

ASTRAZENECA COVID-19 VACCINE- azd1222 injection, suspension
AstraZeneca Pharmaceuticals LP

AstraZeneca COVID-19 Vaccine

Package/Label Display Panel – Vial Label

AstraZeneca COVID-19 Vaccine NDC 0310-1222-10

For use under Emergency Use Authorization

Suspension for Intramuscular Injection

After first use, discard after:

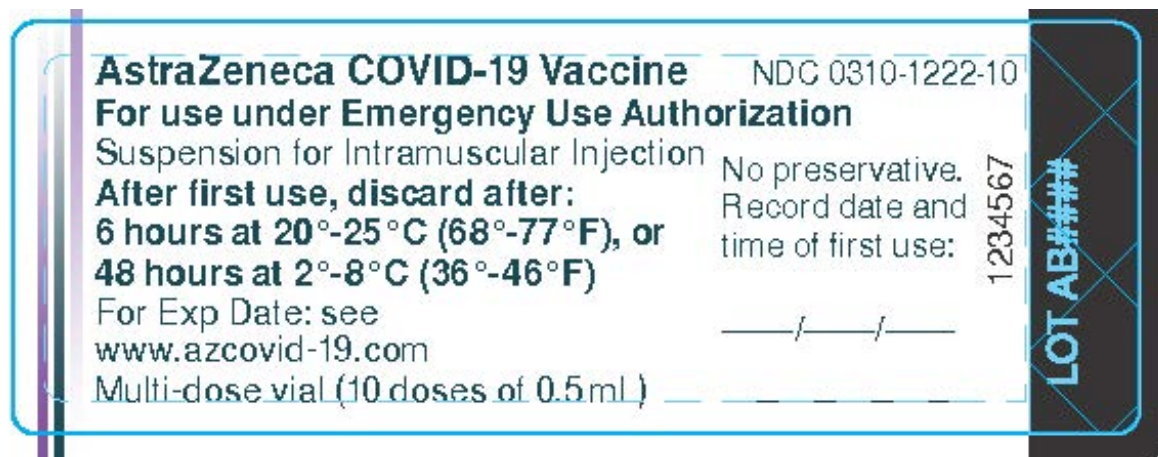
6 hours at 20°-25°C (68°-77°F), or

48 hours at 2°-8°C (36°-46°F)

For Exp Date: see

www.azcovid-19.com

Multi-dose vial (10 doses of 0.5 mL)



Package/Label Display Panel – Vial Carton

NDC 0310-1222-15

AstraZeneca COVID-19 Vaccine

For use under Emergency Use Authorization

Suspension for Intramuscular Injection

Store at 2°-8°C (36°-46°F) in original carton to protect from light.

Do not freeze or shake. **No preservative.**

Discard 6 hours after first use when held at 20°-25°C (68°-77°F).

Discard 48 hours after first use when held at 2°-8°C (36°-46°F).

10 Multi-dose vials

(each vial contains 10 doses of 0.5 mL)

AstraZeneca



Expiration Date: Please see www.azcovid-19.com

See FDA-authorized Fact Sheet for additional information.

Contents: 10 Multi-dose vials (each vial contains 10 doses of 0.5 mL). No preservative.



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ASTRAZENECA COVID-19 VACCINE

azd1222 injection, suspension

Product Information

Product Type	VACCINE	Item Code (Source)	NDC:0310-1222
Route of Administration	INTRAMUSCULAR		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
AZD-1222 (UNII: B5S3K2V0G8) (AZD-1222 - UNII:B5S3K2V0G8)	AZD-1222	50000000000 {VP} in 0.5 mL

Inactive Ingredients

Ingredient Name	Strength
HISTIDINE (UNII: 4QD397987E)	
HISTIDINE MONOHYDROCHLORIDE (UNII: 1D5Q932XM6)	

MAGNESIUM CHLORIDE (UNII: 02F3473H9O)	
POLYSORBATE 80 (UNII: 6OZP39ZG8H)	
ALCOHOL (UNII: 3K9958V90M)	
SUCROSE (UNII: C151H8M554)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	
EDETATE DISODIUM (UNII: 7FLD91C86K)	
WATER (UNII: 059QF0K00R)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:0310-1222-15	10 in 1 CARTON		
1	NDC:0310-1222-10	5 mL in 1 VIAL, MULTI-DOSE; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
Unapproved drug other		12/22/2020	

Labeler - AstraZeneca Pharmaceuticals LP (054743190)

Registrant - AstraZeneca PLC (230790719)

Revised: 12/2020

AstraZeneca Pharmaceuticals LP