PFIZERPEN- penicillin g potassium powder, for solution
Roerig

Buffered
PFIZERPEN
(Penicillin G Potassium for Injection, USP)

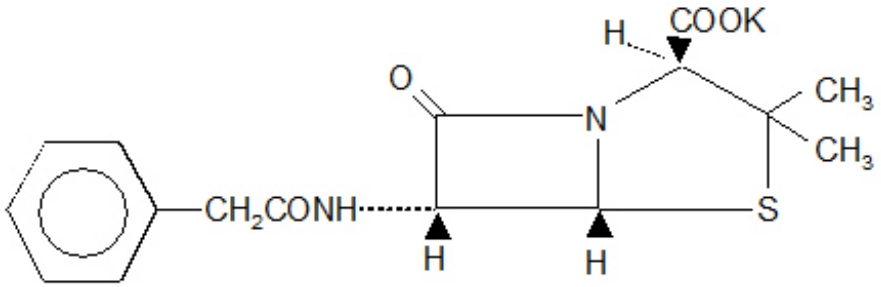
To reduce the development of drug-resistant bacteria and maintain the effectiveness of Penicillin G Potassium and other antibacterial drugs, Buffered Pfizerpen (Penicillin G Potassium for Injection, USP) should be used only to treat or prevent infections that are proven or strongly suspected to be caused by bacteria.

DESCRIPTION

Buffered Pfizerpen (Penicillin G Potassium for Injection, USP) is a sterile, pyrogen-free powder for reconstitution. Buffered Pfizerpen for Injection is an antibacterial agent for intramuscular, continuous intravenous infusion, intrapleural or other local infusion, and intrathecal administration.

Each million units contains approximately 6.8 milligrams of sodium (0.3 mEq) and 65.6 milligrams of potassium (1.68 mEq). Buffered Pfizerpen (Penicillin G Potassium for Injection, USP) is supplied in vials equivalent to 5,000,000 units (5 million units) or 20,000,000 units (20 million units) of penicillin G as the potassium salt.

Chemically, Pfizerpen is monopotassium 3,3-dimethyl-7-oxo-6-(2-phenylacetamido)-4-thia-1-azabicyclo (3.2.0) heptane-2-carboxylate. It has a molecular weight of 372.48 and the following chemical structure:



Formula
$C_{16}H_{17}KN_2O_4S$

Penicillin G potassium is a colorless or white crystal, or a white crystalline powder which is odorless, or practically so, and moderately hygroscopic. Penicillin G potassium is very soluble in water. The pH of the reconstituted product is between 6.0–8.5.

CLINICAL PHARMACOLOGY

Aqueous penicillin G is rapidly absorbed following both intramuscular and subcutaneous injection. Initial blood levels following parenteral administration are high but transient. Penicillins bind to serum proteins, mainly albumin. Therapeutic levels of the penicillins are

easily achieved under normal circumstances in extracellular fluid and most other body tissues. Penicillins are distributed in varying degrees into pleural, pericardial, peritoneal, ascitic, synovial, and interstitial fluids. Penicillins are excreted in breast milk. Penetration into the cerebrospinal fluid, eyes, and prostate is poor. Penicillins are rapidly excreted in the urine by glomerular filtration and active tubular secretion, primarily as unchanged drug. Approximately 60 percent of the total dose of 300,000 units is excreted in the urine within this 5-hour period. For this reason, high and frequent doses are required to maintain the elevated serum levels desirable in treating certain severe infections in individuals with normal kidney function. In neonates and young infants, and in individuals with impaired kidney function, excretion is considerably delayed.

After an intravenous infusion of penicillin G, peak serum concentrations are attained immediately after completion of the infusion. In a study of ten patients administered a single 5 million unit dose of penicillin G intravenously over 3–5 minutes, the mean serum concentrations were 400 mcg/mL, 273 mcg/mL and 3.0 mcg/mL at 5–6 minutes, 10 minutes and 4 hours after completion of the injection, respectively. In a separate study, five healthy adults were administered one million units of penicillin G intravenously, either as a bolus over 4 minutes or as an infusion over 60 minutes. The mean serum concentration eight minutes after completion of the bolus was 45 mcg/mL and eight minutes after completion of the infusion was 14.4 mcg/mL. The mean β-phase serum half-life of penicillin G administered by the intravenous route in ten patients with normal renal function was 42 minutes, with a range of 31–50 minutes.

The clearance of penicillin G in normal individuals is predominantly via the kidney. The renal clearance, which is extremely rapid, is the result of glomerular filtration and active tubular transport, with the latter route predominating. Urinary recovery is reported to be 58–85% of the administered dose. Renal clearance of penicillin is delayed in premature infants, neonates and in the elderly due to decreased renal function. The serum half-life of penicillin G correlates inversely with age and clearance of creatinine and ranges from 3.2 hours in infants 0 to 6 days of age to 1.4 hours in infants 14 days of age or older.

Nonrenal clearance includes hepatic metabolism and, to a lesser extent, biliary excretion. The latter routes become more important with renal impairment.

Probenecid blocks the renal tubular secretion of penicillin. Therefore, the concurrent administration of probenecid prolongs the elimination of penicillin G and, consequently, increases serum concentrations.

Penicillin G is distributed to most areas of the body including lung, liver, kidney, muscle, bone and placenta. In the presence of inflammation, levels of penicillin in abscesses, middle ear, pleural, peritoneal and synovial fluids are sufficient to inhibit most susceptible bacteria. Penetration in the eye, brain, cerebrospinal fluid (CSF) or prostate is poor in the absence of inflammation. With inflamed meninges, the penetration of penicillin G into the CSF improves, such that the CSF/serum ratio is 2–6%. Inflammation also enhances its penetration into the pericardial fluid. Penicillin G is actively secreted into the bile resulting in levels at least 10 times those achieved simultaneously in serum. Penicillin G penetrates poorly into human polymorphonuclear leukocytes.

In the presence of impaired renal function, the β-phase serum half-life of penicillin G is prolonged. β-phase serum half-lives of one to two hours were observed in azotemic patients with serum creatinine concentrations <3 mg/100 mL and ranged as high as 20 hours in anuric patients. A linear relationship, including the lowest range of renal function, is found between the serum elimination rate constant and renal function as

measured by creatinine clearance.

In patients with altered renal function, the presence of hepatic insufficiency further alters the elimination of penicillin G. In one study, the serum half-lives in two anuric patients (excreting <400 mL urine/day) were 7.2 and 10.1 hours. A totally anuric patient with terminal hepatic cirrhosis had a penicillin half-life of 30.5 hours, while another patient with anuria and liver disease had a serum half-life of 16.4 hours. The dosage of penicillin G should be reduced in patients with severe renal impairment, with additional modifications when hepatic disease accompanies the renal impairment. Hemodialysis has been shown to reduce penicillin G serum levels.

Microbiology

Penicillin G exerts a bactericidal action against penicillin-susceptible microorganisms during the stage of active multiplication. It acts through the inhibition of biosynthesis of cell wall peptidoglycan rendering the cell wall osmotically unstable. It is not active against the penicillinase-producing bacteria, which include many strains of staphylococci or against organisms resistant to beta-lactams because of alterations in the penicillin-binding proteins, such as methicillin-resistant staphylococci. Penicillin G is highly active *in vitro* against streptococci (groups A, B, C, G, H, L, and M), and *Neisseria meningitidis*.

Other organisms susceptible to penicillin G are *N. gonorrhoeae*, *Corynebacterium diphtheriae*, *Bacillus anthracis*, *Clostridium* spp, *Actinomyces* species, "*Spirillum minus*", *Streptobacillus moniliformis*, *Listeria monocytogenes* and *Leptospira* spp; *Treponema pallidum* is extremely sensitive to the bactericidal action of penicillin G.

Some species of gram-negative bacilli were previously considered susceptible to very high intravenous doses of penicillin G (up to 80 million units/day) including some strains of *Escherichia coli, Proteus mirabilis*, *Salmonella* spp. and *Shigella* spp.; *Enterobacter aerogenes* (formerly *Aerobacter aerogenes)* and *Alcaligenes faecalis*. Penicillin G is no longer considered a drug of choice for infections caused by these organisms.

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

Therapy

Penicillin G Potassium for Injection is indicated in the treatment of serious infections caused by susceptible strains of the designated microorganisms in the conditions listed below.

Appropriate culture and susceptibility tests should be done before treatment in order to isolate and identify organisms causing infection and to determine their susceptibility to penicillin G.

Therapy with Penicillin G Potassium for Injection may be initiated before results of such tests are known when there is reason to believe the infection may involve any of the organisms listed below; however, once these results become available, appropriate

therapy should be continued.

CLINICAL INDICATION	INFECTING ORGANISM
Septicemia, empyema, pneumonia, pericarditis, endocarditis, meningitis	*Streptococcus pyogenes* (group A β-hemolytic streptococcus), other β-hemolytic streptococci including groups C, H, G, L and M, *Streptococcus pneumoniae* and *Staphylococcus* species (non-penicillinase producing strains)
Anthrax	*Bacillus anthracis*
Actinomycosis (cervico-facial disease and thoracic and abdominal disease)	*Actinomyces israelii*
Botulism (adjunctive therapy to antitoxin), gas gangrene, and tetanus (adjunctive therapy to human tetanus immune globulin)	*Clostridium* species
Diphtheria (adjunctive therapy to antitoxin and prevention of the carrier state)	*Corynebacterium diphtheriae*
Erysipelothrix endocarditis	*Erysipelothrix rhusiopathiae*
Fusospirochetosis (severe infections of the oropharynx [Vincent's], lower respiratory tract and genital area)	*Fusobacterium* species and spirochetes
Listeria infections including meningitis and endocarditis	*Listeria monocytogenes*
Pasteurella infections including bacteremia and meningitis	*Pasteurella multocida*
Haverhill fever	*Streptobacillus moniliformis*
Rat bite fever	*Spirillum minus* or *Streptobacillus moniliformis*
Disseminated gonococcal infections	*Neisseria gonorrhoeae* (Penicillin-susceptible)
Syphilis (congenital and neurosyphilis)	*Treponema pallidum*
Meningococcal meningitis and/or septicemia	*Neisseria meningitidis*
Gram-negative bacillary infections (bacteremias) **Penicillin G is not the drug of choice in the treatment of Gram-negative bacillary infections.**	Gram-negative bacillary organisms (*i.e. Enterobacteriaceae)*

To reduce the development of drug-resistant bacteria and maintain effectiveness of Penicillin G Potassium and other antibacterial drugs, Penicillin G Potassium 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 history of a hypersensitivity (anaphylactic) reaction to any penicillin is a contraindication.

WARNINGS

Anaphylaxis

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 G, careful inquiry should be made concerning previous hypersensitivity reactions to penicillins, cephalosporins, or other allergens. If an allergic reaction occurs, penicillin G should be discontinued and appropriate therapy instituted.

Severe cutaneous adverse reactions

Severe cutaneous adverse reactions (SCAR), such as Stevens-Johnson syndrome (SJS), toxic epidermal necrolysis (TEN), drug reaction with eosinophilia and systemic symptoms (DRESS), and acute generalized exanthematous pustulosis (AGEP) have been reported in patients taking beta-lactam antibiotics. When SCAR is suspected, Penicillin G Potassium for Injection should be discontinued immediately and an alternative treatment should be considered.

Clostridioides difficile associated diarrhea

Clostridioides difficile associated diarrhea (CDAD) has been reported with use of nearly all antibacterial agents, including Penicillin G Potassium for Injection, 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 (see **Warnings**). Whenever allergic reactions occur, penicillin should be

withdrawn unless, in the opinion of the physician, the condition being treated is life-threatening and amenable only to penicillin therapy. Penicillin G Potassium for Injection by the intravenous route in high doses (above 10 million units) should be administered slowly because of the potential adverse effects of electrolyte imbalance from the potassium content of the penicillin. Penicillin G Potassium for Injection contains 1.68 mEq potassium and 0.3 mEq of sodium per million units. The use of antibiotics may promote overgrowth of nonsusceptible organisms, including fungi. Indwelling intravenous catheters encourage superinfections. Should superinfection occur, appropriate measures should be taken. When indicated, incision and drainage or other surgical procedures should be performed in conjunction with antibiotic therapy.

Prescribing Penicillin G Potassium for Injection in the absence of 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.

Intramuscular and Intravenous Therapy

Care should be taken to avoid intravenous or accidental intraarterial administration, or injection into or near major peripheral nerves or blood vessels, since such injections may produce neurovascular damage. Particular care should be taken with IV administration because of the possibility of thrombophlebitis.

Information for Patients

Patients should be counseled that antibacterial drugs including Penicillin G Potassium for Injection should only be used to treat bacterial infections. They do not treat viral infections (e.g., the common cold). When Penicillin G Potassium for Injection is 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 G Potassium for Injection 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.

Laboratory Tests

Periodic assessment of organ system function, including frequent evaluation of electrolyte balance, hepatic, renal and hematopoietic systems, and cardiac and vascular status should be performed during prolonged therapy with high doses of intravenous penicillin G (see **Adverse Reactions**). If any impairment of function is suspected or known to exist, a reduction in the total dosage should be considered (see **Dosage and Administration**). In suspected staphylococcal infections, proper laboratory studies, including susceptibility tests should be performed. All infections due to Group A beta-hemolytic streptococci should be treated for at least 10 days.

Patients being treated for gonococcal infection should have a serologic test for syphilis before receiving penicillin. All cases of penicillin treated syphilis should receive adequate

follow-up including clinical and serological examinations. The recommended follow-up varies with the stage of syphilis being treated.

Drug Interactions

Bacteriostatic antibacterials (*i.e.*, chloramphenicol, erythromycins, sulfonamides or tetracyclines) may antagonize the bactericidal effect of penicillin, and concurrent use of these drugs should be avoided. This has been documented *in vitro*; however, the clinical significance of this interaction is not well-documented.

Penicillin blood levels may be prolonged by concurrent administration of probenecid which blocks the renal tubular secretion of penicillins. Other drugs may compete with penicillin G for renal tubular secretion and thus prolong the serum half-life of penicillin. These drugs include: aspirin, phenylbutazone, sulfonamides, indomethacin, thiazide diuretics, furosemide and ethacrynic acid.

Drug/Laboratory Test Interactions

After treatment with penicillin G, a false-positive reaction for glucose in the urine may occur with Benedict's solution, Fehling's solution or Clinitest® tablet, but not with the enzyme-based tests, such as Clinistix® and Tes-Tape®.

Penicillin G has been associated with pseudoproteinuria by certain test methods.

Carcinogenesis, Mutagenesis, Impairment of Fertility

No long term animal studies have been conducted with this drug.

Pregnancy

Teratogenic Effects

Reproduction studies performed in the mouse, rat, and rabbit have revealed no evidence of impaired fertility or harm to the fetus due to penicillin G. Human experience with the penicillins during pregnancy has not shown any positive evidence of adverse effects on the fetus. There are, however, no adequate and well controlled studies in pregnant women showing conclusively that harmful effects of these drugs on the fetus can be excluded. Because animal reproduction studies are not always predictive of human response, this drug should be used during pregnancy only if clearly needed.

Nursing Mothers

Penicillins are excreted in human milk. Caution should be exercised when penicillins are administered to a nursing woman.

Pediatric Use

Incompletely developed renal function in newborns may delay elimination of penicillin; therefore, appropriate reductions in the dosage and frequency of administration should be made in these patients. All newborns treated with penicillins should be monitored closely for clinical and laboratory evidence of toxic or adverse effects (see **Precautions**).

Pediatric doses are generally determined on a weight basis and should be calculated for

each patient individually. Recommended guidelines for pediatric dosages are presented in **Dosage and Administration**.

Geriatric Use

Clinical studies of penicillin G injection did not include sufficient numbers of subjects aged 65 and over to determine whether they respond differently from younger subjects. Other reported clinical experience has not identified differences in responses between the elderly and younger patients. In general, dose selection for an elderly patient should be cautious, usually starting at the low end of the dosing range, reflecting the greater frequency of decreased hepatic, renal, or cardiac function, and of concomitant disease or other drug therapy.

This drug is known to be substantially excreted by the kidney, and the risk of toxic reactions to this drug may be greater in patients with impaired renal function. Because elderly patients are more likely to have decreased renal function, care should be taken in dose selection, and it may be useful to monitor renal function.

Penicillin G Potassium for Injection contains 6.8 mg (0.3 mEq) of sodium per million units. At the usual recommended doses of 10 to 20 million units per day patients would receive between 68 and 136 mg/day (3 and 6 mEq) of sodium per day. The geriatric population may respond with a blunted natriuresis to salt loading. This may be clinically important with regard to such diseases as congestive heart failure.

ADVERSE REACTIONS

Body as a whole

The Jarisch-Herxheimer reaction is a systemic reaction that may occur after the initiation of penicillin therapy in patients with syphilis or other spirochetal infections (*i.e.*, Lyme disease and Relapsing fever). The reaction begins one or two hours after initiation of therapy and disappears within 12 to 24 hours. It is characterized by fever, chills, myalgias, headache, exacerbation of cutaneous lesions, tachycardia, hyperventilation, vasodilation with flushing and mild hypotension. The pathogenesis of the Herxheimer reaction may be due to the release from the spirochetes of heat-stable pyrogen.

Hypersensitivity reactions

The reported incidence of allergic reactions to all penicillins ranges from 0.7 to 10 percent in different studies (see **Warnings**). Sensitization is usually the result of previous treatment with a penicillin, but some individuals have had immediate reactions when first treated. In such cases, it is postulated that prior exposure to penicillin may have occurred via trace amounts present in milk or vaccines.

Two types of allergic reactions to penicillin are noted clinically – immediate and delayed.

Immediate reactions usually occur within 20 minutes of administration and range in severity from urticaria and pruritus to angioneurotic edema, laryngospasm, bronchospasm, hypotension, vascular collapse and death (see **Warnings**). Such immediate anaphylactic reactions are very rare and usually occur after parenteral therapy, but a few cases of anaphylaxis have been reported following oral therapy. Another type of immediate reaction, an accelerated reaction, may occur between 20

minutes and 48 hours after administration and may include urticaria, pruritus, fever and, occasionally, laryngeal edema.

Delayed reactions to penicillin therapy usually occur within 1–2 weeks after initiation of therapy. Manifestations include serum sickness-like symptoms, *i.e.*, fever, malaise, urticaria, myalgia, arthralgia, abdominal pain and various skin rashes, ranging from maculopapular eruptions to exfoliative dermatitis.

Contact dermatitis has been observed in individuals who prepare penicillin solutions.

Gastrointestinal system

Pseudomembranous colitis has been reported with the onset occurring during or after penicillin G treatment. Nausea, vomiting, stomatitis, black or hairy tongue, and other symptoms of gastrointestinal irritation may occur, especially during oral therapy.

Hematologic system

Reactions include neutropenia, which resolves after penicillin therapy is discontinued; Coombs-positive hemolytic anemia, an uncommon reaction, occurs in patients treated with intravenous penicillin G in doses greater than 10 million units/day and who have previously received large doses of the drug; and with large doses of penicillin, a bleeding diathesis can occur secondary to platelet dysfunction.

Metabolic

Penicillin G Potassium for Injection (1 million units contains 1.68 mEq of potassium ion) may cause serious and even fatal electrolyte disturbances, *i.e.*, hyperkalemia, when given intravenously in large doses.

Nervous system

Neurotoxic reactions including hyperreflexia, myoclonic twitches, seizures and coma have been reported following the administration of massive intravenous doses, and are more likely in patients with impaired renal function.

Urogenital system

Renal tubular damage and interstitial nephritis have been associated with large intravenous doses of penicillin G. Manifestations of this reaction may include fever, rash, eosinophilia, proteinuria, eosinophiluria, hematuria and a rise in serum urea nitrogen.

Discontinuation of penicillin G results in resolution in the majority of patients.

Local reactions

Phlebitis and thrombophlebitis may occur, and pain at the injection site has been reported with intravenous administration.

Immune system disorders

Acute myocardial ischemia with or without myocardial infarction may occur as part of an allergic reaction.

To report SUSPECTED ADVERSE EVENTS, contact Pfizer Inc. at 1-800-438-1985 or FDA

at 1-800-FDA-1088 or http://www.fda.gov/ for voluntary reporting of adverse reactions.

OVERDOSAGE

Dose related toxicity may arise with the use of massive doses of intravenous penicillins (40 to 100 million units per day), particularly in patients with severe renal impairment (see **Precautions**). The manifestations may include agitation, confusion, asterixis, hallucinations, stupor, coma, multifocal myoclonus, seizures and encephalopathy. Hyperkalemia is also possible (see **Adverse Reactions-Metabolic**).

In case of overdosage, discontinue penicillin, treat symptomatically, and institute supportive measures as required. If necessary, hemodialysis may be used to reduce blood levels of Penicillin G, although the degree of effectiveness of this procedure is questionable.

DOSAGE AND ADMINISTRATION

Buffered Penicillin G Potassium for Injection, USP may be given intravenously or intramuscularly. The usual dose recommendations are as follows:

Adult patients

CLINICAL INDICATION	DOSAGE
Serious infections due to susceptible strains of streptococci (including *S. pneumoniae*)	12 to 24 million units/day depending on the infection and its severity administered in equally divided doses every 4–6 hours.
-septicemia, empyema, pneumonia, pericarditis, endocarditis and meningitis	
Serious infections due to susceptible strains of staphylococci	5 to 24 million units/day depending on the infection and its severity administered in equally divided doses every 4–6 hours.
-septicemia, empyema, pneumonia, pericarditis, endocarditis and meningitis	
Anthrax	Minimum of 8 million units/day in divided doses every 6 hours. Higher doses may be required depending on susceptibility of organism.
Actinomycosis	
Cervicofacial disease	1 to 6 million units/day(*)
Thoracic and abdominal disease	10 to 20 million units/day(*)
Clostridial infections	
Botulism (adjunctive therapy to antitoxin)	20 million units/day(*)
Gas gangrene (debridement and/or surgery as indicated)	
Tetanus (adjunctive therapy to human tetanus immune globulin)	
Diphtheria (adjunctive therapy to antitoxin and for the prevention of the carrier state)	2 to 3 million units/day in divided doses for 10–12 days(*)

Erysipelothrix endocarditis	12 to 20 million units/day for 4–6 weeks(*)
Fusospirochetosis (severe infections of the oropharynx [Vincent's], lower respiratory tract and genital area)	5 to 10 million units/day(*)
Listeria infections	
Meningitis	15 to 20 million units/day for 2 weeks(*)
Endocarditis	15 to 20 million units/day for 4 weeks(*)
Pasteurella infections including bacteremia and meningitis	4 to 6 million units/day for 2 weeks(*)
Haverhill fever; Rat-bite fever	12 to 20 million units/day for 3–4 weeks(*)
Disseminated gonococcal infections, such as meningitis, endocarditis, arthritis, etc., caused by penicillin – susceptible organisms	10 million units/day(*); duration depends on the type of infection
Syphilis (neurosyphilis)	12 to 24 million units/day, as 2–4 MU every 4 hours for 10–14 days; many experts recommend additional therapy with Benzathine PCN G 2.4 MU IM weekly for 3 doses after completion of IV therapy
Meningococcal meningitis and/or septicemia	24 million units/day as 2 million units every 2 hours

* Because of its short half-life, Penicillin G is administered in divided doses, usually every 4–6 hours with the exception of meningococcal meningitis/septicemia, *i.e.*, every 2 hours.

Pediatric patients

This product should not be administered to patients requiring less than one million units per dose (see **Precautions-Pediatric Use**).

CLINICAL INDICATION	DOSAGE
Serious infections, such as pneumonia and endocarditis, due to susceptible strains of streptococci (including *S. pneumoniae*) and meningococcus	150,000–300,000 units/kg/day divided in equal doses every 4–6 hours; duration depends on infecting organism and type of infection
Meningitis caused by susceptible strains of pneumococcus and meningococcus	250,000 units/kg/day divided in equal doses every 4 hours for 7–14 days depending on the infecting organism (maximum dose of 12–20 million units/day)
Disseminated Gonococcal Infections (penicillin-susceptible strains)	Weight less than 45 kg:
Arthritis	100,000 units/kg/day in 4 equally divided doses for 7–10 days
Meningitis	250,000 units/kg/day in equal doses every 4 hours for 10–14 days
Endocarditis	250,000 units/kg/day in equal doses every 4 hours for 4 weeks
Arthritis, meningitis, endocarditis	Weight 45 kg or greater: 10 million units/day in equally divided doses with the duration of therapy depending on the type

	of infection
Syphilis (congenital and neurosyphilis) after the newborn period	200,000–300,000 units/kg/day (administered as 50,000 units/kg every 4–6 hours) for 10–14 days
Diphtheria (adjunctive therapy to antitoxin and for prevention of the carrier state)	150,000–250,000 units/kg/day in equal doses every 6 hours for 7–10 days
Rat-bite fever; Haverhill fever (with endocarditis caused by *S. moniliformis*)	150,000–250,000 units/kg/day in equal doses every 4 hours for 4 weeks

Renal Impairment

Penicillin G is relatively nontoxic, and dosage adjustments are generally required only in cases of severe renal impairment. The recommended dosage regimens are as follows:

Creatinine clearance less than 10 mL/min/1.73 m^2; administer a full loading dose (see recommended dosages in the tables above) followed by one-half of the loading dose every 8–10 hours.

Uremic patients with a creatinine clearance greater than 10 mL/min/1.73 m^2; administer a full loading dose (see recommended dosages in the tables above) followed by one-half of the loading dose every 4–5 hours. Additional dosage modification should be made in patients with hepatic disease and renal impairment.

For most acute infections, treatment should be continued for at least 48 to 72 hours after the patient becomes asymptomatic. Antibiotic therapy for Group A β-hemolytic streptococcal infections should be maintained for atleast 10 days to reduce the risk of rheumatic fever.

Parenteral drug products should be inspected visually for particulate matter and discoloration prior to administration whenever solution and container permit.

Reconstitution

The following table shows the amount of solvent required for solution of various concentrations:

Approx. Desired Concentration (units/mL)	Volume (mL) Solvent for Vial of 5,000,000 units	Infusion Only Volume (mL) Solvent for Vial of 20,000,000 units
50,000	–	–
100,000	–	–
250,000	18.2	75.0
500,000	8.2	33.0
1,000,000	3.2	11.5

When the required volume of solvent is greater than the capacity of the vial, the penicillin can be dissolved by first injecting only a portion of the solvent into the vial, then withdrawing the resultant solution and combining it with the remainder of the solvent in a larger sterile container.

Buffered Pfizerpen (Penicillin G Potassium for Injection, USP) is highly water soluble. It may be dissolved in small amounts of Water for Injection, or Sterile Isotonic Sodium Chloride Solution for Parenteral Use. All solutions should be stored in a refrigerator. When refrigerated, penicillin solutions may be stored for seven days without significant loss of potency.

Buffered Pfizerpen for Injection may be given intramuscularly or by continuous intravenous infusion for dosages of 500,000, 1,000,000, or 5,000,000 units. It is also suitable for intrapleural, intraarticular, and other local instillations.

THE 20,000,000 UNIT DOSAGE MAY BE ADMINISTERED BY INTRAVENOUS INFUSION ONLY.

(1) Intramuscular Injection

Keep total volume of injection small. The intramuscular route is the preferred route of administration. Solutions containing up to 100,000 units of penicillin per mL of diluent may be used with a minimum of discomfort. Greater concentration of penicillin G per mL is physically possible and may be employed where therapy demands. When large dosages are required, it may be advisable to administer aqueous solutions of penicillin by means of continuous intravenous infusion.

(2) Continuous Intravenous Infusion

Determine the volume of fluid and rate of its administration required by the patient in a 24 hour period in the usual manner for fluid therapy, and add the appropriate daily dosage of penicillin to this fluid. For example, if an adult patient requires 2 liters of fluid in 24 hours and a daily dosage of 10 million units of penicillin, add 5 million units to 1 liter and adjust the rate of flow so the liter will be infused in 12 hours.

(3) Intrapleural or Other Local Infusion

If fluid is aspirated, give infusion in a volume equal to ¼ or ½ the amount of fluid aspirated, otherwise, prepare as for intramuscular injection.

(4) Intrathecal Use

The intrathecal use of penicillin in meningitis must be highly individualized. It should be employed only with full consideration of the possible irritating effects of penicillin when used by this route. The preferred route of therapy in bacterial meningitides is intravenous, supplemented by intramuscular injection.

Parenteral drug products should be inspected visually for particulate matter and discoloration prior to administration, whenever solution and container permit.

Sterile solution may be left in refrigerator for one week without significant loss of potency.

HOW SUPPLIED

Buffered Pfizerpen® (Penicillin G Potassium for Injection, USP) is available in vials containing respectively 5,000,000 units × 10's (NDC 0049-0520-22) and 20,000,000 units × 1's (NDC 0049-0530-22); buffered with sodium citrate and citric acid to an

optimum pH.

Each million units contains approximately 6.8 milligrams of sodium (0.3 mEq) and 65.6 milligrams of potassium (1.68 mEq).

Store the dry powder below 86°F (30°C).

This product's labeling may have been updated. For the most recent prescribing information, please visit www.pfizer.com.

For medical information about Buffered Pfizerpen, please visit www.pfizermedinfo.com or call 1-800-438-1985.

Rx only

Distributed by
Roerig
Division of Pfizer Inc,
New York, NY 10001



Novaplus is a registered trademark of Vizient, Inc.

LAB-0391-13.0
Revised: 06/2024

PRINCIPAL DISPLAY PANEL - 5,000,000 Unit Vial Label

NDC 0049-0520-23

Buffered

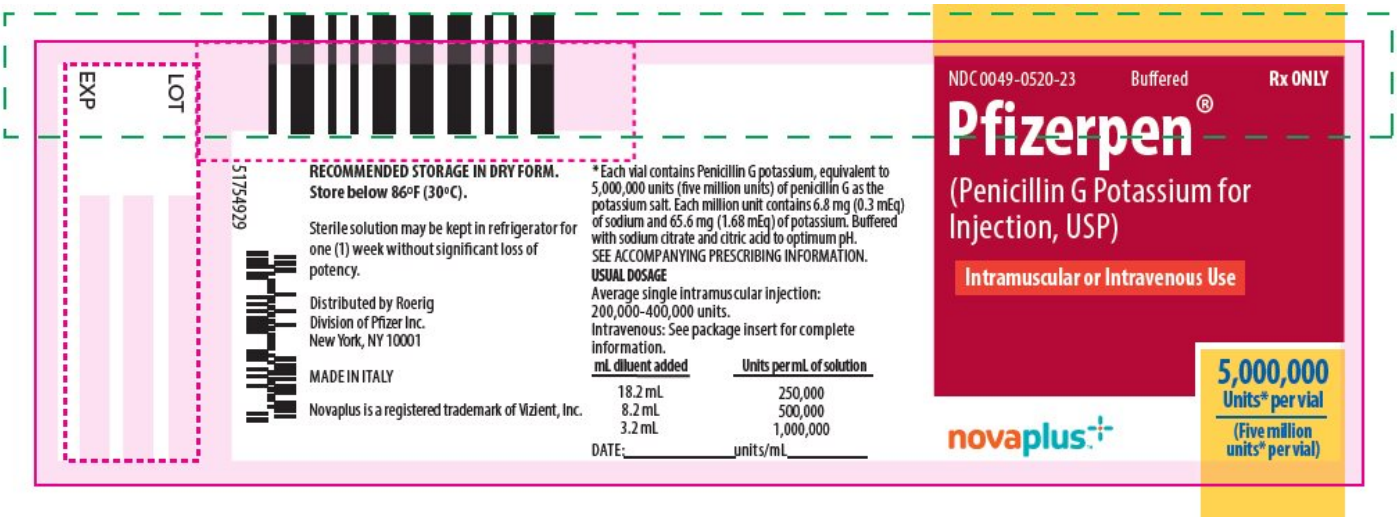
Rx ONLY

Pfizerpen®
(Penicillin G Potassium for
Injection, USP)

Intramuscular or Intravenous Use

novaplus™ +

5,000,000
Units* per vial
(Five million
units* per vial)



PRINCIPAL DISPLAY PANEL - 5,000,000 Unit Vial Carton

NDC 0049-0520-22
contains 10 of NDC 0049-0520-23

Buffered

Pfizerpen®
(Penicillin G potassium for Injection, USP)

5,000,000
Units* per vial
(Five million
units* per vial)

Intramuscular or Intravenous Use

10 Vials

Rx ONLY

novaplus™ +

S20x10K-INC

NDC 0049-0520-22
contains 10 of NDC 0049-0520-23

Buffered

Pfizerpen®

(Penicillin G Potassium for Injection, USP)

5,000,000 Units* per vial

(Five million units* per vial)

Intramuscular or Intravenous Use

10 Vials

Rx ONLY

novaplus

NDC 0049-0520-22
contains 10 of NDC 0049-0520-23

Buffered

Pfizerpen®

(Penicillin G Potassium for Injection, USP)

5,000,000 Units* per vial

(Five million units* per vial)

Intramuscular or Intravenous Use

novaplus

51754927

SEE ACCOMPANYING PRESCRIBING INFORMATION.

USUAL DOSAGE

Average single intramuscular injection: 200,000–400,000 units.

Intravenous: See package insert for complete information.

mL diluent added	Units per mL of solution
18.2 mL	250,000
8.2 mL	500,000
3.2 mL	1,000,000

RECOMMENDED STORAGE IN DRY FORM. Store below 86°F (30°C).

STERILE SOLUTION MAY BE KEPT IN REFRIGERATOR FOR ONE (1) WEEK WITHOUT SIGNIFICANT LOSS OF POTENCY.

Consult package insert for instructions for use of this vial.

GTIN: 00300490520225

Lot

EXP

* Each vial contains Penicillin G potassium, equivalent to 5,000,000 units (five million units) of penicillin G as the potassium salt. Each million unit contains 6.8 mg (0.3 mEq) of sodium and 65.6 mg (1.68 mEq) of potassium. Buffered with sodium citrate and citric acid to optimum pH. The pH of the reconstituted product is between 6.0 - 8.5.

Distributed by Roerig
Division of Pfizer Inc.
New York, NY 10001

MADE IN ITALY

Novaplus is a registered trademark of Vizient, Inc.

300490520225

PRINCIPAL DISPLAY PANEL - 20,000,000 Unit Vial Label

NDC 0049-0530-22

Buffered

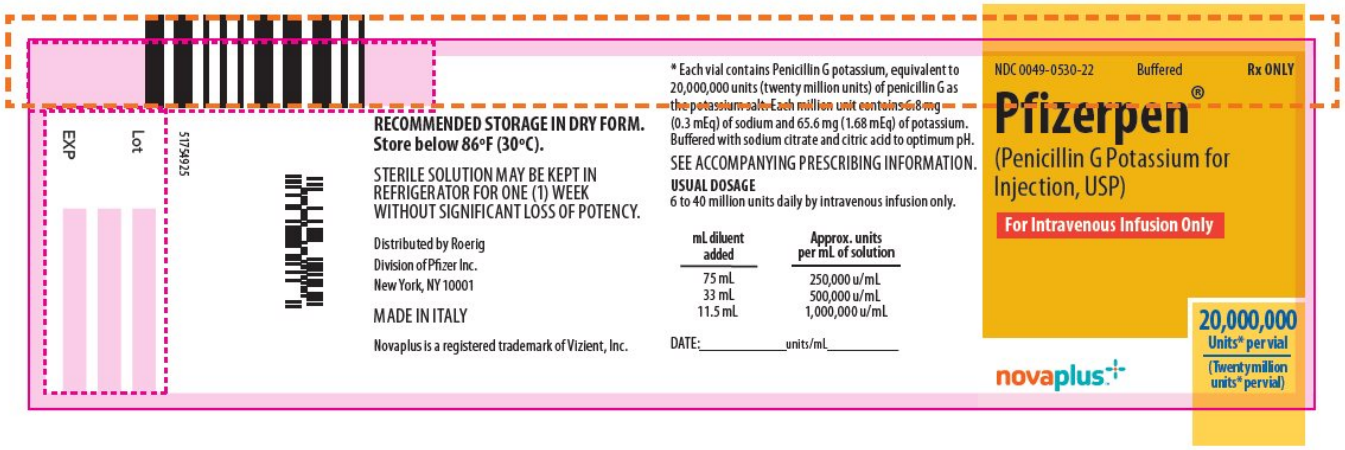
Rx ONLY

Pfizerpen®
(Penicillin G Potassium for
Injection, USP)

For Intravenous Infusion Only

novaplus™ +

**20,000,000
Units* per vial
(Twenty million
units* per vial)**



PRINCIPAL DISPLAY PANEL - 20,000,000 Unit Vial Carton

NDC 0049-0530-22

Buffered

Pfizerpen®
(Penicillin G Potassium for Injection, USP)

**20,000,000
Units* per vial
(Twenty million
units* per vial)**

For Intravenous Infusion Only

1 Vial

Rx ONLY

novaplus™ +



PFIZERPEN
penicillin g potassium powder, for solution

Product Information			
Product Type	HUMAN PRESCRIPTION DRUG	**Item Code (Source)**	NDC:0049-0520
Route of Administration	INTRAVENOUS, INTRAMUSCULAR, INTRAPLEURAL, INTRATHECAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
PENICILLIN G POTASSIUM (UNII: VL775ZTH4C) (PENICILLIN G - UNII:Q42T66VG0C)	PENICILLIN G	5000000 [iU]

Inactive Ingredients	
Ingredient Name	**Strength**
SODIUM POLYMETAPHOSPHATE (UNII: P1BM4ZH95L)	
SODIUM CITRATE, UNSPECIFIED FORM (UNII: 1Q73Q2JULR)	
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)	

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:0049-0520-22	10 in 1 CARTON	06/01/2010	
1		1 in 1 VIAL; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
ANDA	ANDA060657	06/01/2010	

PFIZERPEN

penicillin g potassium powder, for solution

Product Information			
Product Type	HUMAN PRESCRIPTION DRUG	**Item Code (Source)**	NDC:0049-0530
Route of Administration	INTRAVENOUS		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
PENICILLIN G POTASSIUM (UNII: VL775ZTH4C) (PENICILLIN G -		

PENICILLIN G POTASSIUM (UNII: VL7752TH4C) (PENICILLIN G - UNII:Q42T66VG0C)	PENICILLIN G	20000000 [iU]

Inactive Ingredients

Ingredient Name	Strength
SODIUM POLYMETAPHOSPHATE (UNII: P1BM4ZH95L)	
SODIUM CITRATE, UNSPECIFIED FORM (UNII: 1Q73Q2JULR)	
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:0049-0530-22	1 in 1 CARTON	06/01/2010	03/31/2026
1		1 in 1 VIAL; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
ANDA	ANDA060657	06/01/2010	03/31/2026

Labeler - Roerig (829076996)

Establishment

Name	Address	ID/FEI	Business Operations
Pharmacia & Upjohn Company LLC		618054084	ANALYSIS(0049-0520, 0049-0530)

Revised: 11/2026 Roerig