#### 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>VaxText<sup>sm</sup> 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	<ul> <li><u>Health care providers</u></li> <li><u>Recipients/caregivers</u></li> </ul>	<ul> <li><u>Health care providers</u></li> <li><u>Recipients/caregivers</u></li> </ul>	<ul> <li><u>Health care providers</u></li> <li><u>Recipients/caregivers</u></li> </ul>
ACIP	Interim recommendation for use: Persons aged ≥ <b>16 years</b> for prevention of COVID-19	Interim recommendation for use: Persons aged ≥ <b>18 years</b> for prevention of COVID-19	Interim recommendation for use: Persons aged ≥ <b>18 years</b> 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	Dosing and Administration				
Vaccine type	mF	RNA	Viral Vector		
Administer	Intramuscular (I.M.)				
Dose	30 mcg ( <b>0.3 mL each</b> )	100 mcg ( <b>0.5 mL each</b> )	5x10 <sup>10</sup> viral particles ( <b>0.5 mL each</b> )		
Doses per vial	6	10-11 dose vial or 13-15 dose vial	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		
Administration Errors	Refer to CDC's <u>COVID-19 Vaccine Administration Errors of Deviations</u> guide for information about how to handle these situations.				



Vaccine	Pfizer-BioNTech (BNT162b2)	Moderna (mRNA-1273)	Janssen (Ad26.CoV2.S)
Storage*			
How product arrives	Frozen liquid. N	No preservative.	Liquid suspension. No preservative.
Long-term storage	Ultra-low freezing until expiry date OR store frozen between -25°C to -15°C (-13°F to 5°F) for up to 2 weeks	Store frozen between -50°C to -15°C (-58°F to 5°F) 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	24 hours	12 hours

\*Temperature Key:

- Ultra-low Frozen Temperature: -80°C to -60°C (-112°F to 76°F)
- Pfizer-BioNTech Frozen Temperature: -25°C to -15°C (-13°F to 5°F)
- Moderna Frozen Temperature: -50°C to -15°C (-58°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 c	liluted.
Coloring	Off-white	suspension	Colorless to slightly yellow, clear very opalescent suspension
Handling	Do NOT shake; invert only	Do NOT shake; <b>swirl b</b>	efore drawing up dose
Max time refrigerated after first punctured	6 hours after dilution	12 hours	6 hours
Max time at room temperature after first punctured	6 hours after dilution	12 hours	2 hours
Efficacy and Safety Infor	mation		
Publications	Dagan, et al. <i>NEJM</i> . Feb 24, 2021 Polack, et al. <i>NEJM</i> . Dec 31, 2020	Baden, et al. <i>NEJM</i> . Feb 4, 2021 Anderson, et al. <i>NEJM</i> . Dec 17, 2020	Sadoff, et al. NEJM. Jan 13, 2021
Overall efficacy; prevention of COVID-19 infection	Walsh, et al. NEJM. Dec 17, 2020 <b>95%</b> beginning 7 days after second dose: primary analysis of Phase III trial data in 43,538 volunteers	Jackson, et al. NEJM. Nov 12, 2020 <b>94%</b> beginning 14 days after second dose: primary analysis of Phase III trial data in >30,000 volunteers	<b>67%</b> 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	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	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
	and older	and older	older
Postvaccination symptoms	• Injection site: pain, swelling, erythema at injection site, localized axillary lymphadenopathy (80%-89% of vaccinated persons*)		• Injection site: pain, swelling, erythema
	• <b>Systemic:</b> fever, fatigue, headache, chills, myalgia, arthralgia (55%–83% of vaccinated persons*; acetaminophen or ibuprofen		<ul> <li>Systemic: headache, fatigue, muscle ache, nausea, fever</li> <li>Severe allergic reactions, including anaphylaxis, were reported in clinical studies</li> </ul>
	<ul> <li>may be used)</li> <li>These symptoms tend to be more common after the second dose and resolve 1-3 days after vaccination</li> </ul>		
<ul> <li>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></li> </ul>		:/million for Pfizer-BioNTech for Moderna as of 1/18/21; f vaccination outweighs risk of	• Access a comprehensive summary of local reactions, systemic reactions, adverse events, and serious adverse events for the <u>Janssen</u>
	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		COVID-19 vaccine
	* 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	ohylaxis) to any component of the va	ccine			
		n to mRNA COVID-19 vaccines (inclu ve a precaution to Janssen COVID-19				
	<ul> <li>Persons with a contraindication to Janssen COVID-19 vaccine (including due to a known aller polysorbate) have a precaution to mRNA COVID-19 vaccines</li> </ul>					
<ul> <li>Immediate (within 4 hours) allergic reaction of any severity after a previous dose or know allergy to a component of the vaccine (see ingredients below)</li> </ul>			vious dose or known (diagnosed)			
<ul> <li>Persons with contraindication to one mRNA vaccine should not receive doses of either (Pfizer-BioNTech or Moderna)</li> </ul>						
• If screen positive for a contraindication, do not vaccinate and consider referral to allergist-imm			referral to allergist-immunologist			
Precautions• Among persons without a contraindication, a history of any immediate (within 4 hours) to other vaccines or injectable therapies						
	<ul> <li>Persons with a contraindication to mRNA COVID-19 vaccines (Pfizer-BioNTech or Moderna) have a precaution to Janssen COVID-19 vaccine, and vice versa</li> </ul>					
	If screen positive for a precaution, complete a risk assessment, consider referral to allergist- immunologist, and observe for 30 minutes postvaccination					



Vaccine	Pfizer-BioNTech (BNT162b2)	Moderna (mRNA-1273)	Janssen (Ad26.CoV2.S)	
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			
	<ul> <li>Nucleoside-modified mRNA encoding the viral spike (S) glycoprotein of SARS-CoV-2</li> <li>2[(polyethylene glycol)*-2000]- N,N-ditetradecylacetamide</li> <li>1,2-distearoyl-sn-glycero-3- phosphocholine</li> <li>Cholesterol</li> <li>(4-hydroxybutyl)azanediyl) bis(hexane-6,1-diyl)bis(2- hexyldecanoate)</li> <li>Potassium chloride</li> <li>Monobasic potassium phosphate</li> <li>Sodium chloride</li> <li>Dibasic sodium phosphate dihydrate</li> </ul>	<ul> <li>Nucleoside-modified mRNA encoding the viral spike (S) glycoprotein of SARS-CoV-2</li> <li>Polyethylene glycol (PEG)* 2000 dimyristoyl glycerol (DMG)</li> <li>1,2-distearoyl-sn-glycero-3- phosphocholine</li> <li>Cholesterol</li> <li>SM-102 (proprietary to Moderna)</li> <li>Tromethamine</li> <li>Tromethamine hydrochloride</li> <li>Acetic acid</li> <li>Sodium acetate</li> <li>Sucrose</li> </ul>	<ul> <li>5×10<sup>10</sup> virus particles</li> <li>Citric acid</li> <li>Trisodium citrate</li> <li>Ethanol</li> <li>2-hydroxypropyl-β-cyclodextrin</li> <li>Polysorbate-80*</li> <li>Sodium chloride</li> </ul>

\*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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