

PENICILLIN V POTASSIUM- penicillin v potassium tablet
PENICILLIN V POTASSIUM- penicillin v potassium powder, for solution
Teva Pharmaceuticals USA, Inc.

Penicillin V Potassium Tablets, USP
Penicillin V Potassium for Oral Solution, USP

Rx only

To reduce the development of drug-resistant bacteria and maintain the effectiveness of penicillin V potassium tablets, penicillin V potassium for oral solution, and other antibacterial drugs, penicillin V potassium tablets and penicillin V potassium for oral solution should be used only to treat or prevent infections that are proven or strongly suspected to be caused by bacteria.

DESCRIPTION

Penicillin V is the phenoxymethyl analog of penicillin G.

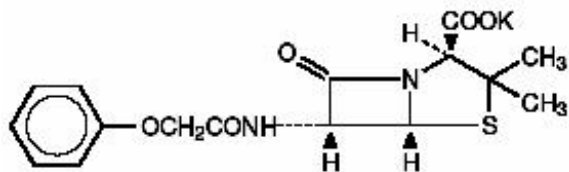
Penicillin V potassium is the potassium salt of penicillin V.

Each Penicillin V Potassium Tablet, USP contains penicillin V potassium equivalent to 250 mg (400,000 units) or 500 mg (800,000 units) penicillin V. The inactive ingredients present are dibasic calcium phosphate, magnesium stearate, microcrystalline cellulose, and sodium citrate.

Each Penicillin V Potassium Tablet USP, 250 mg contains 0.71 mEq (27.9 mg) of potassium and each Penicillin V Potassium Tablet USP, 500 mg contains 1.43 mEq (55.8 mg) of potassium.

Penicillin V Potassium for Oral Solution, USP is an off-white to pinkish colored powder, which when reconstituted as directed, yields a red colored solution with cherry flavor in which each 5 mL contains penicillin V potassium equivalent to 125 mg (200,000 units) or 250 mg (400,000 units) penicillin V. The inactive ingredients present are cherry flavor, FD&C Red #40, saccharin sodium, sodium benzoate, and sucrose.

Each 5 mL of reconstituted Penicillin V Potassium for Oral Solution USP, 125 mg (200,000 units) per 5 mL contains 0.36 mEq (13.9 mg) of potassium. Each 5 mL of reconstituted Penicillin V Potassium for Oral Solution USP, 250 mg (400,000 units) per 5 mL contains 0.71 mEq (27.9 mg) of potassium.



$C_{18}H_{17}KN_2O_5S$ MW 388.49

Monopotassium (2*S*,5*R*,6*R*)-3,3-dimethyl-7-oxo-6-(2-phenoxyacetamido)-4-thia-1-azabicyclo[3.2.0]heptane-2-carboxylate.

CLINICAL PHARMACOLOGY

Penicillin V exerts a bactericidal action against penicillin-sensitive microorganisms during the stage of active multiplication. It acts through the inhibition of biosynthesis of cell-wall mucopeptide. It is not active against the penicillinase-producing bacteria, which include many strains of staphylococci. The drug exerts high *in vitro* activity against staphylococci (except penicillinase-producing strains), streptococci (groups A, C, G, H, L, and M), and pneumococci. Other organisms sensitive *in vitro* to penicillin V are *Corynebacterium diphtheriae*, *Bacillus anthracis*, *Clostridia*, *Actinomyces bovis*, *Streptobacillus moniliformis*, *Listeria monocytogenes*, *Leptospira*, and *Neisseria gonorrhoeae*. *Treponema pallidum* is extremely sensitive.

The potassium salt of penicillin V has the distinct advantage over penicillin G in resistance to inactivation by gastric acid. It may be given with meals; however, blood levels are slightly higher when the drug is given on an empty stomach. Average blood levels are two to five times higher than the levels following the same dose of oral penicillin G and also show much less individual variation.

Once absorbed, penicillin V is about 80% bound to serum protein. Tissue levels are highest in the kidneys, with lesser amounts in the liver, skin, and intestines. Small amounts are found in all other body tissues and the cerebrospinal fluid. The drug is excreted as rapidly as it is absorbed in individuals with normal kidney function; however, recovery of the drug from the urine indicates that only about 25% of the dose given is absorbed. In neonates, young infants, and individuals with impaired kidney function, excretion is considerably delayed.

MICROBIOLOGY

Susceptibility Testing

For specific information regarding susceptibility test interpretive criteria and associated test methods and quality control standards recognized by FDA for this drug, please see: <https://www.fda.gov/STIC>.

INDICATIONS AND USAGE

Penicillin V potassium tablets and penicillin V potassium for oral solution are indicated in the treatment of mild to moderately severe infections due to penicillin G-sensitive microorganisms. Therapy should be guided by bacteriological studies (including sensitivity tests) and by clinical response.

NOTE: Severe pneumonia, empyema, bacteremia, pericarditis, meningitis, and arthritis should not be treated with penicillin V during the acute stage. Indicated surgical procedures should be performed.

The following infections will usually respond to adequate dosage of penicillin V.

Streptococcal infections (without bacteremia). Mild-to-moderate infections of the upper respiratory tract, scarlet fever, and mild erysipelas.

NOTE: Streptococci in groups A, C, G, H, L, and M are very sensitive to penicillin. Other groups, including group D (enterococcus), are resistant.

Pneumococcal infections. Mild to moderately severe infections of the respiratory tract.

Staphylococcal infections—penicillin G-sensitive. Mild infections of the skin and soft tissues.

NOTE: Reports indicate an increasing number of strains of staphylococci resistant to penicillin G, emphasizing the need for culture and sensitivity studies in treating suspected staphylococcal infections.

Fusospirochetosis (Vincent's gingivitis and pharyngitis)—Mild to moderately severe infections of the oropharynx usually respond to therapy with oral penicillin.

NOTE: Necessary dental care should be accomplished in infections involving the gum tissue.

Medical conditions in which oral penicillin therapy is indicated as prophylaxis: For the prevention of recurrence following rheumatic fever and/or chorea: Prophylaxis with oral penicillin on a continuing basis has proven effective in preventing recurrence of these conditions.

Although no controlled clinical efficacy studies have been conducted, penicillin V has been suggested by the American Heart Association and the American Dental Association for use as an oral regimen for prophylaxis against bacterial endocarditis in patients who have congenital heart disease or rheumatic or other acquired valvular heart disease when they undergo dental procedures and surgical procedures of the upper respiratory tract.¹ Oral penicillin should not be used in those patients at particularly high risk for endocarditis (e.g., those with prosthetic heart valves or surgically constructed systemic pulmonary shunts). Penicillin V should not be used as adjunctive prophylaxis for genitourinary instrumentation or surgery, lower-intestinal tract surgery, sigmoidoscopy, and childbirth. Since it may happen that *alpha* hemolytic streptococci relatively resistant to penicillin may be found when patients are receiving continuous oral penicillin for secondary prevention of rheumatic fever, prophylactic agents other than penicillin may be chosen for these patients and prescribed in addition to their continuous rheumatic fever prophylactic regimen.

NOTE: When selecting antibiotics for the prevention of bacterial endocarditis, the physician or dentist should read the full joint statement of the American Heart Association and the American Dental Association.¹

To reduce the development of drug-resistant bacteria and maintain the effectiveness of penicillin V potassium tablets, penicillin V potassium for oral solution, and other antibacterial drugs, penicillin V potassium tablets and penicillin V potassium for oral solution should be used only to treat or prevent infections that are proven or strongly suspected to be caused by susceptible bacteria. When culture and susceptibility information are available, they should be considered in selecting or modifying antibacterial therapy. In the absence of such data, local epidemiology and susceptibility patterns may contribute to the empiric selection of therapy.

CONTRAINDICATIONS

A previous hypersensitivity reaction to any penicillin is a contraindication.

WARNINGS

SERIOUS AND OCCASIONALLY FATAL HYPERSENSITIVITY (anaphylactic) REACTIONS HAVE BEEN REPORTED IN PATIENTS ON PENICILLIN THERAPY. THESE REACTIONS ARE MORE LIKELY TO OCCUR IN INDIVIDUALS WITH A HISTORY OF PENICILLIN HYPERSENSITIVITY AND/OR A HISTORY OF SENSITIVITY TO MULTIPLE ALLERGENS. THERE HAVE BEEN REPORTS OF INDIVIDUALS WITH A HISTORY OF PENICILLIN HYPERSENSITIVITY WHO HAVE EXPERIENCED SEVERE REACTIONS WHEN TREATED WITH CEPHALOSPORINS. BEFORE INITIATING THERAPY WITH PENICILLIN V POTASSIUM, CAREFUL INQUIRY SHOULD BE MADE CONCERNING PREVIOUS HYPERSENSITIVITY REACTIONS TO PENICILLINS, CEPHALOSPORINS, OR OTHER ALLERGENS. IF AN ALLERGIC REACTION OCCURS, PENICILLIN V POTASSIUM SHOULD BE DISCONTINUED AND APPROPRIATE THERAPY INSTITUTED. **SERIOUS ANAPHYLACTIC REACTIONS REQUIRE IMMEDIATE EMERGENCY TREATMENT WITH EPINEPHRINE. OXYGEN, INTRAVENOUS STEROIDS, AND AIRWAY MANAGEMENT, INCLUDING INTUBATION, SHOULD ALSO BE ADMINISTERED AS INDICATED.**

Clostridium difficile associated diarrhea (CDAD) has been reported with use of nearly all antibacterial agents, including penicillin V potassium tablets and penicillin V potassium for oral solution, and may range in severity from mild diarrhea to fatal colitis. Treatment with antibacterial agents alters the normal flora of the colon leading to overgrowth of *C. difficile*.

C. difficile produces toxins A and B which contribute to the development of CDAD. Hypertoxin producing strains of *C. difficile* cause increased morbidity and mortality, as these infections can be refractory to antimicrobial therapy and may require colectomy. CDAD must be considered in all patients who present with diarrhea following antibiotic use. Careful medical history is necessary since CDAD has been reported to occur over two months after the administration of antibacterial agents.

If CDAD is suspected or confirmed, ongoing antibiotic use not directed against *C. difficile* may need to be discontinued. Appropriate fluid and electrolyte management, protein supplementation, antibiotic treatment of *C. difficile*, and surgical evaluation should be instituted as clinically indicated.

PRECAUTIONS

General

Penicillin should be used with caution in individuals with histories of significant allergies and/or asthma.

Prescribing penicillin V potassium tablets or penicillin V potassium for oral solution in the absence of a proven or strongly suspected bacterial infection or a prophylactic indication is unlikely to provide benefit to the patient and increases the risk of the development of drug-resistant bacteria.

The oral route of administration should not be relied upon in patients with severe illness, or with nausea, vomiting, gastric dilatation, cardiospasm, or intestinal hypermotility.

Occasionally patients will not absorb therapeutic amounts of orally administered penicillin.

In streptococcal infections, therapy must be sufficient to eliminate the organism (10 day minimum); otherwise the sequelae of streptococcal disease may occur. Cultures should be taken following completion of treatment to determine whether streptococci have been eradicated.

Prolonged use of antibiotics may promote the overgrowth of nonsusceptible organisms, including fungi. Should superinfection occur, appropriate measures should be taken.

Information for Patients

Patients should be counseled that antibacterial drugs, including penicillin V potassium tablets and penicillin V potassium for oral solution, should only be used to treat bacterial infections. They do not treat viral infections (e.g., the common cold). When penicillin V potassium tablets or penicillin V potassium for oral solution are prescribed to treat a bacterial infection, patients should be told that although it is common to feel better early in the course of therapy, the medication should be taken exactly as directed. Skipping doses or not completing the full course of therapy may (1) decrease the effectiveness of the immediate treatment, and (2) increase the likelihood that bacteria will develop resistance and will not be treatable by penicillin V potassium tablets, penicillin V potassium for oral solution, or other antibacterial drugs in the future.

Diarrhea is a common problem caused by antibiotics which usually ends when the antibiotic is discontinued. Sometimes after starting treatment with antibiotics, patients can develop watery and bloody stools (with or without stomach cramps and fever) even as late as two or more months after having taken the last dose of the antibiotic. If this occurs, patients should contact their physician as soon as possible.

ADVERSE REACTIONS

Although the incidence of reactions to oral penicillins has been reported with much less frequency than following parenteral therapy, it should be remembered that all degrees of hypersensitivity, including fatal anaphylaxis, have been reported with oral penicillin.

The most common reactions to oral penicillin are nausea, vomiting, epigastric distress, diarrhea, and black hairy tongue. The hypersensitivity reactions reported are skin eruptions (maculopapular to exfoliative dermatitis), urticaria and other serum-sicknesslike reactions, laryngeal edema, and anaphylaxis.

Fever and eosinophilia may frequently be the only reaction observed. Hemolytic anemia, leukopenia, thrombocytopenia, neuropathy, and nephropathy are infrequent reactions and usually associated with high doses of parenteral penicillin.

To report SUSPECTED ADVERSE REACTIONS, contact Teva at 1-888-838-2872 or FDA at 1-800-FDA-1088 or www.fda.gov/medwatch.

DOSAGE AND ADMINISTRATION

The dosage of penicillin V potassium tablets and penicillin V potassium for oral solution should be determined according to the sensitivity of the causative microorganisms and the severity of infection, and adjusted to the clinical response of the patient.

The usual dosage recommendations for adults and children 12 years and over are as follows:

Streptococcal infections—mild to moderately severe—of the upper respiratory tract and including scarlet fever and erysipelas: 125 to 250 mg (200,000 to 400,000 units) every 6 to 8 hours for 10 days.

Pneumococcal infections—mild to moderately severe—of the respiratory tract, including otitis media: 250 to 500 mg (400,000 to 800,000 units) every 6 hours until the patient has been afebrile for at least 2 days.

Staphylococcal infections—mild infections of skin and soft tissue (culture and sensitivity tests should be performed): 250 to 500 mg (400,000 to 800,000 units) every 6 to 8 hours.

Fusospirochetosis (Vincent's infection) of the oropharynx. Mild to moderately severe infections: 250 to 500 mg (400,000 to 800,000 units) every 6 to 8 hours.

For the prevention of recurrence following rheumatic fever and/or chorea: 125 to 250 mg (200,000 to 400,000 units) twice daily on a continuing basis.

For prophylaxis against bacterial endocarditis¹ in patients with congenital heart disease or rheumatic or other acquired valvular heart disease when undergoing dental procedures or surgical procedures of the upper respiratory tract: 2 gram of penicillin V (1 gram for children under 60 lbs.) 1 hour before the procedure, and then, 1 gram (500 mg for children under 60 lbs.) 6 hours later.

Directions for Mixing Oral Solution

Do not add water until you dispense. When dispensing, tap bottle until all powder flows freely, slowly add the total amount of water for reconstitution (see table below). After partially filling bottle, replace cap and shake vigorously. Add remaining water and repeat shaking. **After reconstitution, solution must be stored in a refrigerator. Discard any unused portion after 14 days.**

125 mg/5 mL	
Bottle size	Total Amount of Water Required for Reconstitution
100 mL	75 mL
200 mL	150 mL

The resulting solution (red in color) will contain penicillin V potassium equivalent to penicillin V 125 mg (200,000 units) in each 5 mL (teaspoonful).

250 mg/5 mL

Bottle size	Total Amount of Water Required for Reconstitution
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100 mL	75 mL
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200 mL	150 mL
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The resulting solution (red in color) will contain penicillin V potassium equivalent to penicillin V 250 mg (400,000 units) in each 5 mL (teaspoonful).

HOW SUPPLIED

Penicillin V Potassium Tablets, USP are available as follows:

250 mg (400,000 units): biconvex, oval, mottled, white to off-white, uncoated tablets, debossed "93" on one side and "1172" on the other side in bottles of 100 (NDC 0093-1172-01) and 1000 (NDC 0093-1172-10).

500 mg (800,000 units): biconvex, oval, mottled, white to off-white, uncoated, scored tablets, debossed "9" to the left of partial bisect and "3" to the right on one side and "1174" on the other side in bottles of 100 (NDC 0093-1174-01) and 1000 (NDC 0093-1174-10).

Store at 20° to 25°C (68° to 77°F) [See USP Controlled Room Temperature]. Keep tightly closed.

Dispense in a tight container as defined in the USP, with a child-resistant closure as required.

Penicillin V Potassium for Oral Solution, USP is available as follows:

125 mg (200,000 Units) per 5 mL: bottles of 100 mL (NDC 0093-4125-73) and 200 mL (NDC 0093-4125-74).

250 mg (400,000 Units) per 5 mL: bottles of 100 mL (NDC 0093-4127-73) and 200 mL (NDC 0093-4127-74).

Keep tightly closed. Store dry powder at 20° to 25°C (68° to 77°F) [See USP Controlled Room Temperature]. After reconstitution, solution must be stored in a refrigerator. Discard any unused portion after 14 days.

Keep this and all medications out of the reach of children.

REFERENCES

1. American Heart Association. 1984. Prevention of bacterial endocarditis. Circulation 70 (6): 1123A-1127A.

Manufactured In Canada By:

Teva Canada Limited

Toronto, Canada M1B 2K9

Manufactured For:

Teva Pharmaceuticals

Parsippany, NJ 07054

Rev. P 1/2023

Package/Label Display Panel

NDC 0093-1172-01

Penicillin V
Potassium
Tablets, USP
250 mg (400,000 Units)

Rx only

100 Oval Tablets

Each tablet contains penicillin V potassium equivalent to 250 mg (400,000 units) penicillin V. Each tablet contains 0.71 mEq (27.9 mg) of potassium.
Usual Dosage: 250 mg (400,000 units) every 6 to 8 hours. See accompanying literature.
Store at 20° to 25°C (68° to 77°F) [See USP Controlled Room Temperature]. **Keep tightly closed.**
Dispense in a tight container as defined in the USP, with a child-resistant closure (as required).
Keep this and all medications out of the reach of children.
Manufactured In Canada By: Teva Canada Limited
Toronto, Canada M1B 2K9
Manufactured For: Teva Pharmaceuticals
Parsippany, NJ 07054
361-32- 787096240A Rev 04 Rev. L 1/2023

NDC 0093-1172-01

**Penicillin V
Potassium
Tablets, USP**
250 mg (400,000 Units)

GTIN 00300931172013

3 0093-1172-01 3

Serialization Coding Area

Rx only
100 Oval Tablets

teva

Package/Label Display Panel

NDC 0093-1174-01

Penicillin V
Potassium
Tablets, USP
500 mg (800,000 Units)

Rx only

100 Oval Tablets



Each tablet contains penicillin V potassium equivalent to 500 mg (800,000 units) penicillin V. Each tablet contains 1.43 mEq (55.8 mg) of potassium.

Usual Dosage: 250 mg to 500 mg (400,000 to 800,000 units) every 6 to 8 hours. See accompanying literature.

Store at 20° to 25°C (68° to 77°F) [See USP Controlled Room Temperature]. **Keep tightly closed.**

Dispense in a tight container as defined in the USP, with a child-resistant closure (as required).

Keep this and all medications out of the reach of children.

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361-32- 787126240A Rev 04 Rev. K 1/2023

GTIN 00300931174017

NDC 0093-1174-01

Penicillin V Potassium Tablets, USP

500 mg (800,000 Units)

Rx only
100 Oval Tablets



Serialization Coding Area

Package/Label Display Panel

NDC 0093-4125-73

Penicillin V
Potassium for
Oral Solution, USP
125 mg (200,000 U) per 5 mL

WARNING: NOT FOR INJECTION

Rx only
100 mL (when mixed)



When reconstituted as directed each 5 mL contains penicillin V potassium equivalent to 125 mg (200,000 units) of penicillin V. Each 5 mL contains 0.36 mEq (13.9 mg) of potassium.

Usual Dosage: 125 mg to 250 mg (200,000 to 400,000 Units) every 6 to 8 hours. See accompanying literature.

Each bottle contains penicillin V potassium equivalent to 2.5 g penicillin V.

IMPORTANT
Keep in refrigerator. Discard any unused portion after two weeks.

Date of reconstitution: _____

Shake well before using.

Keep this and all medications out of the reach of children.

Store dry powder at 20° to 25°C (68° to 77°F) [See USP Controlled Room Temperature].

Pharmacist Directions for Mixing: Do not add water until you dispense. When dispensing tap bottle until all powder flows freely, slowly add 75 mL of water. After partially filling bottle, replace cap and shake vigorously. Add remaining water and repeat shaking. The resulting solution is red in color.

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Manufactured For: Teva Pharmaceuticals
Parsippany, NJ 07054

361-32- 787146240A Rev 05 Rev. I 1/2023

GTIN 00300934125733

NDC 0093-4125-73

Penicillin V Potassium for Oral Solution, USP

125 mg (200,000 U) per 5 mL

WARNING: NOT FOR INJECTION

Rx only
100 mL (when mixed)



Serialization Coding Area

Package/Label Display Panel

NDC 0093-4127-73

Penicillin V
Potassium for
Oral Solution, USP
250 mg (400,000 U) per 5 mL

WARNING: NOT FOR INJECTION

Rx only

100 mL (when mixed)

GTIN 00300934127737

NDC 0093-4127-73

Penicillin V Potassium for Oral Solution, USP

250 mg (400,000 U) per 5 mL

WARNING: NOT FOR INJECTION

Rx only
100 mL (when mixed)

teva

When reconstituted as directed each 5 mL contains penicillin V potassium equivalent to 250 mg (400,000 units) of penicillin V. Each 5 mL contains 0.71 mEq (27.9 mg) of potassium. **Usual Dosage:** 250 mg (400,000 Units) every 6 to 8 hours. See accompanying literature. Each bottle contains penicillin V potassium equivalent to 5 g penicillin V.

IMPORTANT
Keep in refrigerator. Discard any unused portion after two weeks.

Date of reconstitution: _____
Shake well before using.
Keep this and all medications out of the reach of children.

Store dry powder at 20° to 25°C (68° to 77°F) [See USP Controlled Room Temperature].

Pharmacist: Directions for Mixing: Do not add water until you dispense. When dispensing, tap bottle until all powder flows freely, slowly add 7.5 mL of water. After partially filling bottle, replace cap and shake vigorously. Add remaining water and repeat shaking. The resulting solution is red in color.

Manufactured in Canada By:
Teva Canada Limited
Toronto, Canada M1B 2K9

Manufactured for:
Teva Pharmaceuticals
 Parsippany, NJ 07054

361-32- 787156240A Rev 05 Rev. H 1/2023

0093-4127-73

3 7

Serialization Coding Area

PENICILLIN V POTASSIUM

penicillin v potassium tablet

Product Information

Product Type	HUMAN PRESCRIPTION DRUG	Item Code (Source)	NDC:0093-1172
Route of Administration	ORAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
PENICILLIN V POTASSIUM (UNII: 146T0TU1JB) (PENICILLIN V - UNII:Z61I075U2W)	PENICILLIN V	250 mg

Inactive Ingredients

Ingredient Name	Strength
CALCIUM PHOSPHATE, DIBASIC, ANHYDROUS (UNII: L11K75P92J)	
MAGNESIUM STEARATE (UNII: 70097M6130)	

CELLULOSE, MICROCRYSTALLINE (UNII: OP1R32D61U)	
ANHYDROUS TRISODIUM CITRATE (UNII: RS7A450LGA)	

Product Characteristics

Color	white (white to off-white)	Score	no score
Shape	OVAL	Size	13mm
Flavor		Imprint Code	93;1172
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:0093-1172-01	100 in 1 BOTTLE; Type 0: Not a Combination Product	09/30/1990	
2	NDC:0093-1172-10	1000 in 1 BOTTLE; Type 0: Not a Combination Product	09/30/1990	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
ANDA	ANDA060711	09/30/1990	

PENICILLIN V POTASSIUM

penicillin v potassium tablet

Product Information

Product Type	HUMAN PRESCRIPTION DRUG	Item Code (Source)	NDC:0093-1174
Route of Administration	ORAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
PENICILLIN V POTASSIUM (UNII: 146T0TU1JB) (PENICILLIN V - UNII:Z61I075U2W)	PENICILLIN V	500 mg

Inactive Ingredients

Ingredient Name	Strength
CALCIUM PHOSPHATE, DIBASIC, ANHYDROUS (UNII: L11K75P92J)	
MAGNESIUM STEARATE (UNII: 70097M6I30)	
CELLULOSE, MICROCRYSTALLINE (UNII: OP1R32D61U)	
ANHYDROUS TRISODIUM CITRATE (UNII: RS7A450LGA)	

Product Characteristics

Color	white (white to off-white)	Score	2 pieces
Shape	OVAL	Size	17mm
Flavor		Imprint Code	9;3;1174
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:0093-1174-01	100 in 1 BOTTLE; Type 0: Not a Combination Product	09/30/1990	
2	NDC:0093-1174-10	1000 in 1 BOTTLE; Type 0: Not a Combination Product	09/30/1990	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
ANDA	ANDA060711	09/30/1990	

PENICILLIN V POTASSIUM

penicillin v potassium powder, for solution

Product Information

Product Type	HUMAN PRESCRIPTION DRUG	Item Code (Source)	NDC:0093-4125
Route of Administration	ORAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
PENICILLIN V POTASSIUM (UNII: 146T0TU1JB) (PENICILLIN V - UNII:Z61I075U2W)	PENICILLIN V	125 mg in 5 mL

Inactive Ingredients

Ingredient Name	Strength
FD&C RED NO. 40 (UNII: WZB9127XOA)	
SACCHARIN SODIUM (UNII: SB8ZUX40TY)	
SODIUM BENZOATE (UNII: OJ245FE5EU)	
SUCROSE (UNII: C151H8M554)	

Product Characteristics

Color		Score	
Shape		Size	

Flavor	CHERRY	Imprint Code		
Contains				
Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:0093-4125-73	100 mL in 1 BOTTLE; Type 0: Not a Combination Product	09/30/1990	
2	NDC:0093-4125-74	200 mL in 1 BOTTLE; Type 0: Not a Combination Product	09/30/1990	
Marketing Information				
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
ANDA	ANDA060456	09/30/1990		

PENICILLIN V POTASSIUM			
penicillin v potassium powder, for solution			
Product Information			
Product Type	HUMAN PRESCRIPTION DRUG	Item Code (Source)	NDC:0093-4127
Route of Administration	ORAL		
Active Ingredient/Active Moiety			
Ingredient Name	Basis of Strength	Strength	
PENICILLIN V POTASSIUM (UNII: 146T0TU1JB) (PENICILLIN V - UNII:Z61I075U2W)	PENICILLIN V	250 mg in 5 mL	
Inactive Ingredients			
Ingredient Name	Strength		
FD&C RED NO. 40 (UNII: WZB9127XOA)			
SACCHARIN SODIUM (UNII: SB8ZUX40TY)			
SODIUM BENZOATE (UNII: OJ245FE5EU)			
SUCROSE (UNII: C151H8M554)			
Product Characteristics			
Color		Score	
Shape		Size	
Flavor	CHERRY	Imprint Code	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:0093-4127-73	100 mL in 1 BOTTLE; Type 0: Not a Combination Product	09/30/1990	
2	NDC:0093-4127-74	200 mL in 1 BOTTLE; Type 0: Not a Combination Product	09/30/1990	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
ANDA	ANDA060456	09/30/1990	

Labeler - Teva Pharmaceuticals USA, Inc. (001627975)

Revised: 1/2023

Teva Pharmaceuticals USA, Inc.