

COVID-19 vaccine immunization schedule

Updated September 2022



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Supplier	Recipient age	Immunization Schedule					
		Primary series dose	No. of doses in primary seriesBitInterval between doses	Booster dose	Booster interval since last dose	Product	Preparation instructions BUD
Moderna	6 mo – 5 y	25 mcg (0.25 mL)	2 ^a or 3 ^b • D1 to D2: ≥ 4 ^b -8 ^c wks • D2 to D3: ≥ 4 wks	Not authorized	NA	 Blue vial cap with magenta border (NDC: 80777-279-99) Primary series only (monovalent) 25 mcg/0.25 mL 	 Do not dilute Discard 12 h after first puncture
	6 – 11 y	50 mcg (0.5 mL)	2ª or 3 ^b	Not authorized	NA	Dark blue vial cap with purple border (NDC: 80777-277-99)	Do not dilute
			 D1 to D2: ≥ 4^b-8^c wks D2 to D3: ≥ 4 wks 			 Primary series only (monovalent). Labeling still states for, "booster doses only." This product used to be used as a booster dose in adults and for the primary series in patients 6-11 y old – use in this pediatric population is now the only approved use for this product following EUA revisions and EUA for bivalent booster in adults. 50 mcg/0.5 mL 	 Discard 12 h after first puncture
	12-17 у	100 mcg (0.5 mL)	2 ^a or 3 ^b • D1 to D2: ≥ 4 ^b -8 ^c wks • D2 to D3: ≥ 4 wks	Not authorized	NA	 Red vial cap with light blue border (NDC: 80777-273-99, 80777-273-98) Primary series only (monovalent) 100 mcg/0.5 mL 	 Do not dilute Discard 12 h after first puncture
	≥ 18 y	100 mcg (0.5 mL)	2ª or 3 ^b	25 mcg original, 25 mcg	Last dose*: ≥ 2 mos *Last dose includes either the final dose in the primary vaccine series in patients who have <u>not</u> yet received a booster or the last monovalent booster.	 Red vial cap with light blue border (NDC: 80777-273-99, 80777-273-98, 80777-100-99^d, 80777-100-98^d) Primary series only (monovalent) 100 mcg/0.5 mL Dark blue vial cap with gray border (NDC: 80777-282-99) Booster dose only (bivalent) 50 mcg/0.5 mL 	 Do not dilute Discard 12 h after first puncture
			 D1 to D2: ≥ 4^b-8^c wks D2 to D3: ≥ 4 wks 				
Pfizer	6 mo – 4 y	3 mcg (0.2 mL)	3 ^{a,b}	Not authorized	NA	 Maroon vial cap with maroon border (NDC: 59267-0078-4) Primary series only (monovalent) 3 mcg/0.2 mL 	• Dilute with 2.2 mL sterile
			 D1 to D2: ≥ 3^b-8^c wks D2 to D3: ≥ 8 wks 				0.9% sodium chloride injectionBUD: 12 h after dilution
	5 – 11 y	10 mcg (0.2 mL)	2 ^a or 3 ^b		D2 to B1: ≥ 5 mos ^a D3 to B1: ≥ 3 mos ^b	 Orange vial cap with orange border (NDC: 59267-1055-4) Primary series and booster dose (monovalent) 10 mcg/0.2 mL 	 Dilute with 1.3 mL sterile 0.9% sodium chloride injection BUD: 12 h after dilution
			 D1 to D2: ≥ 3^b-8^c wks D2 to D3: ≥ 4 wks 				
	≥ 12 y	30 mcg (0.3 mL)	2ª or 3 ^b	Bivalent booster 15 mcg original, 15 mcg Omicron BA.4/BA.5 [30 mcg (0.3 mL) total]	Last dose*: ≥ 2 mos *Last dose includes either the final dose in the primary vaccine series in patients who have <u>not</u> yet received a booster or the last monovalent booster.	Purple vial cap with purple border (NDC: 59267-1000-3, 59267-1000-2,	Purple vial cap with purple
			 D1 to D2: ≥ 3^b-8^c wks D2 to D3: ≥ 4 wks 			 0069-1000-03^d, 0069-1000-02^d) Primary series only (monovalent) 30 mcg/0.3 mL Gray vial cap with gray border (NDC: 59267-1025-4, 59267-1025-3, 0069-2025-10^d, 0069-2025-25^d) Primary series only (monovalent) 30 mcg/0.3 mL Gray vial cap with gray border (NDC: 59267-1401-1, 59267-1404-2) Booster dose only (bivalent) 30 mcg/0.3 mL 	 border Dilute with 1.8 mL sterile 0.9% sodium chloride injection BUD: 6 h after dilution Gray vial cap with gray border (monovalent and bivalent products) Do not dilute BUD: 12 h after puncture
Novavax	≥ 12 y	5 mcg (0.5 mL)	2 ^e	Not authorized	NA	Multidose vial (NDC: 80631-100-01)	Do not dilute
			• D1 to D2: 3-8 ^f wks			 Primary series only (monovalent) 5 mcg/0.5 mL 	Discard 6 h after first puncture

Footnotes: ^a For most people; ^b For those who are moderately or severely immunocompromised; ^c An 8 wk interval may be optimal for people who are not moderately or severely immunocompromised and ages 6 mos-64 y, especially for males ages 12-39 years; ^d FDA-approved product; ^e No recommendations are available for using this vaccine in persons who are immunocompromised; ^b For those who are moderately or severely immunocompromised; ^c An 8 wk interval may be optimal for people who are not moderately or severely immunocompromised and ages 6 mos-64 y, especially for males ages 12-39 years; ^d FDA-approved product; ^e No recommendations are available for using this vaccine in persons who are immunocompromised; patients with significant immunocompromising conditions were excluded from clinical studies; ^f An 8-wk interval may be considered to reduce the risk of myocarditis, extrapolated from mRNA vaccine data . **Abbreviations**: B = booster; BUD = beyond use date; D = dose; EUA = emergency use authorization; NA = not applicable



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