• Systemic: fever, fatigue, headache, chills, myalgia, arthralgia (55%–83% of vaccinated persons*; acetaminophen or ibuprofen may be used)		 Systemic: headache, fatigue, muscle ache, nausea, fever Severe allergic reactions, including anaphylaxis, were reported in clinical studies
	• 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 <u>postmarket</u> <u>surveillance</u> 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 <u>Managing Anaphylaxis</u>		 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 summary of local reactions, systemic reactions, adverse events, and serious adverse events for the <u>Pfizer</u> or <u>Moderna</u> COVID-19 vaccines		
	* Depending on the vaccine, age gr	oup, and vaccine dose	



Vaccine	Pfizer-BioNTech (BNT162b2)	Moderna (mRNA-1273)	Janssen (Ad26.CoV2.S)			
Efficacy and Safety Info	Efficacy and Safety Information (continued)					
Contraindications	• Severe allergic reaction (e.g., ana	phylaxis) to any component of the va	ccine			
	 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 					
		n to Janssen COVID-19 vaccine (inclu n to mRNA COVID-19 vaccines	iding due to a known allergy to			
	• Immediate (within 4 hours) allergic reaction of any severity after a previous dose or known (diagnosed) allergy to a component of the vaccine (see ingredients below)					
	• Persons with contraindication to (Pfizer-BioNTech or Moderna)	one mRNA vaccine should not receive	e doses of either mRNA vaccine			
If screen positive for a contraindication, do not vaccinate and consider referral to allergist-in			referral to allergist-immunologist			
Precautions	• Among persons without a contraindication, a history of any immediate (within 4 hours) allergic reaction 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 of vaccination outweigh the risks of v	vaccination a minimum of 14 days fror vaiting to vaccinate	m other vaccines, unless the benefits	
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			
	 Nucleoside-modified mRNA encoding the viral spike (S) glycoprotein of SARS-CoV-2 2[(polyethylene glycol)*-2000]- N,N-ditetradecylacetamide 1,2-distearoyl-sn-glycero-3- phosphocholine Cholesterol (4-hydroxybutyl)azanediyl) bis(hexane-6,1-diyl)bis(2- hexyldecanoate) Potassium chloride Monobasic potassium phosphate Sodium chloride Dibasic sodium phosphate dihydrate 	 Nucleoside-modified mRNA encoding the viral spike (S) glycoprotein of SARS-CoV-2 Polyethylene glycol (PEG)* 2000 dimyristoyl glycerol (DMG) 1,2-distearoyl-sn-glycero-3- phosphocholine Cholesterol SM-102 (proprietary to Moderna) Tromethamine Tromethamine hydrochloride Acetic acid Sodium acetate Sucrose 	 5x10¹⁰ virus particles Citric acid Trisodium citrate Ethanol 2-hydroxypropyl-β-cyclodextrin Polysorbate-80* Sodium chloride

*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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