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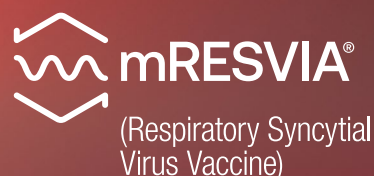
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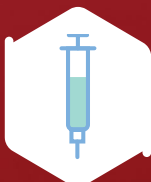
The only ACIP-recommended Ready-to-use
RSV Vaccine in a Pre-filled Syringe^{1,2}

For adults aged 60 years or older, mRESVIA is the only
RSV protection in a ready-to-use, pre-filled syringe.^{1,3,4}

mRESVIA is ready to use once thawed to room temperature.^{1*}



mRESVIA Provides Exceptional Convenience¹



Pre-filled syringe presentation



No reconstitution required



Flexible storage and handling options[†]

Indication

mRESVIA (Respiratory Syncytial Virus Vaccine) is a vaccine indicated for active immunization for the prevention of lower respiratory tract disease (LRTD) caused by respiratory syncytial virus (RSV) in adults 60 years of age and older.

Important Safety Information

Contraindications

Do not administer mRESVIA to individuals with a history of severe allergic reaction (e.g., anaphylaxis) to any component of mRESVIA.

Warnings and Precautions

- **Management of Acute Allergic Reactions:** Appropriate medical treatment must be immediately available to manage potential anaphylactic reactions following administration of mRESVIA.
- **Syncope:** Syncope (fainting) may occur in association with administration of injectable vaccines, including mRESVIA. Procedures should be in place to avoid injury from fainting.
- **Altered Immunocompetence:** Immunocompromised individuals, including those receiving immunosuppressive therapy, may have a diminished immune response to mRESVIA.

Adverse Reactions

In a clinical trial, the most commonly reported ($\geq 10\%$) adverse reactions were injection-site pain (55.9%), fatigue (30.8%), headache (26.7%), myalgia (25.6%), arthralgia (21.7%), axillary (underarm) swelling or tenderness (15.2%), and chills (11.6%).

To report suspected adverse reactions, contact ModernaTX, Inc. at 1-866-663-3762 or VAERS at 1-800-822-7967 or <https://vaers.hhs.gov>.

Click [here](#) for full mRESVIA Prescribing Information.

^{*}Stored frozen between -40°C to -15°C (-40°F to 5°F), with the option to thaw at refrigeration or room temperature. A carton of 1 pre-filled syringe can be thawed at refrigeration 2°C to 8°C (36°F to 46°F) for 100 minutes or at room temperature 15°C to 25°C (59°F to 77°F) for 40 minutes. A carton of 10 pre-filled syringes can be thawed at refrigeration 2°C to 8°C (36°F to 46°F) for 160 minutes or at room temperature 15°C to 25°C (59°F to 77°F) for 80 minutes.¹

[†]Please see mRESVIA Full Prescribing Information for details on how to store mRESVIA.

ACIP, Advisory Committee on Immunization Practices; RSV, respiratory syncytial virus.

References: 1. mRESVIA Prescribing Information. ModernaTX, Inc. 2. Centers for Disease Control and Prevention. CDC updates RSV vaccination recommendation for adults. Updated June 26, 2024. Accessed September 5, 2024. <https://www.cdc.gov/media/releases/2024/s-0626-vaccination-adults.html> 3. AREXVY Prescribing Information. GlaxoSmithKline Biologics SA. 4. ABRYVVO Product Information. Pfizer Inc.

For Colorado and Connecticut price disclosure, please visit <https://modernadirect.com/wac-disclosure>.

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* FFF Enterprises, Inc. aligns our shipping expectations with manufacturers' estimated shipping commitments.

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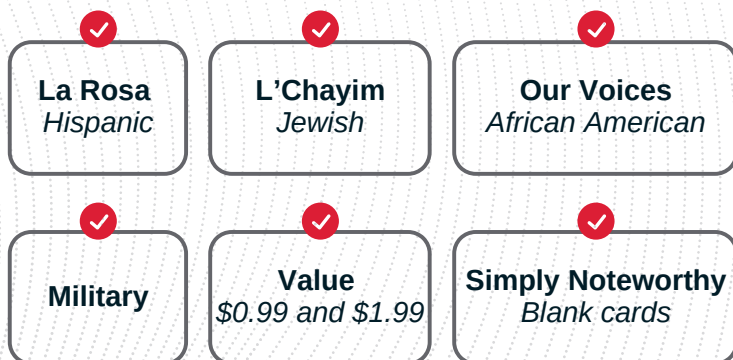
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Rabies is still present and almost always fatal if left untreated¹

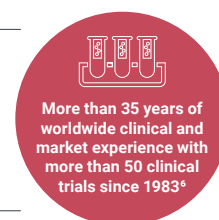
Every year 30,000 to 60,000 people come in contact with potentially rabid animals in the US, resulting in the need to administer rabies post-exposure prophylaxis to those patients²

Even though rabies is almost always fatal, it is vaccine preventable³



RabAvert® (Rabies Vaccine) is the market-leading rabies vaccine in the US⁴

RabAvert® is indicated for pre-exposure vaccination, in both primary series and booster dose, and for post-exposure prophylaxis against rabies in all age groups⁵



EFFICACY

- When administered according to the then current WHO-recommended immunization schedule for pre-exposure vaccination, 100% of subjects in clinical trials attained a protective titer.⁵
- The immunogenicity of RabAvert® was demonstrated in clinical trials conducted in different countries such as the US, the UK, Croatia, and Thailand.⁵
- When administered according to the then current WHO-recommended immunization schedule for post-exposure vaccination, >99% of subjects attained a protective titer.⁵
- Among the 203 patients in post-exposure trials followed for at least 10 months, no cases of rabies were observed.⁵
- Clinical studies of RabAvert® have shown it to be generally well-tolerated.⁵
- Anaphylaxis, meningitis, neuromuscular events such as encephalitis, transient paralysis, Guillain-Barré syndrome, myelitis, retrobulbar neuritis, and multiple sclerosis have been reported to be temporally associated with the use of RabAvert®. A patient's risk of acquiring rabies must be carefully considered before discontinuing vaccination.⁵
- In clinical trials with RabAvert®, the most common adverse reactions were injection site reactions (erythema, induration, and pain); flu-like symptoms (asthenia, fatigue, fever, headache, myalgia, and malaise); arthralgia; dizziness; lymphadenopathy; nausea; and rash.⁵

IMPORTANT SAFETY INFORMATION

- Patients considered to be at risk of a severe hypersensitivity reaction (e.g. anaphylaxis) to RabAvert® or any of its components should receive an alternative rabies vaccine if a suitable product is available. However, in view of the almost invariably fatal outcome of rabies, there is no contraindication to post-exposure prophylaxis, including pregnancy.
- RabAvert® is contraindicated for pre-exposure vaccination in anyone with a history of a severe allergic reaction (eg, anaphylaxis) to the vaccine or any of its components, which include residues of egg and chicken proteins to which some individuals may be hypersensitive.
- Review immunization history for possible vaccine sensitivity and previous vaccination-related adverse reactions, as well as medical history for emergence of clinical symptoms of anaphylaxis after exposure to egg or chicken proteins, which RabAvert® contains. Reconstituted RabAvert® also contains processed bovine gelatin and trace amounts of neomycin, chlortetracycline, and amphotericin B, to which some individuals may be hypersensitive. Appropriate medical treatment—including immediately available epinephrine injection (1:1,000), and readily available volume replacement, corticosteroids, and oxygen—must be close at hand to manage possible anaphylactic reactions following administration of RabAvert®.
- Development of active immunity after vaccination may be impaired in immune-compromised individuals. Radiation therapy, antimalarials, corticosteroids, other immunosuppressive agents, and immunosuppressive illnesses can interfere with the development of active immunity after vaccination and may diminish the protective efficacy of RabAvert®. Pre-exposure vaccination should be administered to such persons with the awareness that the immune response may be inadequate. If persons receiving corticosteroids or other immunosuppressive therapy, or who are immunosuppressed, are vaccinated post-exposure, it is important that a serum sample on Day 14 (the day of the fourth vaccination) be tested for rabies antibody to ensure that an acceptable antibody response has been induced.
- Pre-exposure vaccination should be postponed in the case of sick and convalescent persons and those considered to be in the incubation stage of an infectious disease.

Please see continued Important Safety Information throughout and enclosed full Prescribing Information.



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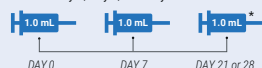


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Dosing schedule for RabAvert® (Rabies Vaccine)

Pre-exposure prophylaxis (PrEP)

According to the FDA-approved prescribing information, RabAvert® is administered intramuscularly in 3 doses on Day 0, Day 7, and Day 21 or 28.⁵



*Please note that current CDC/ACIP dosing recommendations (2022) differ from those in the prescribing information, no longer calling for a third dose in the primary series.

A **BOOSTER DOSE** may be advisable depending on the results of serological testing for the antibody titer level.^{5,7,8}

- for those with **continuous rabies exposure**, as often as every **6 months**
- for those with **frequent rabies exposure**, as often as every **2 years**

Pre-exposure vaccination does not eliminate the need for additional treatment after a known rabies exposure.⁵

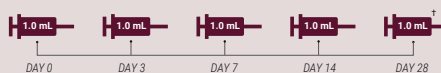
Post-exposure prophylaxis (PEP)

The essential components of rabies PEP are prompt local treatment of wounds and administration of both human rabies immune globulin (HRIG) and vaccine.⁵

PEP should begin the same day exposure occurred or as soon after exposure as possible.⁵

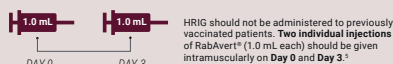
Ensure patients understand the need to promptly initiate and complete the full course of doses.

Previously unvaccinated



According to the FDA-approved prescribing information, the HRIG injection is followed by a series of **5 individual injections** of RabAvert® (1.0 mL each) given intramuscularly on **days 0, 3, 7, 14, and 28**.⁵

Previously vaccinated



HRIG should not be administered to previously vaccinated patients. **Two individual injections** of RabAvert® (1.0 mL each) should be given intramuscularly on **Day 0 and Day 3**.⁵

[†]Please note the CDC/ACIP recommendations differ, calling for a 5th dose on Day 28 only for patients who are immunocompromised.⁵

IMPORTANT SAFETY INFORMATION (CONTINUED)

- This product contains albumin, a derivative of human blood. It is present in RabAvert® at concentrations of ≤0.3 mg/dose. Based on effective donor screening and product manufacturing processes, it carries an extremely remote risk for transmission of viral diseases. A theoretical risk for transmission of Creutzfeldt-Jakob disease (CJD) also is considered extremely remote. No cases of transmission of viral diseases or CJD have ever been identified for albumin.
- Because animal reproductive studies or studies in pregnant or lactating woman have not been conducted with RabAvert®, it is not known whether RabAvert® can cause fetal harm when administered to a pregnant woman, whether it can affect reproduction capacity, or whether it is excreted in animal or human milk with consequent risk to breastfed infants (but many drugs are excreted in human milk). Use RabAvert® in pregnant or lactating women only if clearly needed.
- Pre-exposure vaccination does not eliminate the need for additional therapy after a known rabies exposure.
- For intramuscular use only. Unintentional intravascular injection may result in systemic reactions, including shock.
- Syncope (fainting) can occur in association with administration of injectable vaccines, including RabAvert®. Procedures should be in place to avoid falling injury and to restore cerebral perfusion following syncope.
- No data are available regarding the concurrent administration of RabAvert® with other vaccines.

Please see continued Important Safety Information throughout and enclosed full Prescribing Information.



Ordering information for RabAvert® (Rabies Vaccine)

There is typically a seasonal increase in rabies exposures during the months of June through September due to increased opportunity for human and animal interactions. Consider this when planning your inventory.

To learn more, visit: [Bavarian-Nordic.com](https://www.bavarian-nordic.com) or [BNVaccines.com](https://www.bnvaccines.com)

PRODUCT ORDERING INFORMATION

NDC number product description
50632-010-01 RabAvert® (Rabies Vaccine)

Package size	Case size
Single Dose Kit	140

BILLING CODE

90675 – Rabies Vaccine, for intramuscular use

SINGLE DOSE KIT INCLUDES:

- One vial of freeze-dried vaccine containing a single dose
- Disposable prefilled syringe of sterile diluent for reconstitution of RabAvert®
- Two identical needles: one for reconstitution, one for injection

IMPORTANT SAFETY INFORMATION (CONTINUED)

- Administration of Human Rabies Immune Globulin (HRIG), which along with prompt local cleaning of wounds should take place before post-exposure prophylaxis, must not exceed the recommended dose, since active immunization to the vaccine may be impaired. HRIG should not be administered to previously vaccinated persons as it may blunt their rapid memory response to rabies antigen.

Please see enclosed full Prescribing Information.

References

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To report SUSPECTED ADVERSE REACTIONS, contact Bavarian Nordic at 1-844-4BAVARIAN or the US Department of Health and Human Services by either visiting www.vaers.hhs.gov/reportevent.html or calling 1-800-822-7967.



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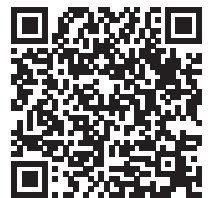
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Vaccine effectiveness may be reduced in adults 65+ as a result of a weakened immune response to vaccines.^{1,2}

Strain Mismatch

Strain mismatch, which occurred in 7 out of 10 flu seasons from 2010-2011 through 2019-2020, may further reduce vaccine effectiveness.^{3,-13}

FLUAD



Approved for Use in Adults 65 Years and Older

For active immunization against influenza disease¹⁴



ACIP Preferentially Recommended

ACIP preferentially recommends FLUAD over standard-dose influenza vaccines for adults 65+.¹⁵



Adjuvant Technology

Adding an adjuvant strengthens, broadens, and lengthens the immune response more than antigen alone.¹⁶⁻¹⁸



Robust Response & Demonstrated Safety Profile

FLUAD produced a robust immune response in clinical trials and has a demonstrated safety profile.^{14,32}



Clinically Effective

20+ years of real-world evidence (RWE) in over 59 million patients supports the clinical effectiveness of the only adjuvanted flu vaccine in the U.S. for adults 65+.^{7, 19-31}

CONTRAINDICATIONS

Do not administer FLUAD to anyone with a history of a severe allergic reaction (e.g., anaphylaxis) to any component of the vaccine, including egg protein, or to a previous influenza vaccine.

Please see the full Important Safety Information on the back of this page and the accompanying full [US Prescribing Information](#) for FLUAD.



FLUAD® (Influenza Vaccine, Adjuvanted)

INDICATION AND IMPORTANT SAFETY INFORMATION

INDICATIONS AND USAGE

FLUAD is a vaccine indicated for active immunization for the prevention of influenza disease caused by influenza virus subtypes A and type B contained in the vaccine. FLUAD is approved for use in adults 65 years of age and older.

This indication is approved under accelerated approval based on the immune response elicited by FLUAD. Continued approval for this indication may be contingent upon verification and description of clinical benefit in a confirmatory trial.

IMPORTANT SAFETY INFORMATION

CONTRAINDICATIONS

Do not administer FLUAD to anyone with a history of a severe allergic reaction (e.g., anaphylaxis) to any component of the vaccine, including egg protein, or to a previous influenza vaccine.

WARNINGS AND PRECAUTIONS

If Guillain-Barré Syndrome (GBS) has occurred within six weeks of previous influenza vaccination, the decision to give FLUAD should be based on careful consideration of the potential benefits and risks.

Appropriate medical treatment must be immediately available to manage potential anaphylactic reactions following administration of FLUAD.

Syncope (fainting) may occur in association with administration of injectable vaccines including FLUAD. Procedures should be in place to avoid injury from fainting.

The immune response to FLUAD in immunocompromised persons, including individuals receiving immunosuppressive therapy, may be lower than in immunocompetent individuals.

Vaccination with FLUAD may not protect all vaccine recipients against influenza disease.

ADVERSE REACTIONS

The most common ($\geq 10\%$) local and systemic adverse reactions in adults 65 years of age and older who received FLUAD were injection site pain (25%), injection site tenderness (21%), myalgia (15%), fatigue (13%) and headache (13%).

Other adverse events may occur.

To report SUSPECTED ADVERSE REACTIONS, contact CSL Seqirus at 1-855-358-8966 or VAERS at 1-800-822-7967 and www.vaers.hhs.gov.

Before administration, please see the full Prescribing Information for FLUAD.

References:

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USA-FLUD-25-0007

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01

Rx Returns

Serving more than 50,000 retail pharmacies nationwide, Inmar's industry leading solution increases speed of credit with minimal user effort. Onsite and offsite service options combined with RxReturns Intel™ dashboards ensure robust reporting, enhanced visibility and manufacturer policy compliance.

02

Consumer Drug Take-Back

This customer-friendly, secure medication take-back program generates publicity, community goodwill and store traffic while supporting efforts to combat prescription drug abuse and helping to protect the environment from the effects of improper drug disposal.

03

RxTransparent

Protect staff and patients from illegitimate medication distribution and safe handling standards for hazardous drugs with Drug Supply Chain Security Act (DSCSA) and USP<800> solutions.

04

Rx Reconciliation

Inmar's Rx Reconciliation Service mitigates receivables risk, improves cash flow and delivers critical business-building intelligence.

05

Healthcare Analytics

Inmar's analytics offering delivers critical business intelligence via dashboards enabling you to make better financial decisions.

06

Commercial Floor Mats

Inmar offers a broad range of customizable matting solutions built with the latest technology and materials to ensure durability.

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Consistent tangible marketing is important.

With PPRB's signature line of printed wall and desk calendars as well as additional print products, like coloring books, pocket planners, and magnets, staying connected with your current & future customers is easy.



Helping pharmacies build relationships, generate sales, and win loyal customers across North America.

Your store's information will be prominently displayed on the generous bottom flap imprint area of PPRB's wall calendars.

Include your business contact info such as your store address, phone and fax numbers, store hours and website links so your customers can connect with you in-store and on your digital platforms too.



KNOWLEDGEABLE & DEPENDABLE

Remind your customers that you are available to support their well-being by providing the best health care advice and products available to them.



VALUABLE CONTENT

Customers will refer to your calendars for recipes, health tips, day-to-day planning, organizing, and more. Each view/impression encourages them to return to your store for health and wellness inquiries and products.



STAY RELEVANT & BUILD TRUST

Foster long-term connections with current and prospective customers in your community.

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Headquartered in Bastian, V.A., InSource, Inc. is an NABP® Accredited Drug Distributor serving Independent Medical Distributors and Independent Retail Pharmacies for over three decades.

InSource operates through a centralized and automated distribution network, with a large selection of Brand and Generic injectables, vaccines, gloves, med surg, disposables, consumables, lab, equipment, and other medical supplies commonly used in the practitioner, pharmacy, occupational health, industrial health, and home health care space.



VAST SELECTION

- Med-Surg, Diagnostics, Disposables, Pharmaceuticals, Equipment, and so much more!
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VALUE-ADDED SERVICES

- Personal Account Managers
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Since its inception in 1991, InSource has been providing independent distributors and pharmacies with quality products and unparalleled service!

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Return Solutions
[go with what works][™]

The Independent's Choice for Rx Returns

If you're in need of a new returns provider, Return Solutions is one of the oldest and most trusted in the industry. With over 30 years of experience and a focus on independent pharmacies and regional chains, we put you first so you can focus on your business and your patients.



Optimize your Rx returns process and payment.



Simplified Reimbursement

Credit due through the OneCheck Select program is consolidated into a single check.



Faster Credit

Choose to receive your check within 10, 30, or 90 days after your return.

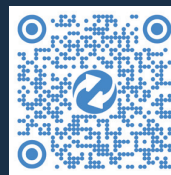


No Hidden Charges

The fee is an all-inclusive percentage of your returnable product value.

Return Solutions offers the fastest credit in the industry.

- ✓ On-Site service is completely turnkey
- ✓ Mail-In service is easy and economical
- ✓ Fastest payment as quickly as 10 days
- ✓ No additional charges



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INDEPENDENT PHARMACIES NEEDED FOR CLINICAL TRIAL RECRUITMENT

SiteLabs, with the support of local Health Systems, is partnering with independent pharmacies to qualify and enroll new patients for critical clinical trials. Your pharmacy can make a profound difference with early detection and specialized care for the patients in your community.

WHY PARTICIPATE?

- Improve community health
- Expand your pharmacy services
- Support early detection
- Compensation per patient enrolled



LOW STOCK ON COVID TEST KITS?

SiteLabs is FPN's preferred partner for COVID, Point of Care, and wellness testing supplies. Shop online at [SiteLabsglobal.com](https://www.sitelabsglobal.com).



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UPDATED CDC recommendations

All eligible adults aged **50+** are now recommended for pneumococcal vaccination^{1,2}



Actor Portrayal

- **A single dose of PCV (PCV15, PCV20, or PCV21) is recommended** for all adults aged ≥50 years and for adults aged 19 to 49 years with certain underlying conditions or risk factors* who have not received a PCV or whose vaccination history is unknown^{1,2†}

- **If PCV15 is administered**, a single dose of PPSV23[‡] should be administered ≥1 year after the PCV15 dose. A minimum interval of 8 weeks can be considered if PCV15 is used in adults with an immunocompromising condition,[§] cochlear implant, or CSF leak¹

PCV13-experienced adults who completed the recommended vaccine series

- Shared clinical decision-making is recommended regarding the use of a supplemental PCV20 or PCV21 dose for adults aged ≥65 years who have completed their recommended vaccine series with both PCV13 and PPSV23¹

PCV13-experienced adults who have not completed the recommended vaccine series

- A single dose of either PCV20 or PCV21 is recommended for adults aged ≥19 years who have started their pneumococcal vaccine series with PCV13 but have not received all recommended pneumococcal vaccine doses¹

Vaccinate all eligible adult patients for pneumococcal pneumonia and IPD today.

CDC=Centers for Disease Control and Prevention; CSF=cerebrospinal fluid; HIV=human immunodeficiency virus; IPD=invasive pneumococcal disease; PCV7=7-valent pneumococcal conjugate vaccine; PCV13=13-valent pneumococcal conjugate vaccine; PCV15=15-valent pneumococcal conjugate vaccine; PCV20=20-valent pneumococcal conjugate vaccine; PCV21=21-valent pneumococcal conjugate vaccine; PPSV23=23-valent pneumococcal polysaccharide vaccine.

*Alcoholism; chronic heart, liver, or lung disease; chronic renal failure; cigarette smoking; cochlear implant; congenital or acquired asplenia; cerebrospinal fluid leak; diabetes mellitus; generalized malignancy; HIV; Hodgkin disease; immunodeficiency; iatrogenic immunosuppression; leukemia, lymphoma, or multiple myeloma; nephrotic syndrome; solid organ transplant; sickle cell disease; or other hemoglobinopathies.¹

[†]Also applies to people who received PCV7 at any age and no other pneumococcal vaccines.¹

[‡]For adults who have received PCV15 but have not completed their recommended pneumococcal vaccine series with PPSV23, 1 dose of PCV21 or PCV20 may be used if PPSV23 is not available.¹

[§]Chronic renal failure, nephrotic syndrome, immunodeficiency, iatrogenic immunosuppression, generalized malignancy, HIV infection, Hodgkin disease, leukemia, lymphoma, multiple myeloma, solid organ transplant, congenital or acquired asplenia, or sickle cell disease or other hemoglobinopathies.¹

Learn more about the latest CDC recommendations



CDC=Centers for Disease Control and Prevention.

References: **1.** Kobayashi M. Summary of Work Group interpretation of ETR and policy options: PCV use in adults aged ≥50 years. Presented at: Advisory Committee on Immunization Practices; October 23, 2024; Atlanta, GA. Accessed October 23, 2024. <https://www.cdc.gov/acip/downloads/slides-2024-10-23-24/04-Kobayashi-Pneumococcal-508.pdf> **2.** Centers for Disease Control and Prevention. Pneumococcal vaccination. Accessed October 25, 2024. <https://www.cdc.gov/pneumococcal/vaccines/index.html>



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northeastpharmacy.com

PBA Health
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Pharmacy Owners Alliance
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rxpoa.org

PPSC
Tallahassee, FL
ppsconline.com

Quality Care Pharmacies
Macedon, NY
qcpharmacies.com

Sav-Mor Pharmacy Services
Southfield, MI
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Smart-Fill Management Group
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smart-fill.com

The Pharmacist Choice
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Doing business with our preferred vendor partners strengthens the purchasing alliance of independent pharmacies and provides increased opportunities for future negotiations. Your participation leverages our collective buying power to offer the most competitive contracts. Thank you for your continued commitment to independent pharmacy and for your patronage of our preferred vendor partners.

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