

MINOCIN- minocycline hydrochloride injection

Melinta Therapeutics, LLC

MINOCIN®
Minocycline For Injection
100 Mg/Vial

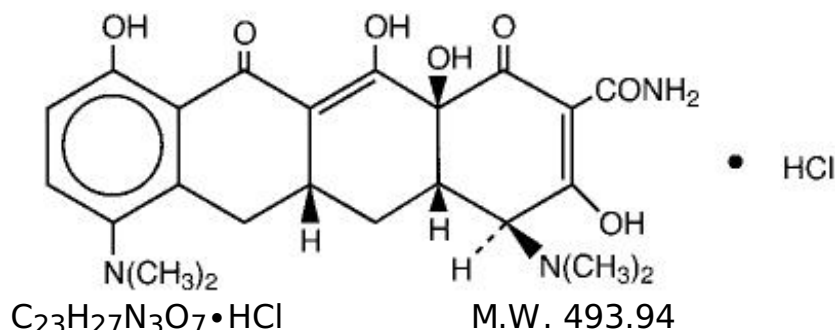
Rx Only

To reduce the development of drug-resistant bacteria and maintain the effectiveness of MINOCIN® (minocycline) for Injection and other antibacterial drugs, MINOCIN® (minocycline) for injection should be used only to treat or prevent infections that are proven or strongly suspected to be caused by bacteria.

DESCRIPTION

MINOCIN (minocycline) for Injection, is a sterile formulation of a semisynthetic derivative of tetracycline. The chemical name of minocycline is 4,7-Bis(dimethylamino)-1,4,4a,5,5a,6,11,12a-octahydro-3,10,12,12a-tetrahydroxy-1,11-dioxo-2-naphthacenecarboxamide monohydrochloride.

Its structural formula is:



MINOCIN is supplied as a sterile yellow to amber lyophilized powder for intravenous infusion. Each vial contains minocycline HCl equivalent to 100 mg minocycline, 269 mg magnesium sulfate heptahydrate (2.2 mEq of magnesium) (an inactive ingredient) and sodium hydroxide (to adjust pH). When reconstituted with 5 mL of Sterile Water for Injection USP the pH ranges from 4.5 to 5.0.

CLINICAL PHARMACOLOGY

Following a single dose of Minocin 200 mg administered intravenously to 10 healthy male subjects, serum concentrations of minocycline ranged from 2.52 to 6.63 mcg/mL (average 4.18 mcg/mL) at the end of infusion and 0.82 to 2.64 mcg/mL (average 1.38 mcg/mL) after 12 hours. In a group of 5 healthy male subjects, serum concentrations of minocycline ranged from 1.4 to 1.8 mcg/mL at the end of the dosing interval following administration of Minocin 100 mg every 12 hours for three days. When Minocin 200 mg once daily was administered for three days, serum concentrations of minocycline were approximately 1 mcg/mL at 24 hours. The serum elimination half-life of minocycline

following administration of either Minocin 100 mg every 12 hours or 200 mg once daily was not significantly different and ranged from 15 to 23 hours. The serum elimination half-life of minocycline ranged from 11 to 16 hours in subjects with hepatic impairment (n=7) and 18 to 69 hours in subjects with renal impairment (n=5). In comparison, the serum elimination half-life of minocycline ranged from 11 to 17 hours following a single dose of oral minocycline 200 mg in healthy subjects (n=12).

Microbiology

Mechanism of Action

The tetracyclines are primarily bacteriostatic and are thought to exert their antimicrobial effect by the inhibition of protein synthesis. The tetracyclines, including minocycline, have a similar antimicrobial spectrum of activity against a wide range of Gram-positive and Gram-negative bacteria. Cross-resistance of these bacteria to tetracyclines is common.

List of Microorganisms

Minocycline has been shown to be active against most isolates of the following bacteria, both *in vitro* and in clinical infections as described in the **INDICATIONS AND USAGE** section:

Gram-positive Bacteria

Bacillus anthracis
Listeria monocytogenes
Staphylococcus aureus
Streptococcus pneumoniae

Gram-negative Bacteria

Bartonella bacilliformis
Brucella species
Klebsiella granulomatis
Campylobacter fetus
Francisella tularensis
Vibrio cholerae
Yersinia pestis
Acinetobacter species
Enterobacter aerogenes
Escherichia coli
Haemophilus influenzae
Klebsiella species
Neisseria meningitidis
Shigella species

Other Microorganisms

Actinomyces species
Borrelia recurrentis
Chlamydophila psittaci
Chlamydia trachomatis
Clostridium species
Entamoeba species

Fusobacterium nucleatum subspecies fusiforme
Mycobacterium marinum
Mycoplasma pneumoniae
Propionibacterium acnes
Rickettsiae
Treponema pallidum subspecies pallidum
Treponema pallidum subspecies pertenue
Ureaplasma urealyticum

Susceptibility Test Methods

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

MINOCIN® Intravenous is indicated in the treatment of the following infections due to susceptible isolates of the designated bacteria:

Rocky Mountain spotted fever, typhus fever and the typhus group, Q fever, rickettsialpox and tick fevers caused by *rickettsiae*.
Respiratory tract infections caused by *Mycoplasma pneumoniae*.
Lymphogranuloma venereum caused by *Chlamydia trachomatis*.
Psittacosis (Ornithosis) due to *Chlamydophila psittaci*.
Trachoma caused by *Chlamydia trachomatis*, although the infectious agent is not always eliminated, as judged by immunofluorescence.
Inclusion conjunctivitis caused by *Chlamydia trachomatis*.
Nongonococcal urethritis, endocervical, or rectal infections in adults caused by *Ureaplasma urealyticum* or *Chlamydia trachomatis*.
Relapsing fever due to *Borrelia recurrentis*.
Plague due to *Yersinia pestis*.
Tularemia due to *Francisella tularensis*.
Cholera caused by *Vibrio cholerae*.
Campylobacter fetus infections caused by *Campylobacter fetus*.
Brucellosis due to *Brucella species* (in conjunction with streptomycin).
Bartonellosis due to *Bartonella bacilliformis*.
Granuloma inguinale caused by *Klebsiella granulomatis*.

Minocycline is indicated for the treatment of infections caused by the following Gram-negative bacteria when bacteriologic testing indicates appropriate susceptibility to the drug:

Escherichia coli.
Enterobacter aerogenes.
Shigella species.
Acinetobacter species.
Respiratory tract infections caused by *Haemophilus influenzae*.
Respiratory tract and urinary tract infections caused by *Klebsiella species*.

MINOCIN® Intravenous is indicated for the treatment of infections caused by the following Gram-positive bacteria when bacteriologic testing indicates appropriate

susceptibility to the drug:

Upper respiratory tract infections caused by *Streptococcus pneumoniae*.

Skin and skin structure infections caused by *Staphylococcus aureus* (Note: Minocycline is not the drug of choice in the treatment of any type of staphylococcal infection.)

When penicillin is contraindicated, minocycline is an alternative drug in the treatment of the following infections:

Meningitis due to *Neisseria meningitidis*.

Syphilis caused by *Treponema pallidum subspecies pallidum*.

Yaws caused by *Treponema pallidum subspecies pertenue*.

Listeriosis due to *Listeria monocytogenes*.

Anthrax due to *Bacillus anthracis*.

Vincent's infection caused by *Fusobacterium fusiforme*.

Actinomycosis caused by *Actinomyces israelii*.

Infections caused by *Clostridium species*.

In *acute intestinal amebiasis*, minocycline may be a useful adjunct to amebicides.

In severe *acne*, minocycline may be useful adjunctive therapy.

To reduce the development of drug-resistant bacteria and maintain the effectiveness of MINOCIN® (minocycline) for Injection and other antibacterial drugs, MINOCIN® (minocycline) for Injection 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

This drug is contraindicated in persons who have shown hypersensitivity to any of the tetracyclines or to any of the components of the product formulation.

WARNINGS

Tooth Development

MINOCIN, like other tetracycline-class antibacterials, can cause fetal harm when administered to a pregnant woman. If any tetracycline is used during pregnancy, or if the patient becomes pregnant while taking these drugs, the patient should be apprised of the potential hazard to the fetus. The use of drugs of the tetracycline class during tooth development (last half of pregnancy, infancy, and childhood to the age of 8 years) may cause permanent discoloration of the teeth (yellow-gray-brown).

This adverse reaction is more common during long-term use of the drugs but has been observed following repeated short-term courses. Enamel hypoplasia has also been reported. Tetracycline drugs, therefore, should not be used during tooth development unless other drugs are not likely to be effective or are contraindicated.

Skeletal Development

All tetracyclines form a stable calcium complex in any bone-forming tissue. A decrease in the fibula growth rate has been observed in premature human infants given oral tetracycline in doses of 25 mg/kg every six hours. This reaction was shown to be reversible when the drug was discontinued.

Results of animal studies indicate that tetracyclines cross the placenta, are found in fetal tissues, and can have toxic effects on the developing fetus (often related to retardation of skeletal development). Evidence of embryotoxicity has been noted in animals treated early in pregnancy.

Dermatologic Reaction

Drug Rash with Eosinophilia and Systemic Symptoms (DRESS) including fatal cases have been reported with minocycline use. Fixed drug eruptions have occurred with minocycline and other tetracyclines. Worsening severity upon subsequent administrations, including generalized bullous fixed drug eruption, has been observed with other tetracyclines (see **ADVERSE REACTIONS**). If severe skin reactions occur, discontinue MINOCIN immediately and institute appropriate therapy.

Anti-anabolic Action

The anti-anabolic action of the tetracyclines may cause an increase in BUN. While this is not a problem in those with normal renal function, in patients with significantly impaired function, higher serum levels of tetracycline may lead to azotemia, hyperphosphatemia, and acidosis. Under such conditions, monitoring of creatinine and BUN is recommended, and the total daily dosage should not exceed 200 mg in 24 hours (See **DOSAGE AND ADMINISTRATION**). If renal impairment exists, even usual oral or parenteral doses may lead to systemic accumulation of the drug and possible liver toxicity.

Photosensitivity

Photosensitivity manifested by an exaggerated sunburn reaction has been observed in some individuals taking tetracyclines. This has been reported with minocycline.

Central Nervous System Effects

Central nervous system side effects including light-headedness, dizziness or vertigo have been reported. Patients who experience these symptoms should be cautioned about driving vehicles or using hazardous machinery while on minocycline therapy. These symptoms may disappear during therapy and usually disappear rapidly when the drug is discontinued.

***Clostridium difficile* Associated Diarrhea**

Clostridium difficile associated diarrhea (CDAD) has been reported with use of nearly all antibacterial agents, including MINOCIN[®], 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 antibacterial

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 antibacterial use not directed against *C. difficile* may need to be discontinued. Appropriate fluid and electrolyte management, protein supplementation, antibacterial treatment of *C. difficile*, and surgical evaluation should be instituted as clinically indicated.

Intracranial Hypertension

Intracranial hypertension (IH, pseudotumor cerebri) has been associated with the use of tetracyclines including Minocin. Clinical manifestations of IH include headache, blurred vision, diplopia, and vision loss; papilledema can be found on fundoscopy. Women of childbearing age who are overweight or have a history of IH are at greater risk for developing tetracycline associated IH. Concomitant use of isotretinoin and Minocin should be avoided because isotretinoin is also known to cause pseudotumor cerebri.

Although IH typically resolves after discontinuation of treatment, the possibility for permanent visual loss exists. If visual disturbance occurs during treatment, prompt ophthalmologic evaluation is warranted. Since intracranial pressure can remain elevated for weeks after drug cessation patients should be monitored until they stabilize.

PRECAUTIONS

General

As with other antibacterial preparations, use of this drug may result in overgrowth of nonsusceptible organisms, including fungi. If superinfection occurs, the antibacterial should be discontinued and appropriate therapy instituted.

Hepatotoxicity has been reported with minocycline; therefore, minocycline should be used with caution in patients with hepatic dysfunction and in conjunction with other hepatotoxic drugs.

Incision and drainage or other surgical procedures should be performed in conjunction with antibacterial therapy when indicated.

MINOCIN[®] (minocycline) for Injection contains magnesium sulfate heptahydrate (See **DESCRIPTION**). Because magnesium is excreted primarily by the kidney, serum levels of magnesium should be monitored in patients with renal impairment (See **DOSAGE AND ADMINISTRATION**).

Because MINOCIN[®] (minocycline) for Injection contains magnesium (See **DESCRIPTION**), close monitoring is recommended in patients with heart block or myocardial damage.

Prescribing MINOCIN[®] for Injection 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.

Information for Patients

Patients should be counseled that antibacterial drugs including MINOCIN[®] (minocycline) for Injection should only be used to treat bacterial infections. They do not treat viral

infections (e.g., the common cold). When MINOCIN[®] (minocycline) 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 MINOCIN[®] (minocycline) for Injection or other antibacterial drugs in the future.

Diarrhea is a common problem caused by antibacterials which usually ends when the antibacterial is discontinued. Sometimes after starting treatment with antibacterial drugs, 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 antibacterial. If this occurs, patients should contact their physician as soon as possible.

Laboratory Tests

Periodic laboratory evaluation of organ systems, including hematopoietic, renal, and hepatic studies should be performed.

All patients with gonorrhea should have a serologic test for syphilis at the time of diagnosis. Patients treated with minocycline should have a follow-up serologic test for syphilis after 3 months.

Drug Interactions

Because tetracyclines have been shown to depress plasma prothrombin activity, patients who are on anticoagulant therapy may require downward adjustment of their anticoagulant dosage.

Since bacteriostatic drugs may interfere with the bactericidal action of penicillin, it is advisable to avoid giving tetracyclines in conjunction with penicillin.

Concurrent use of tetracyclines with oral contraceptives may render oral contraceptives less effective.

Administration of isotretinoin should be avoided shortly before, during, and shortly after minocycline therapy. Each drug alone has been associated with pseudotumor cerebri (See **PRECAUTIONS**).

Increased risk of ergotism when ergot alkaloids or their derivatives are given with tetracyclines.

MINOCIN[®] (minocycline) for Injection contains magnesium sulfate heptahydrate (See **DESCRIPTION**). Potentially serious drug interactions may occur when intravenous magnesium sulfate heptahydrate is given concomitantly with CNS depressants, neuromuscular blocking agents and cardiac glycosides.

Drug/Laboratory Test Interactions

False elevations of urinary catecholamine levels may occur due to interference with the fluorescence test.

Carcinogenesis, Mutagenesis, Impairment of Fertility

Dietary administration of minocycline in long-term tumorigenicity studies in rats resulted

in evidence of thyroid tumor production. Minocycline has also been found to produce thyroid hyperplasia in rats and dogs. In addition, there has been evidence of oncogenic activity in rats in studies with a related antibacterial, oxytetracycline (i.e., adrenal and pituitary tumors). Likewise, although mutagenicity studies of minocycline have not been conducted, positive results in *in vitro* mammalian cell assays (i.e., mouse lymphoma and Chinese hamster lung cells) have been reported for related antibacterials (tetracycline hydrochloride and oxytetracycline). Segment I (fertility and general reproduction) studies have provided evidence that minocycline impairs fertility in male rats.

Pregnancy

Teratogenic Effects

(See **WARNINGS**)

All pregnancies have a background risk of birth defects, loss, or other adverse outcome regardless of drug exposure. There are no adequate and well-controlled studies on the use of minocycline in pregnant women. Minocycline, like other tetracycline-class antibacterials, crosses the placenta and may cause fetal harm when administered to a pregnant woman. Rare spontaneous reports of congenital anomalies including limb reduction have been reported in post-marketing experience. Only limited information is available regarding these reports; therefore, no conclusion on causal association can be established. If minocycline is used during pregnancy or if the patient becomes pregnant while taking this drug, the patient should be apprised of the potential hazard to the fetus.

Nonteratogenic Effects

(See **WARNINGS**)

Labor and Delivery

The effect of tetracyclines on labor and delivery is unknown.

Nursing Mothers

Tetracyclines are excreted in human milk. Because of the potential for serious adverse reactions in nursing infants from tetracyclines, a decision should be made whether to discontinue nursing or discontinue the drug, taking into account the importance of the drug to the mother. (See **WARNINGS**).

Pediatric Use

Minocycline is not recommended for use in children below 8 years of age unless the expected benefits of therapy outweigh the risks (See **WARNINGS**).

Geriatric Use

Clinical studies of intravenous minocycline did not include sufficient numbers of subjects aged 65 and over to determine whether they respond differently from younger subjects. 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 (See

WARNINGS, DOSAGE AND ADMINISTRATION).

ADVERSE REACTIONS

The following adverse reactions have been observed in patients receiving tetracyclines.

Body as a whole: Fever, and discoloration of secretions.

Gastrointestinal: Anorexia, nausea, vomiting, diarrhea, dyspepsia, stomatitis, glossitis, dysphagia, enamel hypoplasia, enterocolitis, pseudomembranous colitis, pancreatitis, inflammatory lesions (with monilial overgrowth) in the oral and anogenital regions. These reactions have been caused by both the oral and parenteral administration of tetracyclines.

Genitourinary: Vulvovaginitis.

Hepatic toxicity: Hyperbilirubinemia, hepatic cholestasis, increases in liver enzymes, fatal hepatic failure, and jaundice. Hepatitis, including autoimmune hepatitis, and liver failure have been reported (See **PRECAUTIONS**).

Skin: Alopecia, erythema nodosum, hyperpigmentation of nails, pruritus, toxic epidermal necrolysis, and vasculitis, maculopapular and erythematous rashes. Exfoliative dermatitis has been reported. Fixed drug eruptions have been reported. Lesions occurring on the glans penis have caused balanitis. Erythema multiforme and Stevens-Johnson syndrome have been reported. Photosensitivity is discussed above (See **WARNINGS**). Pigmentation of the skin and mucous membranes has been reported.

Local Reactions: Injection site erythema and injection site pain.

Respiratory: Cough, dyspnea, bronchospasm, exacerbation of asthma, and pneumonitis.

Renal toxicity: Interstitial nephritis. Elevations in BUN have been reported and are apparently dose related (See **WARNINGS**). Acute renal failure has been reported.

Musculoskeletal: Arthralgia, arthritis, bone discoloration, myalgia, joint stiffness, and joint swelling.

Hypersensitivity reactions: Urticaria, angioneurotic edema, polyarthralgia, anaphylaxis/anaphylactoid reaction (including shock and fatalities), anaphylactoid purpura, myocarditis, pericarditis, exacerbation of systemic lupus erythematosus, and pulmonary infiltrates with eosinophilia have been reported. A lupus-like syndrome and serum sickness-like reactions also have been reported.

Blood: Agranulocytosis, hemolytic anemia, thrombocytopenia, leukopenia, neutropenia, pancytopenia, and eosinophilia have been reported.

Central Nervous System: Convulsions, dizziness, hypesthesia, paresthesia, sedation, and vertigo. Pseudotumor cerebri (benign intracranial hypertension) in adults and bulging fontanels in infants (See **PRECAUTIONS - General**). Headache has also been reported.

Other: Thyroid cancer has been reported in the post-marketing setting in association with minocycline products. When minocycline therapy is given over prolonged periods, monitoring for signs of thyroid cancer should be considered. When given over prolonged periods, tetracyclines have been reported to produce brown-black microscopic discoloration of the thyroid gland. Cases of abnormal thyroid function have

been reported.

Tooth discoloration in pediatric patients less than 8 years of age (See **WARNINGS**) and in adults has been reported.

Oral cavity discoloration (including tongue, lip, and gum) have been reported.

Tinnitus and decreased hearing have been reported in patients on MINOCIN® (minocycline for injection).

The following syndromes have been reported. In some cases involving these syndromes, death has been reported. As with other serious adverse reactions, if any of these syndromes are recognized, the drug should be discontinued immediately:

Hypersensitivity syndrome consisting of cutaneous reaction (such as rash or exfoliative dermatitis), eosinophilia, and one or more of the following: hepatitis, pneumonitis, nephritis, myocarditis, and pericarditis. Fever and lymphadenopathy may be present.

Lupus-like syndrome consisting of positive antinuclear antibody; arthralgia, arthritis, joint stiffness, or joint swelling; and one or more of the following: fever, myalgia, hepatitis, rash, and vasculitis.

Serum sickness-like syndrome consisting of fever; urticaria or rash; and arthralgia, arthritis, joint stiffness, or joint swelling. Eosinophilia may be present.

MINOCIN® (minocycline) for Injection contains magnesium sulfate heptahydrate (See **DESCRIPTION**). Adverse effects that may be associated with magnesium intoxication include flushing, sweating, hypotension, depressed reflexes, flaccid paralysis, hypothermia, circulatory collapse, cardiac and CNS depression proceeding to respiratory paralysis (See **PRECAUTIONS**).

OVERDOSAGE

The adverse events more commonly seen in overdose are dizziness, nausea, and vomiting.

No specific antidote for minocycline is known.

In case of overdosage, discontinue medication, treat symptomatically, and institute supportive measures. Minocycline is not removed in significant quantities by hemodialysis or peritoneal dialysis.

DOSAGE AND ADMINISTRATION

THE USUAL DOSAGE AND FREQUENCY OF ADMINISTRATION OF MINOCYCLINE DIFFERS FROM THAT OF THE OTHER TETRACYCLINES. EXCEEDING THE RECOMMENDED DOSAGE MAY RESULT IN AN INCREASED INCIDENCE OF SIDE EFFECTS.

Note: Rapid administration is to be avoided. Parenteral therapy is indicated only when oral therapy is not adequate or tolerated. Oral therapy should be instituted as soon as possible. If intravenous therapy is given over prolonged periods of time, thrombophlebitis may result.

For Pediatric Patients above 8 years of Age

Usual pediatric dose: Initial dose of 4 mg/kg, then 2 mg/kg administered over 60 minutes every 12 hours, not to exceed the usual adult dose.

Adults

Usual adult dose: Initial dose of 200 mg, then 100 mg administered over 60 minutes every 12 hours and should not exceed 400 mg in 24 hours.

The lyophilized powder should be reconstituted with 5 mL Sterile Water for Injection USP and immediately further diluted in 100 mL to 1,000 mL with Sodium Chloride Injection USP, Dextrose Injection USP, or Dextrose and Sodium Chloride Injection USP, or in 250 mL to 1000 mL Lactated Ringer's Injection USP, but not with other solutions containing calcium because a precipitate may form especially in neutral and alkaline solutions.

When diluted in compatible solutions, the pH usually ranges from 4.5 to 6.0.

Once diluted into an intravenous bag, MINOCIN[®] for Injection may be stored either at room temperature for up to 4 hours or refrigerated at 2 to 8°C (36 to 46°F) for up to 24 hours. Any unused portions must be discarded after that period.

The pharmacokinetics of minocycline in patients with renal impairment ($CL_{CR} < 80$ mL/min) have not been fully characterized. Current data are insufficient to determine if a dosage adjustment is warranted. The total daily dosage should not exceed 200 mg in 24 hours in patients with renal impairment. However, due to the anti-anabolic effect of tetracyclines, BUN and creatinine should be monitored (See **WARNINGS**). Because MINOCIN (minocycline) for Injection contains magnesium sulfate heptahydrate, serum levels of magnesium should be monitored in patients with renal impairment (See **DESCRIPTION, PRECAUTIONS**).

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

Incompatibilities

Additives or other medications should not be added to MINOCIN single-dose vials or infused simultaneously through the same intravenous line including Y-connectors. If the same intravenous line is used for sequential infusion of additional medications, the line should be flushed before and after infusion of MINOCIN with Sodium Chloride Injection USP, Dextrose Injection USP, Dextrose and Sodium Chloride Injection USP, or Lactated Ringer's Injection USP.

HOW SUPPLIED

MINOCIN[®] (minocycline) for Injection is supplied as 100 mg single-dose vials of sterile lyophilized powder.

NDC 70842-160-01: 100 mg single-dose vial

NDC 70842-160-10: units of 10 × 1 single-dose 100 mg vials

Store at Controlled Room Temperature 20° to 25°C (68° to 77°F).

ANIMAL PHARMACOLOGY AND TOXICOLOGY

Minocycline hydrochloride has been observed to cause a dark discoloration of the thyroid in experimental animals (rats, minipigs, dogs, and monkeys). In the rat, chronic treatment with minocycline hydrochloride has resulted in goiter accompanied by elevated radioactive iodine uptake and evidence of thyroid tumor production. Minocycline hydrochloride has also been found to produce thyroid hyperplasia in rats and dogs.

This product's label may have been updated. For current package insert and further product information, please visit www.minociniv.com or call our toll-free number: 1-844-633-6568. Call between 9:00 a.m. and 5:00 p.m. Eastern Time, Monday through Friday.

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Melinta Therapeutics, LLC
Parsippany, New Jersey 07054

3/2025

MEL080-R004

PRINCIPAL DISPLAY PANEL - 100 mg Vial Carton

Rx only
NDC 70842-160-10

10 Vials

Minocin®
(minocycline)
for injection

100 mg per vial

Must be reconstituted
then diluted.
For Intravenous Infusion.
Sterile Single-Dose Vials.
Discard Unused Portion.

Each vial contains minocycline hydrochloride (equivalent to 100 mg minocycline), magnesium sulfate heptahydrate and sodium hydroxide (to adjust pH 4.5-5.0).

Usual adult dose: Initial dose of 200 mg, then 100 mg every 12 hours and should not exceed 400 mg in 24 hours.

Reconstitution: Reconstitute each vial with 5 mL of Sterile Water for Injection.

Dilution: Reconstituted solution must be further diluted in 100 mL to 1,000 mL for infusion.

See package insert for detailed instructions for diluting,

indications and dosage.

Rx only NDC 70842-160-10
10 Vials

**Minocin®
(minocycline)
for injection**

100 mg per vial

Must be reconstituted
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For Intravenous Infusion.
Sterile Single-Dose Vials.
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See package insert for detailed instructions for diluting, indications and dosage.

Rx only NDC 70842-160-10

**Minocin®
(minocycline)
for injection**

100 mg per vial



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Area

Glue



melinta
therapeutics

Marketed by:
Melinta Therapeutics, LLC
Lincolnshire, IL 60069 USA

Rx only NDC 70842-160-10
**Minocin®
(minocycline)
for injection**
100 mg per vial
Must be reconstituted
then diluted.
For Intravenous Infusion.
Sterile Single-Dose Vials.
Discard Unused Portion.

70015668

Rx only NDC 70842-160-10

**Minocin®
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100 mg per vial

Must be reconstituted
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Discard Unused Portion.

Reconstitution: Reconstitute each vial with 5 mL of Sterile Water for Injection.

Dilution: Reconstituted solution must be further diluted in 100 mL to 1,000 mL for infusion.

See package insert for detailed instructions for diluting, indications and dosage.

Store at controlled room temperature:
20°C to 25°C (68°F to 77°F)

Manufactured for:
Rempec Pharmaceuticals, Inc., a wholly owned subsidiary
of Melinta Therapeutics, LLC, Lincolnshire, IL 60069 USA

Marketed by Melinta Therapeutics, LLC
Product of Italy

Rx only NDC 70842-160-10

**Minocin®
(minocycline)
for injection**

100 mg per vial

MEL082-R003

Glue

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MINOCIN

minocycline hydrochloride injection

Product Information

Product Type	HUMAN PRESCRIPTION DRUG	Item Code (Source)	NDC:70842-160
Route of Administration	INTRAVENOUS		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
MINOCYCLINE HYDROCHLORIDE (UNII: 0020414E5U) (MINOCYCLINE - UNII:FYY3R43WGO)	MINOCYCLINE	100 mg

Inactive Ingredients

Ingredient Name	Strength
MAGNESIUM SULFATE HEPTAHYDRATE (UNII: SK47B8698T)	269 mg
SODIUM HYDROXIDE (UNII: 55X04QC32I)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:70842-160-10	10 in 1 CARTON	04/19/2013	
1	NDC:70842-160-01	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
NDA	NDA050444	04/19/2013	

Labeler - Melinta Therapeutics, LLC (034132850)

Registrant - Patheon Italia SpA (338336589)

Establishment

Name	Address	ID/FEI	Business Operations
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Patheon Italia SpA		338336589	ANALYSIS(70842-160) , MANUFACTURE(70842-160) , STERILIZE(70842-160) , PACK(70842-160)
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Establishment

Name	Address	ID/FEI	Business Operations
AndersonBrecon Inc.		053217022	PACK(70842-160) , LABEL(70842-160)

Establishment

Name	Address	ID/FEI	Business Operations
Redox Srl		439198391	ANALYSIS(70842-160)

Establishment

Name	Address	ID/FEI	Business Operations
Curia Italy S.r.l		440365766	ANALYSIS(70842-160) , API MANUFACTURE(70842-160)

Establishment

Name	Address	ID/FEI	Business Operations
IDT Australia Limited		743859243	ANALYSIS(70842-160) , API MANUFACTURE(70842-160)

Establishment

Name	Address	ID/FEI	Business Operations
Pharma Packaging Solutions, LLC dba Tjoapack, LLC		928861723	PACK(70842-160) , LABEL(70842-160)

Establishment

Name	Address	ID/FEI	Business Operations
Eurofins Biolab Srl		429117112	ANALYSIS(70842-160)

Revised: 4/2025

Melinta Therapeutics, LLC