

## **PRODUCT MONOGRAPH**

**Pr RIFADIN®**

**Rifampin capsules USP  
150 mg & 300 mg**

**Antibiotic**

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## PRODUCT MONOGRAPH

<sup>Pr</sup>RIFADIN<sup>®</sup>  
(rifampin)

Capsules  
150 mg & 300 mg

Antibiotic

### CLINICAL PHARMACOLOGY

RIFADIN (rifampin) inhibits DNA-dependent RNA polymerase activity in susceptible cells. Specifically, it interacts with bacterial RNA polymerase, but does not inhibit the mammalian enzyme. Cross-resistance to rifampin has only been shown with other rifamycins. This is the probable mechanism of action by which rifampin exerts its therapeutic effects.

Absorption is more rapid when rifampin is administered one hour before meals. Peak blood levels in normal adults vary widely from individual to individual. Peak levels occur between 2 and 4 hours following the oral administration of a 600 mg dose with average peak values of 7-10 µg/mL.

Rifampin is distributed throughout the body and is detectable in many organs and body fluids, including the cerebrospinal fluid. The highest concentrations are present in the liver and bile.

In normal subjects, the biological half-life of rifampin in serum averages about 3 hours after a 600 mg oral dose, with increases up to 5.1 hours reported after 900 mg dose. Rifampin is eliminated from the blood equally in the urine and feces as unchanged drug and metabolites.

The principal metabolite in man is the biologically active desacetylrifampin. Desacetylation of rifampin in the body does not substantially modify its anti-mycobacterial activity. In Kirschner's medium, the MIC against *M. tuberculosis* varied from 0.1 to 2 µg/mL.

### INDICATIONS AND CLINICAL USE

RIFADIN (rifampin) is indicated as a treatment of tuberculosis.

To achieve a complete kill of the bacillary population and to avoid selection of drug-resistant mutants, RIFADIN must be used concomitantly with at least one other active anti-tuberculous drug. The selection of the specific drug for partner is determined by the *in vitro* sensitivity of the causative organisms, comparative safety and effectiveness, the patient's previous clinical history and the absorption/distribution pattern of the drug.



It is also indicated for the prophylaxis of bacterial meningitis or carriage of *N. meningitidis* or *H. influenza b* in persons exposed to a primary case.

To reduce the development of drug-resistant bacteria and maintain the effectiveness of RIFADIN and other antibacterial drugs, RIFADIN should be used only to treat 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

- Jaundice associated with reduced bilirubin excretion.
- RIFADIN is contraindicated in patients who are hypersensitive to this drug, who have a history of previous sensitivity to any of the rifamycins or to any ingredient in the formulation, including any non-medicinal ingredient, or component of the container. For a complete listing, see AVAILABILITY OF DOSAGE FORMS.
- Premature and newborn infants in whom the liver is not yet capable of functioning with full efficiency.
- RIFADIN (rifampin) passes into the breast milk and therefore should not be used during lactation.
- RIFADIN use is contraindicated when given concurrently with the combination of saquinavir/ritonavir (see PRECAUTIONS).

## WARNINGS

RIFADIN (rifampin) has been shown to produce liver dysfunction. There have been fatalities associated with jaundice in patients with liver disease or receiving RIFADIN concomitantly with other hepatotoxic agents. Since an increased risk may exist for individuals with liver disease, benefits must be weighed carefully against the risk of further liver damage. Patients with impaired liver function should only be given rifampin in case of necessity and then with caution and under strict medical supervision. In these patients, careful monitoring of liver function, especially serum alanine aminotransferase (ALT) and serum aspartate aminotransferase (AST) should be carried out prior to therapy and then every two to four weeks during therapy. If signs of hepatocellular damage occur, RIFADIN should be withdrawn.

In some cases, hyperbilirubinemia resulting from competition between rifampin and bilirubin for excretory pathways of the liver at the cell level can occur in the early days of treatment. An isolated report showing a moderate rise in bilirubin and/or transaminase level is not in itself an indication for interrupting treatment; rather, the decision should be made after repeating the tests, noting trends in the levels, and considering them in conjunction with the patient's clinical condition.



Rifampin has enzyme-inducing properties, including induction of delta aminolevulinic acid synthetase. Isolated reports have associated porphyria exacerbation with rifampin administration.

### **Drug Reaction with Eosinophilia and Systemic Symptoms (DRESS)**

Severe, systemic hypersensitivity reactions, including fatal cases, such as Drug Reaction with Eosinophilia and Systemic Symptoms (DRESS) syndrome have been observed during treatment with anti-tuberculosis therapy (see ADVERSE REACTIONS).

It is important to note that early manifestations of hypersensitivity, such as fever, lymphadenopathy or biological abnormalities (including eosinophilia, liver abnormalities) may be present even though rash is not evident. If such signs or symptoms are present, the patient should be advised to consult immediately their physician.

RIFADIN should be discontinued if an alternative etiology for the signs and symptoms cannot be established.

### **Clostridium difficile-associated disease**

*Clostridium difficile*-associated disease (CDAD) has been reported with use of many antibacterial agents, including rifampin. 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).

### **Susceptibility/Resistance**

#### **Development of Drug Resistant Bacteria**

Prescribing RIFADIN in the absence of a proven or strongly suspected bacterial infection is unlikely to provide benefit to the patient and risks the development of drug-resistant bacteria.



## PRECAUTIONS

### General

A complete blood count (CBC), serum creatinine and liver function tests should be obtained prior to instituting therapy and periodically throughout the course of therapy. Because of a possible transient rise in transaminase and bilirubin values, blood for baseline clinical chemistries should be obtained before RIFADIN dosing.

RIFADIN (rifampin) increases the requirements for anticoagulant drugs of the coumarin type. This effect is not observed until the fifth day following initiation of treatment. The decrease in prothrombin time usually lasts between 5 and 7 days, and is the result of RIFADIN's ability to cause induction of drug metabolizing enzyme systems of the liver. As a result, the rate of metabolism of those drugs which are substrates for these enzymes can be altered, resulting in reduced pharmacological effects of the drugs involved. In patients receiving anticoagulants, it is recommended that daily prothrombin times be performed until the dose of the anticoagulant required has been established. This is particularly important when rifampin administration is either initiated or withdrawn.

The intermittent administration of high doses of RIFADIN >120 mg/dose has been reported to be associated with a hypersensitivity reaction, characterized by fever and myalgia. The incidence of this reaction is greater when RIFADIN is given on a once-a-week basis than on a twice or thrice weekly basis. It is recommended that when resuming treatment with RIFADIN after short or prolonged interruptions, it be given in small, gradually increasing doses. During the transitional period, the renal and hemopoietic systems should be closely monitored. The drug should be stopped immediately if renal failure, thrombocytopenia purpura or hemolytic anaemia develop and should not be re-instituted.

Safe conditions for the use of ethambutol alone or in combination with RIFADIN have not been established for children under the age of thirteen years. Although renal insufficiency does not alter blood levels of RIFADIN, marked increases in ethambutol levels are observed under similar conditions; this, therefore, should be taken into consideration in such patients receiving RIFADIN /ethambutol combination therapy. Caution is recommended when instituting therapeutic regimens in which isoniazid is to be used concurrently with RIFADIN, in patients with impaired liver function, the elderly and malnourished.

From experimental studies, it would appear that bromosulphalein (BSP) and RIFADIN compete with one another at the liver cell-bile canaliculus boundary. Clinically, this phenomenon can be reflected by spurious BSP levels. It is recommended that the BSP test be carried out at least five hours after the last dose of RIFADIN.



RIFADIN and its metabolites may produce a discoloration (yellow, orange, red, brown) of the teeth, urine, feces, saliva, sputum, sweat and tears. Individuals to be treated should be made aware of these possibilities in order to prevent undue anxiety.

Patients should be advised that soft contact lenses may be permanently stained. Teeth discoloration may also be permanent (see ADVERSE REACTIONS).

It has been reported that oral contraceptives have failed to prevent conception in some patients receiving RIFADIN in association with other anti-tuberculosis drugs. It is therefore necessary that alternative or additional contraceptive measures be recommended.

### **Use in pregnancy**

#### Teratogenic Effects

Although rifampin has been reported to cross the placental barrier and appear in the cord blood, the effect of combinations of RIFADIN with other anti-tuberculous drugs on the human fetus is not known. No obvious effect on the fetus was detected after the administration of RIFADIN to 15 pregnant patients. An increase in congenital malformations, primarily spinabifida and cleft palate, has been reported in the offspring of mice and rats given oral doses of RIFADIN 100 mg/kg/day during pregnancy.

RIFADIN should not be used in pregnant women or women with childbearing potential. If RIFADIN therapy is judged to be essential, such treatment should be implemented only after carefully weighing the potential benefits of therapy against the risks which may be involved. In women with childbearing potential, treatment with RIFADIN should be undertaken only when the possibility of pregnancy during therapy is judged to be remote.

#### Non-teratogenic effects

It is not known whether RIFADIN can affect reproduction capacity.

When administered during the last few weeks of pregnancy, rifampin can cause postnatal hemorrhages in the mother and infant. In this case, treatment with vitamin K may be indicated for postnatal hemorrhage.

### **Nursing mothers**

Rifampin is excreted in breast milk. Therefore, RIFADIN should not be used during lactation (see CONTRAINDICATIONS).



## **Carcinogenesis, mutagenesis, impairment of fertility**

There are no known human data on long-term potential for carcinogenicity, mutagenicity, or impairment of fertility. A few cases of accelerated growth of lung carcinoma have been reported in man, but a causal relationship with the drug has not been established. An increase in the incidence of hepatomas in female mice (of a strain known to be particularly susceptible to the spontaneous development of hepatomas) was observed when rifampin was administered in doses two to ten times the average daily human dose for 60 weeks followed by an observation period of 46 weeks. No evidence of carcinogenicity was found in male mice of the same strain, mice of a different strain, or rats under similar experimental conditions.

Rifampin has been reported to possess immunosuppressive potential in rabbits, mice, rats, guinea pigs, human lymphocytes *in vitro*, and humans. Antitumor activity *in vitro* has been shown with rifampin.

There was no evidence of mutagenicity in bacteria, *Drosophila melanogaster*, or mice. An increase in chromatid breaks was noted when whole blood cell cultures were treated with rifampin.

## **Drug interaction**

Rifampin is a potent inducer of certain cytochrome P-450 enzymes. Coadministration of rifampin with other drugs that are metabolized through these cytochrome P-450 enzymes may accelerate the metabolism and reduce the activity of these other drugs. Therefore, caution should be used when prescribing rifampin with drugs metabolized by cytochrome P-450. To maintain optimum therapeutic blood levels, dosages of drugs metabolized by these enzymes may require adjustment when starting or stopping concomitantly administered rifampin.

Examples of drugs metabolized by cytochrome P-450 enzyme are: anticonvulsants (eg, phenytoin), antiarrhythmics (eg, disopyramide, mexiletine, quinidine, tocainide, propafenone), antiestrogens (eg tamoxifen, toremifene), antipsychotics (eg haloperidol), oral anticoagulants (eg warfarin) antifungals (eg, fluconazole, itraconazole, ketoconazole), antiretroviral drugs (e.g. zidovudine, saquinavir, indinavir, efavirenz), barbiturates, beta-blockers, benzodiazepines (eg diazepam), benzodiazepine-related drugs (eg zopiclone, zolpidem), calcium channel blockers (eg, diltiazem, nifedipine, verapamil), chloramphenicol, clarithromycin, corticosteroids, cardiac glycosides preparations, clofibrate, oral contraceptives, dapsone, doxycycline, estrogens, gestrinone, oral hypoglycemic agents (sulfonylureas), immunosuppressive agents (e.g. cyclosporine, tacrolimus), irinotecan, levothyroxine, losartan, narcotic analgesics, methadone, praziquantel, progestins, quinine, riluzole, selective 5-HT<sub>3</sub> receptor antagonists (eg ondansetron), statins metabolized by CYP 3A4, telithromycin, theophylline, thiazolidinediones (e.g. rosiglitazone), tricyclic antidepressants (eg amitriptyline, nortriptyline).

Upon completion of the treatment with RIFADIN, a renewed readjustment of the dosage should be made.



## Other interaction

Atovaquone: when the two drugs are taken concomitantly, decreased concentrations of atovaquone and increased concentrations of rifampicin were observed.

Concurrent use of ketoconazole and rifampin has resulted in decreased serum concentration of both drugs. Concurrent use of rifampin and enalapril has resulted in decreased concentrations of enalaprilat, the active metabolite of enalapril. Dosage adjustments should be made if indicated by the patient's clinical condition.

Concomitant antacid administration may reduce the absorption of rifampin. Daily doses of rifampin should be given at least 1 hour before the ingestion of antacids.

Probenecid and cotrimoxazole have been reported to increase the blood level of rifampin.

When RIFADIN is given concomitantly with the combination saquinavir/ritonavir, the potential for hepatotoxicity is increased. Therefore, concomitant use of RIFADIN with saquinavir/ritonavir is contraindicated (see CONTRAINDICATIONS).

When rifampin is given concomitantly with either halothane or isoniazid the potential for hepatotoxicity is increased. The concomitant use of RIFADIN, which contains rifampin, and halothane should be avoided. Patients receiving both rifampin and isoniazid should be monitored closely for hepatotoxicity (see WARNINGS).

Plasma concentration of sulfapyridine may be reduced following the concomitant administration of sulfasalazine and rifampin. This finding may be the result of alteration in the colonic bacteria responsible for the reduction of sulfasalazine to sulfapyridine and mesalamine.

## Drug/laboratory tests interaction

Therapeutic levels of rifampin have been shown to inhibit standard microbiological assays for serum folate and vitamin B<sub>12</sub>. Therefore, alternate assay methods should be considered.

Transient abnormalities in liver function tests (eg, elevation in serum bilirubin, abnormal bromosulphalein [BSP] excretion, alkaline phosphatase and serum transaminases), and reduced biliary excretion of contrast media used for visualization of the gallbladder have also been observed. Therefore, these tests should be performed before the morning dose of RIFADIN.

Rifampin has enzyme induction properties that can enhance the metabolism of endogenous substrates including adrenal hormones, thyroid hormones, and vitamin D.

Cross-reactivity and false-positive urine screening tests for opiates have been reported in patients receiving rifampicin when using the KIMS (Kinetic Interaction of Microparticles in Solution) method (eg, Abuscreen OnLine opiates assay; Roche Diagnostic Systems). Confirmatory tests, such as gas chromatography/ mass spectrometry, will distinguish rifampicin from opiates.



## ADVERSE REACTIONS

RIFADIN (rifampin) is usually well tolerated at recommended dosage levels.

**Blood and lymphatic system disorders:** Thrombocytopenia, purpura, leukopenia, hemolytic anaemia (may be related to hypersensitivity reactions) and decreased hemoglobin have been observed. Thrombocytopenia with or without purpura has occurred when RIFADIN and ethambutol were administered concomitantly according to an intermittent dose schedule twice weekly and in high doses.

Thrombocytopenia has occurred primarily with high dose intermittent therapy, but has also been noted after resumption of interrupted treatment. It rarely occurs during well-supervised daily therapy. This effect is reversible if the drug is discontinued as soon as purpura occurs.

- Disseminated intravascular coagulation has also been rarely reported.
- Agranulocytosis has been reported very rarely.
- Occasionally, eosinophilia associated to hypersensitivity reactions has been encountered.

**Endocrine disorders:** Rare reports of adrenal insufficiency in patients with compromised adrenal function have been observed.

**Eye disorders:** Visual disturbances have been observed.

- Occasionally, conjunctivitis associated with hypersensitivity reactions has been encountered.

**Gastrointestinal disorders:** heartburn, epigastric distress, nausea, vomiting, gas, cramps, abdominal discomfort and diarrhea, have been noted in some patients.

Occasionally, sore mouth, sore tongue associated with hypersensitivity reactions have been encountered.

**Hepatobiliary disorders:** Rarely, hepatitis or a shock-like syndrome with hepatic involvement and abnormal liver function tests has been reported. Transient abnormalities in liver function tests (elevations of serum bilirubin, BSP, alkaline phosphatase and serum transaminases) have been observed, particularly during the first few weeks of treatment (see WARNINGS section).

A few cases of jaundice with evidence of hepatocellular damage have been reported in patients receiving RIFADIN. In some of them it was possible, under careful laboratory control, to resume treatment after an interval without recurrence of abnormalities.

**Immune system disorders:** Immunological reactions (including anaphylaxis/anaphylactic reaction) with shortness of breath, wheezing, decrease in blood pressure and shock have occurred with intermittent dosage regimens.

Occasionally, hypersensitivity reactions such as: pruritus, urticaria, skin rashes, pemphigoid reaction, eosinophilia, sore mouth, sore tongue, conjunctivitis, acute hemolytic anemia and acute



renal failure (usually due to acute tubular necrosis or to acute interstitial nephritis) have been observed.

**Investigations:** decrease in blood pressure (associated with hypersensitivity reactions)

**Infection and infestations:** Although *Clostridium difficile* has been shown *in vitro* to be sensitive to rifampin, pseudomembranous colitis has been reported with the use of rifampin (and other broad spectrum antibiotics). Therefore, it is important to consider this diagnosis in patients who develop diarrhea in association with antibiotic use (see WARNINGS).

Other reactions which have occurred with intermittent dosage regimens include “flu” syndrome/influenza (such as episodes of fever/pyrexia, chills, headache, dizziness, bone pain and malaise).

The “flu” syndrome/influenza may also appear if rifampin is taken irregularly by the patient or if daily administration is resumed after a drug free interval. It rarely occurs during well-supervised daily therapy.

**Metabolism and nutritional disorders:** anorexia/decreased appetite

**General disorders and administration site conditions:** Edema of the face and extremities has been reported.

**Musculoskeletal and connective tissue disorders:** Rare reports of myopathy have also been observed. Bone pain (may be related to hypersensitivity reactions).

**Nervous system disorders:** headache, drowsiness, fatigue, ataxia, dizziness, inability to concentrate, mental confusion, psychoses, behavioral changes, muscular weakness, fever, pains in extremities and generalized numbness have also been noted.

Cerebral hemorrhage and fatalities have been reported when rifampin administration has been continued or resumed after the appearance of purpura.

**Renal and urinary disorders:** Elevations in BUN and serum uric acid have been reported. Rarely, hemoglobinuria, hematuria, interstitial nephritis, renal insufficiency, tubular necrosis interstitial nephritis and acute kidney injury have been noted.

These are generally considered to be hypersensitivity reactions. They usually occur during intermittent therapy or when treatment is resumed following intentional or accidental interruption of a daily dosage regimen, and are reversible when rifampin is discontinued and appropriate therapy instituted.

#### **Reproductive system and breast disorders**

The following menstrual disturbances/disorders: breakthrough bleeding, spotting, amenorrhea, monthly prolongation of both menstrual interval and menses have been reported.



**Respiratory, thoracic and mediastinal disorders:** shortness of breath/dyspnea and wheezing are generally associated to hypersensitivity reactions.

**Skin and subcutaneous tissue disorders:** cutaneous reactions are mild and self-limiting and do not appear to be hypersensitivity reactions. Typically, they consist of flushing and itching with or without a rash (pruritus/rash pruritic). More serious cutaneous reactions/dermatitis allergic, which may be due to hypersensitivity, occur but are uncommon. Erythema multiforme including Stevens-Johnson syndrome, toxic epidermal necrolysis and vasculitis have been reported on rare occasions. Occasionally, pruritus, urticaria, skin rashes, and pemphigoid reaction are associated with hypersensitivity reactions.

Clinical trials have furnished no evidence to suggest that RIFADIN has any harmful effects on the cochleovestibular system.

### **Post-Market Adverse Drug Reactions**

#### **Congenital, familial and genetic disorders:**

- Porphyria

#### **Eye disorders:**

- Tear discoloration

#### **Gastrointestinal disorders:**

- Gastrointestinal disorder
- Tooth discoloration (which may be permanent)

#### **Investigations:**

- Blood creatinine increased, hepatic enzyme increased
- Blood bilirubin increased, aspartate aminotransferase increased, alanine aminotransferase increased

#### **Musculoskeletal and connective tissue disorders:**

- Muscle weakness

#### **Psychiatric disorders:**

- Psychotic disorder

#### **Pregnancy, puerperium and perinatal conditions:**

- Post-partum hemorrhage, fetal-maternal hemorrhage

#### **Renal and urinary disorders:**

- Chromaturia



**Respiratory, thoracic and mediastinal disorders:**

- sputum discolored

**Skin and subcutaneous tissue disorders:**

- Skin reaction, sweat discoloration
- Drug Reaction with Eosinophilia and Systemic Symptoms (DRESS) syndrome (see WARNINGS)

**SYMPTOMS AND TREATMENT OF OVERDOSAGE**

For management of a suspected drug overdose, contact your regional Poison Control Centre.
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The minimum acute lethal or toxic dose is not well established. However, non fatal acute overdoses in adults have been reported with doses ranging from 9 to 12 g rifampicin. Fatal acute overdoses in adults have been reported with doses ranging from 14-60 g. Alcohol or a history of alcohol abuse was involved in some of the fatal and non-fatal reports.

Nonfatal overdoses in pediatric patients ages 1 to 4 years old of 100 mg/kg for one to two doses has been reported.

**Signs and Symptoms**

Nausea, vomiting, abdominal pain, pruritus, headache and increasing lethargy will probably occur within a short time after acute ingestion; unconsciousness may occur when there is severe hepatic disease. Brownish-red or orange discoloration of the skin, urine, sweat, saliva, tears, teeth and feces will occur, and its intensity is proportional to the amount ingested. Facial or periorbital edema has also been reported in pediatric patients. Hypotension, sinus tachycardia, ventricular arrhythmias, seizures and cardiac arrest were reported in some fatal cases.

Liver enlargement, possibly with tenderness, can develop within few hours after severe overdose; bilirubin levels may increase and jaundice may develop rapidly. Hepatic involvement may be more marked in patients with prior impairment of hepatic function. Other physical findings remain essentially normal. A direct effect upon the hematopoietic system, electrolyte levels, or acid-base balance is unlikely.

**Management**

In the event of an acute oral overdose activated charcoal may be considered. General supportive measures are recommended and individual symptoms should be treated as they arise.

Antiemetic medication may be required to control severe nausea and vomiting. Active diuresis (with measured intake and output) will help promote excretion of the drug. Hemodialysis may be of value in some patients. No specific antidote is known.



## DOSAGE AND ADMINISTRATION

### Treatment of tuberculosis

#### Adults

600 mg in a single daily dose. Should intolerance occur, the daily dosage may be reduced to 450 mg. In patients with impaired liver function, a daily dose of 8 mg/kg should not be exceeded. A daily dosage of 10 mg/kg is recommended for frail and elderly persons.

#### Children

10 to 20 mg/kg not to exceed 600 mg/day. Data is not available for the determination of dosage for children under 5 years of age.

In treatment of pulmonary tuberculosis, RIFADIN (rifampin) must be used in conjunction with at least one other anti-tuberculous agent. In general, therapy should be continued until bacterial conversion has been established and maximum clinical improvement has occurred.

To ensure optimum absorption, RIFADIN should be taken on an empty stomach (1 hour before breakfast).

#### Prophylaxis versus *H. influenza type b*

Adults:	600 mg every 24 hours x 4 days
Children ( $\geq 1$ month):	20 mg/kg (up to 600 mg) every 24 hours x 4 days
Neonates ( $< 1$ month):	10 mg/kg every 24 hours x 4 days

#### Prophylaxis versus *N. meningitidis*

Adults:	600 mg every 12 hours x 2 days
Children ( $\geq 1$ month):	10 mg/kg (up to 600 mg) every 12 hours x 2 days
Neonates ( $< 1$ month):	5 mg/kg every 12 hours x 2 days



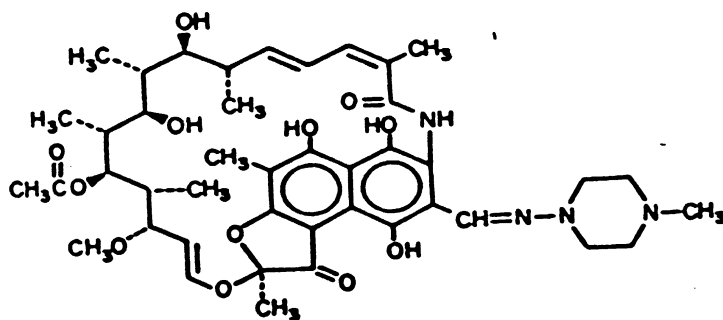
## PHARMACEUTICAL INFORMATION

### Drug Substance

Proper Name: RIFAMPIN

Chemical Name: 5,6,9,17,19,21-hexahydroxy-23-methoxy-2,4,12,16,18,20,22-heptamethyl-8-[N-(4-methyl-1-piperazinyl)formimidoyl]-2,7-(epoxypentadeca [1,11,13] trienimino) naphtho [2,1-*b*]furan-1,11(2H)-dione 21-acetate

Structural formula:  $C_{43}H_{58}N_4O_{12}$



Molecular weight: 822.94

Description: RIFADIN is an orange to red-brown, tasteless, crystalline powder. It is highly soluble in chloroform and methylene chloride, readily soluble in methyl alcohol and ethyl acetate, and poorly soluble in acetone and water. Its melting point is 183-188°C.

### AVAILABILITY OF DOSAGE FORMS

Each capsule contains: RIFADIN (rifampin) 150 mg (opaque maroon and opaque scarlet) or 300 mg (opaque maroon and opaque scarlet).

**Non-medicinal ingredients:** corn starch, D&C Red #28, FD&C Blue #1, FD&C Red #40, gelatin, magnesium stearate, titanium dioxide and white ink. Tartrazine-free.

Available in bottles of 100 capsules.



## MICROBIOLOGY

### Anti-mycobacterial activity

The minimum inhibitory concentration of RIFADIN (rifampin) for *Mycobacterium tuberculosis* *in vitro*, varies considerably with the technique used.

Rifampin inhibited 20 strains of *Mycobacterium tuberculosis* in concentrations of 0.005 to 0.02 µg/mL in 7H-9 broth with Tween 80 and killed all or nearly all of the inoculum in four to eight times greater concentrations. In the same medium without Tween 80, as well as on 7H-10 agar, about 16 to 64 times these amounts were required to produce the same effect (Lorian & Finland, 1969). The susceptibility of a range of typical and atypical *Mycobacterium* to rifampin in Dubos medium is given in the following table:

### IN VITRO

#### ACTIVITY OF RIFAMPIN ON COLLECTION STRAINS OF MYCOBACTERIAL IN DUBOS MEDIUM

STRAIN	MIC µg/mL	STRAIN	MIC µg/mL
<i>M. tuberculosis H<sub>37</sub>Rv</i>	0.2	<i>M. smegmatis ATCC 607</i>	100
<i>M. tuberculosis H<sub>37</sub>Ra</i>	0.2	<i>M. smegmatis 63</i>	200
<i>M. tuberculosis Abate</i>	0.2	<i>M. chelonae NCTC 946</i>	100
<i>M. tuberculosis Schiava</i>	0.1	<i>M. phlei 54</i>	0.1
<i>M. bovis Vallée</i>	0.1	<i>M. phlei Timoteo</i>	0.1
<i>M. bovis BCG</i>	0.05	<i>M. thamnopheos</i>	0.1
<i>M. avium Poulet T<sub>3</sub></i>	10	<i>M. marinum</i>	0.1
<i>M. avium Bang</i>	0.1	<i>M. ranae (fortuitum)</i>	100
<i>M. avium Faisan<sub>2</sub></i>	5	<i>M. kansasii</i>	0.5
<i>M. avium Faisan<sub>4</sub></i>	2	<i>M. scrofulaceum</i>	0.2
<i>M. avium Kirchenberg</i>	20	<i>M. intracellulare</i>	0.1

### Resistance Pattern

The incidence of primary resistance to rifampin in tuberculosis was found not to exceed 0.4 percent (Hobby *et al.*, 1970).

The mutation rates of *M. tuberculosis* and *M. kansasii* toward resistance levels of roughly 10, 50 and 200 times the MIC of rifampin were found to range from 10<sup>-5</sup> to 10<sup>-7</sup>. These relatively high



values show that drug resistance to rifampin is of the streptomycin type. Consequently, monotherapy of human tuberculosis associated with a bacterial count greater than  $10^{-7}$  involves a high risk of selecting rifampin-resistant organisms through the killing of the rifampin-sensitive population. (Manten *et al.*, 1969). Mycobacterial resistant to 80 µg/mL have been observed in patients under treatment for tuberculosis. Resistant strains may sometimes be isolated from patients treated with different drug combinations.

No cross-resistance with any of the known anti-tuberculous drugs or with antibiotics other than those in the rifamycin series has been demonstrated.

## PHARMACOLOGY

### Absorption

Rifampin is readily absorbed from the gastrointestinal tract. Peak blood levels in normal adults vary widely from individual to individual. Peak levels occur between 2 and 4 hours following the oral administration of a 600 mg dose with average peak values of 7-10 µg/mL. Absorption of rifampin is reduced when the drug is ingested with food.

Cumulation was noted upon multiple dosages of RIFADIN (rifampin) 10 mg/kg/day to newborns. Peak values appeared to be delayed in the newborns which were not seen in children up to 18 months of age. It is suggested that the drug is less readily eliminated from the newborn, probably because of the low flow of bile during the first days of life. In all of these children and infants, the main serum level of rifampin corresponded to one-third to one-tenth the levels in adults receiving proportionally the same dose (Acocella *et al.*, 1969).

Absorption is more rapid when rifampin is administered one hour before meals. The following data were reported by Furesz *et al.*, 1967.

### SERUM LEVELS OF RIFAMPIN (µg/mL) AFTER ORAL ADMINISTRATION OF A SINGLE 150 mg DOSE

Pt. No.	2		4		6	
	Fasting	After Meal	Fasting	After Meal	Fasting	After Meal
1	0.37	0	0.32	0	0	0.90
2	2.70	0	1.37	1.60	0.49	1.30
3	1.55	0	0.75	0.14	0.20	0.31
4	1.82	0	1.02	0	0.34	0.16
5	1.77	1.65	0.86	1.00	0.19	0.21

### Distribution

Rifampin is distributed throughout the body and is detectable in many organs and body fluids, including the cerebrospinal fluid. The highest concentrations are present in the liver and bile.



Rifampin is about 80% protein bound. Most of the unbound fraction is not ionized and therefore is diffused freely in tissues.

### Biological Half-Life

In normal subject, the biological half-life of rifampin in serum averages about 3 hours after a 600 mg oral dose, with increases up to 5.1 hours reported after 900 mg dose. Biliary obstruction causes a longer half-life, but kidney blockage does not appear to cause a change. With repeated administration, the half-life decreases and reaches average values of approximately 2 to 3 hours. The half-life does not differ in patients with renal failure at doses not exceeding 600 mg daily and, consequently, no dosage adjustment is required. The half-life of rifampin at a dose of 720 mg daily has not been established in patients with renal failure. Refer to the WARNINGS section for information regarding patients with hepatic insufficiency.

THE FOLLOWING TABLE SUMMARIZES THE HALF-LIFE VALUES REPORTED IN VARIOUS CONDITIONS

<u>SUBJECTS</u>	<u>DOSAGE</u>	<u>BIOLOGICAL HALF-LIFE</u>
Normal Adults*	300 mg	1-3 hours
Obstruction of** common bile duct (1 patient)	300 mg	14 hours before operation 4-5 hours after operation
Anuria** Patient #1 Patient #2 Patient #3	300 mg 300 mg 300 mg	3.1 hours 2.8 hours 1.8 hours
37 children*** 12 children***	10 mg/kg 20 mg/kg	1.2-5.6 hours

\*L. Dettli, 1968; \*\* P. Spring, 1986; \*\*\* B. Krauer, 1968.

### Excretion

Rifampin is eliminated from the blood equally in the urine and feces as unchanged drug and metabolites. Approximately half of the original dose eliminated by the bile is unchanged drug. The proportion of unchanged drug to metabolite is less in the urine than in the bile.

In the presence of complete renal shutdown, the drug is excreted entirely in the bile.

The principal metabolite in man is the biologically active desacetyl rifampin. Its excretion appears to be a dynamically changing picture at all times. This can be seen in the following table, giving the details of fluid levels of desacetyl rifampin after an intravenous dose of 300 mg <sup>14</sup>C-rifampin to a patient with a biliary fistula (Kerberle *et al.*, 1968).



Source	Time After Medication	Rifampin	Percent of total radioactivity in the collection samples as: Desacetyl rifampin
Plasma	0.5 hour	93%	5%
	1 “	77%	12%
	3 hours	62%	18%
Bile	1-1.5 hours	39%	49%
	3-3.5 hours	14%	58%
	4-4.5 hours	8%	79%
	0-24 hours	43% of radioactivity administered is in bile	
Urine	0-4 hours	69%	17%
	0-24 hours	21% of radioactivity administered is in urine	

Desacetylation of rifampin in the body does not substantially modify its anti-mycobacterial activity. In Kirschner's medium, the MIC against M. Tuberculosis varied from 0.1 to 2 µg/mL.

### Transplacental Transfer of Rifampin

Rifampin was administered orally to 20 women at the end of pregnancy in a dose of 300 mg (3.75-5.00 mg/kg) at intervals of 8-12 hours for various lapses of time. Maternal and fetal bloods were collected at parturition. In 5 cases, amniotic fluid was taken. Rifampin (detected by microbiological assay) rapidly crossed both blood-placental barrier and the chorion, being present in large amounts in fetal blood and amniotic fluid. The maternal blood level of rifampin was higher than the known values for male subjects. It was concluded that the fetus eliminates the drug more slowly than does the mother. Nevertheless, the amount of rifampin found in fetal blood is always less than that of the mother and after repeated dosage, the ratio is about 1:3 (Termine et al., 1970).

### TOXICOLOGY

The toxicity of RIFADIN (rifampin) has been determined in various species of laboratory animals. With the exceptions of the dog and rhesus monkey, animals, including the mouse, rat rabbit and cynomolgus monkey satisfactorily tolerated oral doses of rifampin administered over a 2 to 26 week period. These doses were at least ten-fold the therapeutic dose recommended for man.



## Acute Toxicity

The LD<sub>50</sub> within 5 days, in the mouse, rat and rabbit is as follows:

Species	Route	No. of Dose Levels	No. of Animals/ Dose	LD <sub>50</sub> (mg/kg)	95% Confidence Limits (mg/kg)
Mouse	i.p.	4	20	621	595-647
Mouse	oral	5	22	858	829-888
Rat	i.p.	5	16	533	515-552
Rat	oral	6	10	1668	1303-2135
Rabbit	oral	1	8	1550	---

LD<sub>50</sub> based on the statistical method of Litchfield-Wilcoxon i.p. - intraperitoneal (Serralunga, 1967).

Rifampin given in combination with isoniazid or PAS to mice and rats, or with streptomycin sulfate to rats, did not demonstrate any increased acute toxicity of any of the drugs.

## Subacute Toxicity

The cynomolgus monkey, dog and mouse, received oral doses of rifampin for periods of 1-13 weeks and in the rat, both oral and peritoneal administration was given for the same duration. Adverse effects were not observed in mice administered rifampin orally at dose levels inclusive of 100 mg/kg, or in rats administered 100 mg/kg. Parenteral doses of 100 mg/kg similarly did not produce adverse effects in the rat.

In the dog, oral doses of 100 mg/kg were toxic. Abnormalities were seen in behaviour, physical condition, renal and hepatic function tests and hemograms of these animals. After four to six weeks of treatment, death occurred in 50% of the treated animals, characterized pathologically by leukopenia, icterus, ascites, enteritis, and hepatic and renal degeneration. Icterus and moderate renal degeneration were also observed in the remaining animals. Occasional behaviour abnormalities and mild alterations in transaminase, alkaline phosphatase and protein values, were observed following a daily dose of 25 mg/kg of rifampin, administered over a period of four to thirteen weeks. No toxic effect was produced in dosage levels under 25 mg/kg/day.

Short-term (4-13 weeks) studies in the cynomolgus monkey demonstrated mild fatty changes and sporadic sublobar foci of necrosis in the livers of some monkeys; however, this could not be specifically related to drug administration.

A moderate loss of appetite and weight occurred in some monkeys receiving 105 mg/kg over a 90-day period; there were no other apparent toxic effects of the drug.



Rifampin given for a period of thirty days, alone and in combination with ethambutol, isoniazid, PAS or streptomycin sulfate to rats, in drug ratios proposed for human clinical trials, showed that the rat can satisfactorily tolerate oral doses of rifampin and of rifampin combinations without developing significant adverse effects.

### **Chronic Toxicity**

#### **Rats**

Six-month chronic toxicity studies using doses of rifampin up to 200 mg/kg/day had no effect on body weight, hematology or gross and microscopic pathology of the animals.

#### **Dogs**

Six-month studies showed the same toxic effects as those seen in the short-term studies. Oral doses of 25 or 50 mg/kg/day caused emesis, anorexia, diarrhea, slight depression and ataxia; three animals died during the first seven weeks of treatment. Serum transaminase, alkaline phosphatase and bromosulphalein retention values were increased, while the albumin, total protein, erythrocyte and hemoglobin levels were decreased. Gross and histomorphologic examinations revealed that animals given 25 and 50 mg/kg/day had a hemorrhagic enteritis, degenerative hepatic and renal alterations, but these changes were not evident in the group given 12.5 mg/kg/day.

#### **Cynomolgus Monkeys**

In a six month study, oral doses of 15, 45, 75 and 105 mg/kg/day were administered. Some monkeys exhibited emesis and lethargy in the early phase of treatment. Hematologic, clinical chemistry and urinalysis examination results of treated animals were the same as those of the control animals, with the exception of one monkey on 105 mg/kg/day, which had an elevated alkaline phosphatase. The sacrificed animals failed to reveal any evidence of systemic toxic effects from administration of the drug following gross and microscopic examinations.

#### **Rhesus Monkeys**

During the first 30 days of an 18-month study of rifampin, administered orally in doses of 40, 80 and 120 mg/kg, nephrotoxicity developed. Kidney examinations showed many of the characteristics of acute and/or subacute glomerulonephritis, suggestive of target-organ effect. This target-organ-like effect on the kidney seems to be related specifically to the rhesus monkey, as cynomolgus monkeys, studied for a longer period of time, did not exhibit a similar toxicity.

In another study, doses of 40, 80 and 120 mg/kg administered orally to cynomolgus monkeys, failed to induce a renal target-organ effect, or other significant drug-related adverse reactions.



## **Carcinogenicity**

There are no known human data on long-term potential for carcinogenicity, mutagenicity, or impairment of fertility. A few cases of accelerated growth of lung carcinoma have been reported in man, but a causal relationship with the drug has not been established. An increase in the incidence of hepatomas in female mice (of a strain known to be particularly susceptible to the spontaneous development of hepatomas) was observed when rifampicin was administered in doses two to ten times the average daily human dose for 60 weeks followed by an observation period of 46 weeks. No evidence of carcinogenicity was found in male mice of the same strain, mice of a different strain, or rats under similar experimental conditions.

Rifampin has been reported to possess immunosuppressive potential in rabbits, mice, rats, guinea pigs, human lymphocytes *in vitro*, and humans. Antitumor activity *in vitro* has been shown with rifampin.

## **Mutagenicity**

Increased frequency of chromosomal aberrations was observed *in vitro* in lymphocytes obtained from patients treated with combinations of rifampin, isoniazid, and pyrazinamide and combinations of streptomycin, rifampin, isoniazid, and pyrazinamide.

There was no evidence of mutagenicity in bacteria, *Drosophila melanogaster*, or mice. An increase in chromatid breaks was noted when whole blood cell cultures were treated with rifampin.

## **Fetal Toxicity and Teratogenicity**

### **Mice**

No maternal toxic effects were demonstrated when pregnant mice were given oral doses of 50 to 200 mg/kg of rifampin. A decrease in the average weight was noted in the fetuses of test animals receiving oral doses of 150 mg/kg and exencephaly was seen in 4% of the fetuses. An increase in the incidence of resorptions and a decrease in fetal numbers also occurred at an oral dose of 200 mg/kg. Skeletal abnormalities characterized by an absence of ossification also occurred on doses of 200 mg/kg. On oral doses of 50-100 mg/kg and 150-200 mg/kg, dose-dependent cleft palate abnormalities occurred with a range of 0.6%-0.72% and 3.69%-18.2%, respectively.

### **Rats**

A maternal toxic effect occurred with oral doses of 200 mg/kg and over. Reductions in the number of fetuses and fetal weights, with an increase of resorptions, were seen following oral doses of 100 and 200 mg/kg. Spina bifida was observed with oral doses of 150 mg/kg and the incidence of this teratogenic abnormality increased with higher doses. On a dosage of 200 mg/kg, rhachischisis in the lumbosacral region and incomplete ossification centre development, particularly of the sternbrae, were noted.



In one study, following administration of rifampin in doses of 200 mg/kg, microcephaly was detected in 0.4% of the fetuses and microphthalmia in 1.2%. No adverse effect on lactation or on the suckling young was noted when oral doses of 50 and 100 mg/kg had been administered to female rats during the last trimester of pregnancy.

### **Rabbits**

Doses of rifampin, 200 and 300 mg/kg/day produced a high mortality rate of the dams and toxic effects including anorexia, weight loss and abortion were produced in pregnant rabbits on oral doses of 75 and 150 mg/kg. Embryotoxic effects resulting in increased resorptions were evidenced following oral doses of 150 mg/kg. This species did not demonstrate any major malformations which could be attributed to the drug.



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**READ THIS FOR SAFE AND EFFECTIVE USE OF YOUR MEDICINE**  
**PATIENT MEDICATION INFORMATION**

**RIFADIN**  
**Rifampin capsules, 150 mg and 300 mg**

Read this carefully before you start taking **RIFADIN** and each time you get a refill. This leaflet is a summary and will not tell you everything about this drug. Talk to your healthcare professional about your medical condition and treatment and ask if there is any new information about **RIFADIN**.

**What is RIFADIN used for?**

Rifadin capsules contain a medicine called rifampin. It belongs to a group of medicines called anti-bacterials.

Rifadin capsules are used to treat the following bacterial infections:

- Tuberculosis (also known as TB) alongside other medicines

Rifadin capsules can also be used to stop the following infections from developing;

- Meningitis (infection of the membranes in the brain and spinal cord)
- Haemophilus influenza (flu)

Anti-bacterial drugs like Rifadin treat only bacterial infections. They do not treat viral infections such as the common cold. Although you may feel better early in treatment, Rifadin should be used exactly as directed. Misuse or overuse of Rifadin could lead to the growth of bacteria that will not be killed by Rifadin (resistance). This means that Rifadin may not work for you in the future. Do not share your medicine.

**How does RIFADIN work?**

It works by killing the bacteria that cause infections.

**What are the ingredients in RIFADIN?**

Medicinal ingredients: Each capsule contains rifampin

Non-medicinal ingredients: Corn starch, D&C Red #28, FD&C Blue #1, FD&C Red #40, gelatin, magnesium stearate, titanium dioxide and white ink. Tartrazine-free

**RIFADIN comes in the following dosage forms:**

Capsules, 150 mg and 300 mg



**Do not use RIFADIN if:**

- You are allergic (hypersensitive) to
  - Rifampin or any rifamycin antibiotics
  - any of the other ingredients of the Rifadin capsules (See What are the ingredients in Rifadin)
- You have yellowing of the skin and eyes (jaundice)
- You are breastfeeding
- You are taking a saquinavir/ritonavir combination for HIV infection.
- The person taking this medicine is a premature and or a newborn infant

**To help avoid side effects and ensure proper use, talk to your healthcare professional before you take RIFADIN. Talk about any health conditions or problems you may have, including if you:**

- You have liver problems
- You have any kidney problems
- You have diabetes. Your diabetes may become more difficult to control while taking this medicine
- You are under weight or malnourished
- You wear contact lenses. Taking Rifadin capsules may permanently stain soft contact lenses
- You have porphyria (a genetic enzyme disorder that causes symptoms affecting the skin or nervous system)
- The person taking this medicine is a child
- You are using hormonal birth control. The effectiveness of the birth control may be decreased. It may be necessary to use another method of birth control.
- If you are pregnant or planning to become pregnant.

**Other warnings you should know about:****Blood Tests**

Your doctor will need to check your blood before you take this medicine. This will help your doctor know if any changes happen to your blood after taking this medicine. You may also need to have regular blood tests to check how your liver is working.



**Tell your healthcare professional about all the medicines you take, including any drugs, vitamins, minerals, natural supplements or alternative medicines.**

**The following may interact with RIFADIN:**

- Antacids used for indigestion. Take Rifadin at least 1 hour before taking antacids
- Medicines for high blood pressure (e.g. ACE inhibitors, angiotensin II receptor antagonist).
- Medicines for heart problems or to control your heartbeat (e.g., disopyramide, mexiletine, quinidine, tocainide, propafenone)
- Medicines used to thin the blood such as warfarin
- Medicines used to lower cholesterol (e.g., statins, clofibrates)
- Medicines for thought disorders known as “antipsychotics” such as haloperidol
- Medicines to calm or reduce anxiety (hypnotics, anxiolytics)
- Medicines to help you sleep (barbiturates)
- Medicines used for epilepsy such as phenytoin
- Some medicines used for depression such as amitriptyline and nortriptyline
- Riluzole - used for motor neurone disease
- Some medicines used for viral infections such as zidovudine, indinavir, efavirenz and saquinavir
- Medicines used for fungal infections (e.g. antifungals such as fluconazole, itraconazole, ketoconazole)
- Medicines used for bacterial infections (antibiotics such as telithromycin, clarithromycin, doxycycline)
- Medicines used for lowering your immune system such as cyclosporin and tacrolimus
- Praziquantel - used for tapeworm infections
- Atovaquone - used for pneumonia
- Some hormone medicines (estrogen, systemic hormones, progestogens) including birth control pills (oral contraceptives)
- Some hormone medicines (anti-estrogens) used for breast cancer or endometriosis such as tamoxifen, toremifene
- Levothyroxine (thyroid hormone) used for thyroid problems
- Irinotecan - used for cancer
- Medicines used for pain such as codeine, morphine, fentanyl or pethidine
- Corticosteroids used for inflammation such as hydrocortisone, betamethasone and prednisolone
- Methadone - used for heroin withdrawal
- Medicines used for diabetes
- Medicines used to relax muscles before surgery (anesthetics) such as halothane
- Some medicines used for nausea or vomiting such as ondansetron
- Quinine - used for malaria
- Theophylline - used for wheezing or difficulty in breathing



### How to take RIFADIN

Take Rifadin on an empty stomach. This means at least 1 hour before breakfast.

### Usual dose:

### Treatment of tuberculosis

**Adults:** 600 mg in a single daily dose. The dose may need to be reduced in some patients. Follow your doctor's advice about how to take RIFADIN, when to take it, and how long to take it.

**Children ( $\geq 5$  years):** 10 to 20 mg/kg not to exceed 600 mg/day.

### Prophylaxis versus *H. influenza type b*

Adults:	600 mg every 24 hours for 4 days
Children ( $\geq 1$ month):	20 mg per kilogram (up to 600 mg) every 24 hours for 4 days
Neonates ( $< 1$ month):	10 mg per kilogram every 24 hours for 4 days

### Prophylaxis versus *N. meningitidis*

Adults:	600 mg every 12 hours for 2 days
Children ( $\geq 1$ month):	10 mg per kilogram (up to 600 mg) every 12 hours for 2 days
Neonates ( $< 1$ month):	5 mg per kilogram every 12 hours for 2 days

### Overdose:

If you take more Rifadin capsules than you should, tell a doctor or go to a hospital emergency department straight away. Take the medicine pack with you. This is so the doctor knows what you have taken.

You may feel sick (nausea), be sick (vomiting), have stomach pain, itching or a headache. You may also feel tired, sleepy, dizzy or light-headed. Other signs of taking too much includes swelling of the face, eyes or eyelids, fast heartbeat, irregular heartbeats, seizure and heart attack.

If you think you have taken too much RIFADIN, contact your healthcare professional, hospital emergency department or regional poison control centre immediately, even if there are no symptoms.

### Missed Dose:

If you forget a dose, take it as soon as you remember it. However, if it is nearly time for the next dose, skip the missed dose. Do not take a double dose to make up for the forgotten capsules.



## What are possible side effects from using RIFADIN?

These are not all the possible side effects you may feel when taking RIFADIN. If you experience any side effects not listed here, contact your healthcare professional.

You may notice a discoloration (yellow, orange, red, brown) in your urine, sweat, phlegm (sputum), saliva, tears or teeth. This is quite common and you need not worry. However, the discoloration may permanently stain soft contact lenses. The discoloration in tears may last for some time after you have stopped taking Rifadin. Tooth discoloration may be permanent.

Serious side effects and what to do about them			
Symptom / effect	Talk to your healthcare professional		Stop taking drug and get immediate medical help
	Only if severe	In all cases	
<b>RARE</b> You have an allergic reaction. The signs may include: a rash, swallowing or breathing problems, wheezing, swelling of your lips, face, throat or tongue.			✓
You experience symptoms such as severe diarrhea (bloody or watery) with or without fever, abdominal pain, or tenderness, you may have <i>Clostridium difficile</i> colitis (bowel inflammation).			✓
You have a fever and yellowing of the skin or whites of the eyes, feel tired, weak or generally unwell, loss of appetite (anorexia), feeling sick (nausea), being sick (vomiting). These may be early signs of liver problems.			✓
Drug rash with eosinophilia and systemic symptoms (DRESS) including death, with any combination of red itchy rash with blisters and peeling of the skin and /or of the lips, eyes, mouth, nasal passages or genitals. It often goes with fever, chills, headache, cough, body aches or joint pain. You may have less or dark urine, yellow skin or eyes.			✓
You bruise more easily than usual. Or you may have a painful rash of dark red spots under the skin which do not go away when you press on them (purpura). This could be because of a serious blood problem.			✓
You have chills, tiredness, unusually pale skin colour, shortness of breath, fast heartbeat or dark coloured urine. This could be signs of a serious type of anemia.			✓
You have blood in your urine or an increase or decrease in amount of urine you produce. You			✓



Serious side effects and what to do about them			
Symptom / effect	Talk to your healthcare professional		Stop taking drug and get immediate medical help
	Only if severe	In all cases	
may also get swelling, especially of the legs, ankles or feet. This may be caused by serious kidney problems.			
You have a sudden severe headache. This could be a sign of bleeding in the brain.			✓
Shortness of breath and wheezing.			✓
You get confused, sleepy, cold clammy skin, shallow or difficult breathing, a racing heartbeat or your skin is paler than normal. These could be signs of shock.			✓
You get more infections more easily than normal. Signs include fever, sore throat or mouth ulcers. This could be because you have a low number of white blood cells.			✓
You have bleeding from your nose, ear, gums, throat, skin or stomach. Signs may include a feeling of tenderness and swelling in your stomach, purple spots on your skin and black or tar-like stools			✓
Mental problems with unusual thoughts and strange visions (hallucinations)		✓	
Flu-like symptoms including chills, fever, headache, dizziness and bone pains		✓	
Skin flushing or itching	✓		
Irregular periods	✓		
Diarrhea or stomach discomfort	✓		
Loss of appetite (anorexia)	✓		
Headache	✓		

If you have a troublesome symptom or side effect that is not listed here or becomes bad enough to interfere with your daily activities, talk to your healthcare professional.



### **Reporting Side Effects**

You can report any suspected side effects associated with the use of health products to Health Canada by:

- Visiting the Web page on Adverse Reaction Reporting (<http://www.hc-sc.gc.ca/dhp-mps/medeff/report-declaration/index-eng.php>) for information on how to report online, by mail or by fax; or
- Calling toll-free at 1-866-234-2345.

*NOTE: Contact your health professional if you need information about how to manage your side effects. The Canada Vigilance Program does not provide medical advice.*

### **Storage:**

Store between 15-30 °C.

Keep out of reach and sight of children.

### **If you want more information about RIFADIN:**

- Talk to your healthcare professional
- Find the full product monograph that is prepared for healthcare professionals and includes this Patient Medication Information by visiting the Health Canada website (<http://hc-sc.gc.ca/index-eng.php>); the manufacturer's website [www.sanofi.com](http://www.sanofi.com), or by calling 1-800-265-7927.

This leaflet was prepared by sanofi-aventis Canada Inc.

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