



# Rifampicin Sodium PRODUCT DATA SHEET

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<b>Product Name:</b>	Rifampicin Sodium
<b>Product Number:</b>	R040
<b>CAS Number:</b>	38776-75-9
<b>Molecular Formula:</b>	$C_{43}H_{58}N_4NaO_{12}$
<b>Molecular Weight:</b>	844.93
<b>Form:</b>	Powder
<b>Appearance:</b>	Fine or slightly granular red powder
<b>Solubility:</b>	Water: soluble, Ethanol: soluble
<b>Water Content (Karl Fischer):</b>	12.0 - 17.0%
<b>Absorbance:</b>	190 - 210
<b>pH:</b>	4.5 - 8.0
<b>Storage Conditions:</b>	2-8°C
<b>Description:</b>	<p>Rifampicin sodium (Rifampin) is a semisynthetic antimicrobial compound derived from rifamycin SV and originally developed by the Lepetit group in 1965. Rifampicin shows activity against gram-positive bacteria, particularly mycobacteria like tuberculosis, some gram-negative anaerobic bacteria, protozoa, fungi, and poxviruses.</p> <p>Rifampicin sodium inhibits the assembly of DNA and protein into mature virus particles. It inhibits initiation of RNA synthesis by binding to <math>\beta</math>-subunit of RNA polymerase, which results in cell death.</p> <p>Rifampicin sodium has been shown to inhibit <math>\alpha</math>-synuclein fibrillation and disaggregate fibrils in a concentration-dependent manner. Rifampicin can activate pregnane X receptor (PXR), which affects cytochrome P450, and the activity of glucuronosyltransferases and P-glycoprotein. Rifampicin has been shown to enhance CYP2C-mediated metabolism, affect compounds that are transported by P-glycoprotein and metabolized by CYP3A4.</p> <p>Rifampicin sodium is an immunosuppressive agent.</p> <p>For a safer and easier to use Rifampicin, try TOKU-E's ready-made <a href="#">Rifampicin Solution (10 mg/mL in water)</a>.</p> <p>Synonyms: RFP, Rifampin, 3-(((4-Methyl-1-piperazinyl)imino)methyl)rifamycin SV, NIH 10782, NSC 113926</p>
<b>Mechanism of Action:</b>	Rifampicin sodium targets prokaryotic DNA dependent RNA polymerases which prevent subsequent RNA transcription and protein translation.

**Spectrum:**

Rifampicin is a broad-spectrum antibiotic with a wide range of activity including:

- Gram-positive aerobic bacteria, particularly *Staphylococcus spp* and *Rhodococcus equi*
- Brucella and some other fastidious organisms are susceptible but Gram-negative bacteria more generally are resistant
- Gram-positive and Gram-negative anaerobic bacteria are inhibited at low concentrations, including *Bacteroides fragilis*
- *Chlamydomphila* and *Rickettsia* are susceptible
- *Mycobacterium tuberculosis*: activity is high against this organism but most other mycobacteria are resistant
- Some protozoa
- Some fungi and poxviruses

**Microbiology Applications**

Rifampicin sodium is commonly used in bacterial recombinant protein expression to inhibit bacterial RNA polymerase activity and synthesis of host bacterial proteins. Rifampicin sodium can also be used as a selective agent to isolate *Campylobacter jejuni*.

**Plant Biology Applications**

Rifampicin has been tested in Jerusalem artichoke tuber explants by adding 10 to 50 µg/ml to the tissue culture medium. At 50 µg/ml no bacterial infection was detectable, without affecting cell division rates, cytodifferentiation and DNA synthesis. As a result Rifampicin was used as antibacterial in the following experiments of this university department (Philips, 1981).

Rifampicin is a selective inhibitor of chloroplast RNA polymerase and can be used to study chloroplast-level DNA transcription in plants.

**References:**

"Rifampin: Mechanisms of Action and Resistance." *Oxford Journals* (1983): n. pag. *Clinical Infectious Diseases*. Web. 21 Aug. 2012.

"Philips R., Arnott S.M. and K aplan S.E., 1981, Antibiotics in plant tissue culture: rifampicin effectively controls bacterial contaminants without affecting the growth of short-term explant cultures of *Helianthus tuberosus*. *Plant Science Letters*, 21 (1981) 235-240.

Li, T., & Chiang, J. Y. (2006). Rifampicin induction of CYP3A4 requires pregnane X receptor cross talk with hepatocyte nuclear factor 4alpha and coactivators, and suppression of small heterodimer partner gene expression. *Drug metabolism and disposition: the biological fate of chemicals*, 34(5), 756-64.

Jill E Maddison, A David J Watson, Jonathan Elliott (2008) Chapter 8 - Antibacterial drugs, *Small Animal Clinical Pharmacology* (Second Edition), 148-185.

Bassi, L., Berardino, L., Arioli, V., Silvestri, L., & Lignière, E. (1973). Conditions for Immunosuppression by Rifampicin. *The Journal of Infectious Diseases*, 128(6), 736-744.