Respiratory Syncytial Virus (RSV) Vaccine

Vaccine Description	 Brands and types ABRYSVO[™]: Bivalent, recombinant protein subunit AREXVY[™]: Adjuvanted, monovalent, recombinant subunit Neither vaccine contains preservatives or latex but both may have residual host cell proteins See package inserts
Dose & Route	Dose: 0.5 mL Route: IM (Use IM precautions for persons with bleeding disorders or receiving anticoagulant therapy) See package inserts
Indications	 Individuals 60 years and older for the prevention of lower respiratory tract disease caused by RSV, using shared clinical decision making
Administrative Schedule	One dose
Booster	• None
Contraindications	 History of a severe allergic reaction (e.g., anaphylaxis) to any component of ABRYSVO[™] or AREXVY[™]
Precautions	 Vaccination should be delayed for persons experiencing moderate or severe acute illness with or without fever Appropriate medical treatment and supervision must be available to manage possible anaphylactic reactions following administration Syncope (fainting) may occur in association with administration of injectable vaccines. Procedures should be in place to avoid injury from fainting. Vaccines may be less effective in immunocompromised persons, including those receiving immunosuppressive therapy
Special Considerations	 Discard vaccine if not used within 4 hours of reconstitution See package insert for reconstitution instructions for each vaccine See Storage and Handling Section
VIS: https://www.cdc.gov/vaccines/hcp/vis/vis-statements/rsv.html Additional information may be found at: www.health.mil/rsv	