

Prescribing Information

Pr **ERYTHROCIN® I.V.**

(erythromycin lactobionate for injection)

500 mg and 1 g erythromycin/vial

For i.v. use only

Antibiotic

Amdipharm Limited,
Temple Chamber, 3 Burlington Road,
Dublin, Dublin 4, Ireland

DATE OF REVISION:
February 11, 2010

Distributed by: Methapharm Inc.
Brantford, Ontario, N3S 7X6

Submission Control Number: 129342

PRESCRIBING INFORMATION

NAME OF DRUG

Pr **ERYTHROCIN® I.V.**
(erythromycin lactobionate for injection)
500 mg and 1 g erythromycin/vial

THERAPEUTIC CLASSIFICATION

Antibiotic

ACTION AND CLINICAL PHARMACOLOGY

Erythromycin exerts its antibacterial action by binding to the 50S ribosomal subunit of susceptible bacteria and suppressing protein synthesis. Erythromycin is usually bacteriostatic but may be bactericidal in high concentrations or against highly susceptible organisms.

Intravenous infusion of 500 mg erythromycin lactobionate at a constant rate over 1 hour in fasting adults produced a mean serum erythromycin level of approximately 7 µg/mL at 20 minutes, 10 µg/mL at 1 hour, 2.6 µg/mL at 2.5 hours, and 1 µg/mL at 6 hours.

Erythromycin diffuses readily into most body fluids. Only low concentrations are normally achieved in the spinal fluid, but passage of the drug across the blood-brain barrier increases in meningitis.

Erythromycin is largely bound to plasma proteins (over 70%).

The half-life ($t_{1/2}$) for erythromycin is approximately 2 hours.

In the presence of normal hepatic function, erythromycin is concentrated in the liver and is excreted in the bile; the effect of hepatic dysfunction on biliary excretion of erythromycin is not known. From 12 to 15 percent of intravenously administered erythromycin is excreted in active form in the urine.

MICROBIOLOGY

Many strains of *Hemophilus influenzae* are resistant to erythromycin alone.

Staphylococci resistant to erythromycin may emerge during a course of erythromycin therapy. Culture and sensitivity testing should be performed prior to and during therapy.

Erythromycin is usually bacteriostatic but may be bactericidal in high concentrations or against highly susceptible organisms. The bactericidal activity is greatest against a small number of rapidly dividing microorganisms and increases markedly as the pH of the medium is raised over the range of pH 5.5 to 8.5.

Biochemical tests demonstrate that erythromycin inhibits protein synthesis of the pathogen without directly affecting nucleic acid synthesis. Antagonism has been demonstrated between clindamycin, lincomycin and chloramphenicol and erythromycin.

Susceptibility Testing

Disc Susceptibility Tests: Quantitative methods that require measurement of zone diameters give the most precise estimates of antibiotic susceptibility. One recommended procedure uses erythromycin class discs for testing susceptibility; interpretations correlate zone diameters of this disc test with MIC values for erythromycin. With this procedure, a report from the laboratory of "susceptible" indicates that the infecting organism is likely to respond to therapy. A report of "resistant" indicates that the infecting organism is not likely to respond to therapy. A report of "intermediate susceptibility" suggests that the organism would be susceptible if higher doses were used.

NOTE: Many strains of *Hemophilus influenzae* are resistant to erythromycin alone, but are susceptible to erythromycin and sulfonamides together. Staphylococci resistant to erythromycin may emerge during a course of erythromycin therapy. Culture and susceptibility testing should be performed.

The standard single disc susceptibility test (using the 15 µg erythromycin disc) and the dilution susceptibility test should be interpreted according to the criteria in Table 1.

Table 1 Criteria for Interpreting Standard Single Disc Susceptibility Test and The Dilution Susceptibility Test		
	Zone Diameter (mm)	Approximate MIC Correlate (mg/L)
Susceptible	≥ 23	≤ 0.5
Intermediate*	14-22	1-4
Resistant	≤ 13	≥ 8

* Indicates that the test results are equivocal; therefore, dilution tests may be indicated.
N.B.: These criteria and the definition are in agreement with NCCLS Order Code M2A3.

Control limits for monitoring erythromycin susceptibility tests are given in Table 2.

Table 2 Control Limits for Monitoring Erythromycin Susceptibility Tests		
	Zone Diameter (mm)	MIC (mg/L)
<i>S. aureus</i> ATCC 29213	22-30	0.12 - 0.50
<i>S. faecalis</i> ATCC 29212		1.0 - 4.0

INDICATIONS AND CLINICAL USES

ERYTHROCIN[®] I.V. - (erythromycin lactobionate for injection) should be used in the treatment of patients when oral administration is not possible or when it is desirable to obtain higher serum levels of erythromycin than achievable with orally administered preparations. Intravenous erythromycin should be replaced by an oral form of erythromycin as soon as possible.

Erythromycin is indicated for the treatment of infections caused by susceptible strains of the designated microorganisms in the diseases listed below:

1. **Lower respiratory tract infections** of mild to moderate severity caused by *S. pyogenes* (Group A beta-hemolytic streptococci), *S. pneumoniae* and *M. pneumoniae*.
2. **Skin and soft tissue infections** of mild to moderate severity caused by *S. pyogenes* and *S. aureus*.

N.B. Resistance of staphylococci may emerge during treatment.

3. **Legionnaires' disease** caused by *L. pneumophila*. Although no controlled clinical efficacy studies have been conducted, *in vitro* and limited clinical data suggest that erythromycin can be effective in treating Legionnaires' disease. Clinical evidence suggests that erythromycin is the preferred antibiotic for treating Legionnaires' Diseases.
4. Erythromycin should not be used for the treatment of syphilis in pregnancy because it cannot be relied upon to cure an infected fetus (see **PRECAUTIONS, Pregnancy**).

Specimens for bacteriologic culture should be obtained prior to therapy in order to isolate and identify the causative organisms and to determine their susceptibility to erythromycin. Therapy may be instituted before results of susceptibility studies are known; however, antibiotic treatment should be re-evaluated when the results become available or if the clinical response is not adequate.

CONTRAINDICATIONS

ERYTHROCIN[®] I.V. - (erythromycin lactobionate for injection) is contraindicated in patients with known hypersensitivity to erythromycin, clarithromycin or other macrolide antibacterial agents. Erythromycin is also contraindicated as concurrent therapy with astemizole*, terfenadine*, cisapride*, pimozone*, and ergotamine or dihydroergotamine (see **PRECAUTIONS: Drug Interactions**).

ERYTHROCIN[®] I.V. must be administered by continuous or intermittent intravenous infusion only. I.V. bolus/push is an unacceptable route of administration.

WARNINGS

Erythromycin should be administered with caution to any patient who has demonstrated some form of allergy to drugs. If an allergic reaction to erythromycin occurs, administration of the drug

* Astemizole, terfenadine, cisapride and pimozone are no longer marketed in Canada.

should be discontinued. Serious hypersensitivity reactions may require epinephrine, antihistamines, or corticosteroids.

Hepatic dysfunction, including increased liver enzymes and hepatocellular and/or cholestatic hepatitis, with or without jaundice, has been infrequently reported with erythromycin. If findings suggestive of significant hepatic dysfunction occur, therapy with erythromycin products should be discontinued.

There have been reports suggesting erythromycin does not reach the fetus in adequate concentrations to prevent congenital syphilis. Infants born to women treated during pregnancy with oral erythromycin for early syphilis should be treated with an appropriate penicillin regimen.

Prolonged QTc interval and ventricular arrhythmias have rarely been reported in patients receiving erythromycin IV.

Rhabdomyolysis with or without renal impairment has been reported in seriously ill patients receiving erythromycin concomitantly with lovastatin.

***Clostridium difficile*-associated disease**

Clostridium difficile-associated disease (CDAD) has been reported with use of many antibacterial agents, including ERYTHROCIN[®] IV. CDAD may range in severity from mild diarrhea to fatal colitis. It is important to consider this diagnosis in patients who present with diarrhea, or symptoms of colitis, pseudomembranous colitis, toxic megacolon, or perforation of colon subsequent to the administration of any antibacterial agent. CDAD has been reported to occur over 2 months after the administration of antibacterial agents.

Treatment with antibacterial agents may alter the normal flora of the colon and may permit overgrowth of *Clostridium difficile*. *C. difficile* produces toxins A and B, which contribute to the development of CDAD. CDAD may cause significant morbidity and mortality. CDAD can be refractory to antimicrobial therapy.

If the diagnosis of CDAD is suspected or confirmed, appropriate therapeutic measures should be initiated. Mild cases of CDAD usually respond to discontinuation of antibacterial agents not directed against *Clostridium difficile*. In moderate to severe cases, consideration should be given to management with fluids and electrolytes, protein supplementation, and treatment with an antibacterial agent clinically effective against *Clostridium difficile*. Surgical evaluation should be instituted as clinically indicated, as surgical intervention may be required in certain severe cases. (see **ADVERSE REACTIONS**).

PRECAUTIONS

Prolonged or repeated use of erythromycin may result in an overgrowth of non-susceptible bacteria or fungi and organisms initially sensitive to erythromycin. If superinfection occurs, erythromycin should be discontinued and appropriate therapy instituted.

Since erythromycin is principally excreted by the liver, caution should be exercised when erythromycin is administered to patients with impaired hepatic function.

There have been reports erythromycin may aggravate the weakness of patients with myasthenia gravis.

There have been reports of infantile hypertrophic pyloric stenosis (IHPS) occurring in infants following erythromycin therapy. In one cohort of 157 newborns who were given erythromycin for pertussis prophylaxis, seven neonates (5%) developed symptoms of non-bilious vomiting or irritability with feeding and were subsequently diagnosed as having IHPS requiring surgical pyloromyotomy. Since erythromycin may be used in the treatment of conditions in infants which are associated with significant mortality or morbidity (such as pertussis or chlamydia), the benefit of erythromycin therapy needs to be weighed against the potential risk of developing IHPS. Parents should be informed to contact their physician if vomiting or irritability with feeding occurs.

Drug Interactions

Serious Drug Interactions

- Concomitant administration of erythromycin with astemizole*, terfenadine*, cisapride*, pimozone*, and ergotamine or dihydroergotamine is contraindicated. (see **CONTRAINDICATIONS**)
- The use of terfenadine* and astemizole* is contraindicated in conjunction with erythromycin due to the occurrence of rare life threatening cardiovascular adverse events including death, cardiac arrest, torsade de pointes and other ventricular arrhythmias. Quinidine, disopyramide and verapamil have been associated with rare instances of cardiac adverse events and should be used with caution in combination with erythromycin
- Erythromycin is an inhibitor of the CYP 3A4 and CYP 1A2 isoenzymes. Concurrent administration of erythromycin and drugs metabolised by either of these two isoenzymes may lead to an increase in the plasma concentrations of the co-administered drug which could result in clinically significant safety concerns.

* No longer marketed in Canada.

Overview

Many categories of drugs are metabolized by CYP3A4 or CYP1A2. Some drugs may inhibit or induce the activities of these two isoenzymes. Administration of such inhibitors or inducers may impact upon the metabolism of co-administered medication(s). In some cases serum concentrations may be increased and in others decreased. Care must therefore be exercised when co-administering such drugs.

Effects of erythromycin on other drugs

Erythromycin is an inhibitor of the cytochrome P450 isoenzymes CYP1A2 and CYP3A4. This inhibition may lead to increased or prolonged serum levels of those drugs metabolized by either of these two isoenzymes when co-administered with erythromycin. For such drugs the monitoring of their serum concentration may be necessary.

Erythromycin should be used with caution in patients receiving treatment with other drugs known to be CYP3A4 and/or CYP1A2 substrates, especially if the substrate has a narrow safety margin and/or the substrate is extensively metabolized by CYP3A4 or CYP1A2.

Dosage adjustments may be considered, and when possible, serum concentrations of these drugs should be monitored closely in patients concurrently receiving erythromycin.

Effects of other drugs on erythromycin

Drugs that induce CYP3A4 (such as rifampicin, phenytoin, carbamazepine, phenobarbital, St John's Wort) may induce the metabolism of erythromycin. This may lead to sub-therapeutic levels of erythromycin and a decreased effect. The induction decreases gradually during two weeks after discontinued treatment with CYP3A4 inducers. Erythromycin should not be used during and two weeks after treatment with CYP3A4 inducers.

Established or Predicted Drug-Drug Interactions

Proper Name	Ref	Effect	Clinical Comments
CYP3A4 inducers			
Rifabutin	T	↓erythromycin levels	Rifabutin is an inducer of the cytochrome P450 isoenzyme CYP3A4. Rifabutin may therefore affect the pharmacokinetic behaviour of drugs metabolised by the enzymes belonging to this subfamily. Studies with the macrolide antibiotic clarithromycin demonstrated that the level of clarithromycin was reduced by approximately half when the medicines were concomitantly administered. An upward adjustment of the dose of clarithromycin and other antibiotics in the macrolide class (e.g.erythromycin) may be required when the medicines are administered in combination with rifabutin.
CYP3A4 substrates			
Alfentanil	CT	↑alfentanil levels	Alfentanil is metabolised in the liver by cytochrome P450 isoenzyme

			<p>CYP3A4.</p> <p>Increased blood concentrations of alfentanil have been reported in patients co-administered erythromycin. This may increase the risk of prolonged or delayed respiratory depression.</p>
Astemizole*	C	↑astemizole levels	<p>Astemizole is metabolised in the liver by the cytochrome P450 isoenzyme CYP3A4. Concomitant administration of astemizole with erythromycin is contraindicated because erythromycin is known to impair the cytochrome P450 enzyme system which also influences astemizole metabolism. Erythromycin significantly alters the metabolism of astemizole when taken concomitantly. Rare cases of serious cardiovascular adverse events including cardiac arrest, torsade de pointes and other ventricular arrhythmias have been observed. (see <u>CONTRAINDICATIONS</u> and <u>ADVERSE REACTIONS</u>).</p>
Bromocriptine	CT	↑bromocriptine levels	<p>Bromocriptine is metabolised in the liver by cytochrome P450 isoenzyme CYP3A4. Increased blood concentrations of bromocriptine have been reported in patients co-administered erythromycin.</p> <p>This appears to be a result of decreased hepatic metabolism of bromocriptine due to the inhibition of CYP3A4 by erythromycin.</p>
Carbamazepine	C	↑carbamazepine levels	<p>Carbamazepine is a substrate of CYP3A4 and macrolides such as erythromycin have been reported to cause substantial elevations of serum concentrations of carbamazepine and symptoms of carbamazepine toxicity.</p>
Cilostazol	CT	↑cilostazol levels	<p>Cilostazol is metabolised in the liver by cytochrome P450 isoenzyme CYP3A4.</p>

* No longer marketed in Canada.

			<p>Increased blood concentrations of cilostazol have been reported in patients co-administered erythromycin.</p> <p>This appears to be a result of decreased hepatic metabolism of cilostazol due to the inhibition of CYP3A4 by erythromycin.</p>
Cisapride*	C	↑cisapride levels	<p>Elevated cisapride levels have been reported in patients receiving erythromycin and cisapride concomitantly. This may result in QT prolongation and cardiac arrhythmias including ventricular tachycardia, ventricular fibrillation and torsade de pointes.</p>
Colchicine	C	↑colchicine levels	<p>There have been post-marketing reports of colchicine toxicity with concomitant use of erythromycin and colchicine.</p>
Cyclosporin	C	↑cyclosporin levels	<p>Cyclosporin is metabolised in the liver by cytochrome P450 isoenzyme CYP3A4.</p> <p>Increased blood concentrations of cyclosporin have been reported in patients co-administered erythromycin.</p> <p>This appears to be a result of decreased hepatic metabolism of cyclosporin due to the inhibition of CYP3A4 by erythromycin.</p>
Disopyramide	C	↑disopyramide levels	<p>Disopyramide is metabolised in the liver by cytochrome P450 isoenzyme CYP3A4.</p> <p>Increased blood concentrations of disopyramide have been reported in patients co-administered erythromycin.</p> <p>This appears to be a result of decreased hepatic metabolism of disopyramide due to the inhibition of CYP3A4 by erythromycin.</p>
Ergotamine/Dihydroergotamine	C	↑ergotamine/dihydroergotamine	<p>There are reports that ischemic reactions may occur when erythromycin is given concurrently with ergotamine-containing drugs.</p> <p>Post-marketing reports indicate that</p>

* No longer marketed in Canada.

			co-administration of erythromycin with ergotamine or dihydroergotamine has been associated with acute ergot toxicity characterized by vasospasm and ischemia of the extremities and other tissues including the central nervous system. (see CONTRAINDICATIONS).
HMG-CoA Reductase Inhibitors	C,CT	↑statin levels	Erythromycin has been reported to increase concentrations of HMG-CoA Reductase inhibitors (e.g., lovastatin and simvastatin). Rare reports of rhabdomyolysis have been reported in patients taking erythromycin concomitantly with HMG-CoA Reductase inhibitors.
Phenytoin	T	↑phenytoin levels	The metabolism of phenytoin is complex and is believed to be mediated by several cytochrome enzymes, particularly cytochrome P450 CYP2C9 and to a lesser extent the CYP 2C19 and CYP3A4 isoenzymes. Increased blood concentrations of phenytoin have been reported in patients co-administered erythromycin. This appears to be a result of decreased hepatic metabolism of phenytoin due to the inhibition of CYP3A4 by erythromycin.
Pimozide*	T	↑pimozide levels	Pimozide is a substrate for cytochrome P450 isoenzyme CYP3A4. Elevated pimozide levels have been observed with concomitant administration with clarithromycin, another macrolide antibiotic. This may result in QT prolongation and cardiac arrhythmias including ventricular tachycardia, ventricular fibrillation and torsade de pointes..
Quinidine	C	↑quinidine levels	Quinidine is metabolised in the liver by cytochrome P450 isoenzyme CYP3A4. Increased blood concentrations of quinidine have

* No longer marketed in Canada.

			<p>been reported in patients co-administered erythromycin.</p> <p>This appears to be a result of decreased hepatic metabolism of quinidine due to the inhibition of CYP3A4 by erythromycin.</p>
Sildenafil	C	↑sildenafil levels	<p>Sildenafil is metabolised in the liver by cytochrome P450 isoenzyme CYP3A4.</p> <p>Increased blood concentrations of sildenafil have been reported in patients co-administered erythromycin.</p> <p>This appears to be a result of decreased hepatic metabolism of sildenafil due to the inhibition of CYP3A4 by erythromycin.</p>
Tacrolimus	C	↑tacrolimus levels	<p>Tacrolimus is metabolised in the liver by cytochrome P450 isoenzyme CYP3A4. Increased blood concentrations of tacrolimus have been reported in patients co-administered erythromycin.</p>
Terfenadine*	C	↑terfenadine levels	<p>Terfenadine undergoes metabolism in the liver by the specific cytochrome P450 isoenzyme, CYP3A4. This metabolic pathway may be impaired in patients who are taking erythromycin, an inhibitor of this isoenzyme. Interference with this enzyme can lead to elevated terfenadine plasma levels which may be associated with QT prolongation. Rare cases of serious cardiovascular adverse events including death, cardiac arrest, torsades de pointes and other ventricular arrhythmias (such as ventricular tachycardia, and ventricular fibrillation) have been observed. (see CONTRAINDICATIONS and ADVERSE REACTIONS).</p>
Triazolobenzodiazepines (such as triazolam and alprazolam) and related benzodiazepines	C	↑triazolobenzodiazepines levels	<p>Erythromycin has been reported to decrease the clearance of triazolam and midazolam, and related benzodiazepines, and thus may</p>

* No longer marketed in Canada.

			increase the pharmacologic effects of these benzodiazepines.
Verapamil	C	↑verapamil levels	Verapamil is metabolised in the liver by cytochrome P450 isoenzyme CYP3A4. Increased blood concentrations of verapamil have been reported in patients co-administered erythromycin. This appears to be a result of decreased hepatic metabolism of verapamil due to the inhibition of CYP3A4 by erythromycin. Hypotension, bradyarrhythmias, and lactic acidosis have been observed in patients receiving concurrent verapamil, belonging to the calcium channel blockers drug class.
Vinblastine	T	↑vinblastine levels	Vinblastine is metabolised in the liver by cytochrome P450 isoenzyme CYP3A4. Concomitant administration of vinblastine and erythromycin has been demonstrated to increase the toxicity of vinblastine. This appears to be a result of decreased hepatic metabolism of vinblastine due to the inhibition of CYP3A4 by erythromycin.
Zopiclone	CT	↑zopiclone levels	Erythromycin has been reported to decrease the clearance of zopiclone and thus may increase the pharmacodynamic effects of this drug.
CYP1A2 substrates			
Theophylline	CT	↑theophylline levels	Recent data from studies of erythromycin in patients reveal that its use in patients who are receiving high doses of theophylline may be associated with an increase in serum theophylline levels and potential theophylline toxicity. In case of theophylline toxicity and/or elevated serum theophylline levels, the dose of theophylline should be reduced while the patient is receiving concomitant erythromycin therapy.

			There have been published reports suggesting that when oral erythromycin is given concurrently with theophylline there is a significant decrease in erythromycin serum concentrations. This decrease could result in subtherapeutic concentrations of erythromycin.
Other drug interactions			
Digoxin	C	↑digoxin levels	There have been reports that there is a rise in plasma digoxin levels during concomitant administration of erythromycin
Lincomycin, Clindamycin, Chloramphenicol	T	↓effect of coadministered drugs	Erythromycin should be used with caution if administered concomitantly with these drugs. <i>In vitro</i> experiments have demonstrated that binding sites for erythromycin, lincomycin, clindamycin and chloramphenicol overlap and competitive inhibition may occur.
Methylprednisolone	CT	↑methylprednisolone levels	Concurrent administration of erythromycin and methylprednisolone has been shown to decrease the clearance of methylprednisolone and increase the half life. The mechanistic basis of this interaction is uncertain, but may involve enzymes in the Cytochrome P450 system.
Oral Anticoagulants	C	↑prothrombin time	Published reports indicate that caution should be observed when erythromycin and oral anticoagulants are used concurrently since prothrombin time may be prolonged.
Valproate	C	↑valproate levels	Increased blood concentrations of valproate have been reported in patients co-administered erythromycin. The mechanistic basis of this interaction is uncertain, but may involve enzymes in the Cytochrome P450 system.

C=Case Study; CT= Clinical Trial; T= Theoretical

Laboratory Test

Erythromycin interferes with the fluorometric determination of urinary catecholamines.

Carcinogenesis, Mutagenesis, Impairment of Fertility

Long-term (two-year) oral studies conducted in rats up to about 400 mg/kg/day and in mice up to about 500 mg/kg/day with erythromycin stearate did not provide evidence of tumorigenicity. Mutagenicity studies conducted did not show any genotoxic potential, and there was no apparent effect on male or female fertility in rats treated with erythromycin base by oral gavage at 700 mg/kg/day.

Pregnancy

There are no adequate and well-controlled studies in pregnant women. However, observational studies in humans have reported cardiovascular malformations after exposure to medicinal products containing erythromycin during early pregnancy.

Erythromycin should not be used by women during pregnancy unless clearly needed.

Erythromycin has been reported to cross the placental barrier in humans, but fetal plasma levels are generally low.

No evidence of teratogenicity or embryotoxicity was observed in the following studies in animals:

Reproductive toxicity in rats with 350 mg/kg/day (7 times the human dose) and 700 mg/kg/day (14 times the human dose) of erythromycin base prior to and during mating, during gestation, and through weaning.

Reproductive toxicity in Swiss Webster mice with 700 mg/kg/day (14 times the human dose) of erythromycin base during the period of embryo-fetal organogenesis (gestational day 6-15).

Labour and Delivery

The effect of erythromycin on labour and delivery is unknown.

Nursing mothers

The safety of erythromycin for use during breast feeding has not been established. Erythromycin is excreted in breast milk.

Neonates

The safety of erythromycin for use in neonates has not been established.

Pediatric Use

See **DOSAGE AND ADMINISTRATION**.

ADVERSE REACTIONS

- Gastrointestinal** Abdominal cramping, discomfort. Nausea, vomiting, diarrhea and anorexia are also observed but less frequently. Pseudomembranous colitis has been occasionally reported to occur in association with erythromycin therapy (see **WARNINGS**).
- Hepatotoxicity** Symptoms of hepatitis, hepatic dysfunction and/or abnormal liver function test results may occur (see **WARNINGS**).
- Pancreatitis** There have been rare reports of pancreatitis. There has also been a report of a case of erythromycin-induced pancreatitis following erythromycin overdose.
- Allergic reactions** Urticaria, mild skin eruptions and anaphylaxis. Skin reactions ranging from mild eruptions to erythema multiform, Stevens-Johnson syndrome and toxic epidermal necrolysis have rarely been reported.
- Cardiovascular** Occasional case reports of cardiac arrhythmias such as ventricular tachycardia have been documented in patients receiving erythromycin therapy. There have been isolated reports of other cardiovascular symptoms such as chest pain, dizziness, and palpitations; however, a cause and effect relationship has not been established. As with other macrolides, QT prolongation, ventricular tachycardia, and Torsades de Pointes have been reported with erythromycin (see **CONTRAINDICATIONS**).
- Neurological** Central nervous system side effects including seizures, hallucinations, confusion, vertigo and tinnitus have been reported occasionally in patients; however, a cause and effect relationship has not been established. There have also been rare reports of convulsions.
- Miscellaneous** During prolonged or repeated therapy, there is a possibility of overgrowth of nonsusceptible bacteria or fungi and organisms initially sensitive to erythromycin (e.g. *Staphylococcus aureus*, *Hemophilus influenzae*). If such infections occur, erythromycin should be discontinued and appropriate therapy instituted.
- Occasionally there have been reports of reversible hearing loss occurring chiefly in patients with renal insufficiency and patients receiving high doses of erythromycin.
- There have been reports of interstitial nephritis coincident with erythromycin use.
- Venous irritation has been encountered occasionally during intravenous administration of erythromycin. If the infusion is given slowly, in dilute solution, preferably by continuous intravenous infusion or intermittent infusion over no less than 20 to 60 minutes, pain and vessel trauma are minimized.

SYMPTOMS AND TREATMENT OF OVERDOSAGE

For management of a suspected drug overdose contact your regional Poison Control Centre.

Symptoms

Recently there has been a report of a case of erythromycin-induced pancreatitis following erythromycin overdose.

Treatment

There is no specific treatment for overdosage. Erythromycin should be discontinued and gastric lavage considered, if appropriate; otherwise, the treatment should be symptomatic.

Erythromycin is not removed by peritoneal dialysis or hemodialysis.

DOSAGE AND ADMINISTRATION

ERYTHROCIN[®] I.V. - (erythromycin lactobionate for injection) should be used in the treatment of patients when oral administration is not possible or when it is desirable to obtain higher serum levels of erythromycin than achievable with orally administered preparations. Intravenous erythromycin should be replaced by an oral form of erythromycin as soon as possible.

Dosage

The doses are expressed in terms of the base.

Adults: The recommended intravenous dose is 15 to 20 mg/kg/day. In severe infections, doses of up to 4 g of erythromycin may be given daily in divided doses.

For treatment of Legionnaires' Disease: Optimal dosages have not been established; however, the larger doses (eg. 4 g daily) should be considered for known or suspected Legionella infections. Doses utilized in reported clinical data were 1 to 4 grams daily in divided doses.

Children: Age, weight, and severity of the infection are important factors in determining the proper dosage. The recommended dose is 15 to 20 mg/kg/day.

Administration

ERYTHROCIN[®] I.V. must be administered by continuous or intermittent intravenous infusion only. Due to the irritative properties of ERYTHROCIN[®] I.V., intravenous push is not an acceptable method of administration.

Continuous infusion of erythromycin lactobionate is preferable due to the slow infusion rate and lower concentration of erythromycin; however, intermittent infusion at intervals not greater than every 6 hours may also be used.

Continuous Intravenous Infusion: For slow continuous infusion, erythromycin lactobionate solutions should be diluted to give a final concentration of 1 g per litre (1 mg/mL).

Intermittent Intravenous Infusion: One-fourth of the total daily dose of erythromycin lactobionate should be administered by intravenous infusion in 20 to 60 minutes at intervals not greater than every 6 hours. The erythromycin lactobionate solutions should be diluted to give a final concentration of 1 to 5 mg/mL. No less than 100 mL of i.v. diluent should be used. Infusion should be sufficiently slow to minimize pain along the vein.

Reconstitution

Fliptop Vial Reconstitution

Reconstitute with Sterile Water for Injection USP only as indicated in Table 3. Use of other diluents may cause precipitation during reconstitution. Do not use diluents containing preservatives or inorganic salts.

Table 3 Reconstitution of ERYTHROCIN® I.V. vials		
Vial Size	Volume to be Added to Vial	Erythromycin Concentration
500 mg	10 mL	50 mg/mL
1 g	20 mL	50 mg/mL

Stability and Storage Recommendation for stock solution:

The reconstituted stock solution is stable under refrigeration for 96 hours, or for 24 hours between 15° to 25°C. The stock solution must be diluted before use (see Vial Dilution).

Vial Dilution:

For Continuous Intravenous Infusion dilute the reconstituted stock solution in 0.9% Sodium Chloride Injection USP, Lactated Ringer's Injection USP, or NORMOSOL-R* to give a concentration of approximately 1 mg/mL.

For Intermittent Intravenous Infusion dilute the reconstituted stock solution to a final concentration of 1 to 5 mg/mL in 0.9% Sodium Chloride Injection USP, Lactated Ringer's Injection USP, or NORMOSOL-R*.

The following solutions may also be used for continuous or intermittent intravenous infusion provided they are first buffered by the addition of 1 mL of 4% sodium bicarbonate per 100 mL of solution.

- 5% Dextrose Injection USP
- 5% Dextrose and Lactated Ringer's Injection
- 5% Dextrose and 0.9% Sodium Chloride Injection USP

Sodium bicarbonate must be added to these solutions so that their pH is in the optimum range for erythromycin lactobionate stability. Acidic solutions of erythromycin are unstable and lose their potency rapidly. A pH of at least 5.5 is desirable for the final diluted solution of erythromycin lactobionate.

No drug or chemical agent should be added to an ERYTHROCIN[®] - I.V. fluid admixture unless its effect on the chemical and physical stability of the solution has first been determined.

The pH of intravenous solutions in plastic containers tends to be lower than that of the same solutions in glass containers. The concurrent use of additives that will result in an erythromycin lactobionate admixture with a pH below 5.5 should be avoided.

Stability of Solution Prepared Using Fliptop Vials:

The final diluted solution of erythromycin lactobionate is not suitable for storage and should be completely administered **within 8 hours** in order to assure proper potency.

DESCRIPTION

Erythromycin is produced by a strain of *Streptomyces erythraeus* and belongs to the macrolide group of antibiotics. It is chemically known as erythromycin mono-(4-O-β-D-galactopyranosyl-D-glucinate). It is basic and readily forms salts with acids. It occurs as white or slightly yellow, odourless or almost odourless, slightly hygroscopic crystals or powder with a bitter taste. It is freely soluble in methanol, ethanol, acetone and chloroform. It is soluble in water at 2 mg/mL. The melting point is 135 to 140°C.

Erythromycin lactobionate is a white amorphous salt of erythromycin freely soluble in methanol and ethanol. It is soluble in water at 200 mg/mL.

ERYTHROCIN[®] - I.V. (erythromycin lactobionate for injection) **Fliptop vials** contain a soluble salt of erythromycin **without preservative** suitable for intravenous administration.

AVAILABILITY

Each Fliptop vial contains 500 mg or 1 g erythromycin base in the form of erythromycin lactobionate for injection as sterile lyophilized powder. They are available in packages of 10 vials.

Storage Recommendations and Stability:

ERYTHROCIN[®] I.V. (erythromycin lactobionate for injection) powder should be stored between 15 to 25°C and protected from heat.