

Antimicrobial Activity of *Rhus coriaria* L. Leaf Extract

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The antibacterial activity of *Rhus coriaria* leaf methanol extract was assayed against Gram-positive and Gram-negative bacteria; antimycotic activity was assayed against some *Candida* species. MICs were determined by a broth microdilution assay in microlitre plates using Mueller–Hinton medium. MBCs were determined by plating 0.01 mL samples from clear 1 mL tubes onto agar plates. © 1998 John Wiley & Sons, Ltd.

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INTRODUCTION

We report the preliminary results of antibacterial and antimycotic activities of *Rhus coriaria* L. leaf extract.

Rhus coriaria L. (Anacardiaceae) found on the coast, in the hedges and in the woods of the mediterranean region, is an evergreen shrub or sapling up to 4 m tall. Numerous drugs obtained from different species of *Rhus* genus for their medicinal properties are included in the Pharmacopoeias of various countries (Imbesi, 1964).

Among these drugs, *Rhus coriaria* leaves, which contain tannins, flavonoids and an essential oil (El Sissi *et al.*, 1966; Van Loo *et al.*, 1988; Kuruku *et al.*, 1993) are used for their astringent properties, and we decided to examine them for antimicrobial activity.

MATERIALS AND METHODS

Plant material. Leaves of *Rhus coriaria* L. were collected in the neighbourhood of Messina in March 1996.

A voucher specimen of the plant was deposited in the herbarium of the Pharmaco-Biological Department of the University of Messina (Italy).

The fresh material was immediately lyophilized and powdered.

Preparation of extract. The methanol extract of *Rhus coriaria* L. was assayed. For the preparation of methanol extract exhaustive extraction of 10 g of *Rhus coriaria* L. leaves was carried out at 60°C in a water bath using methanol as solvent.

The mixture was filtered and the organic solvent removed under vacuum, dissolved in DMSO and diluted to 250 000 mg/L in volumetric flasks.

Microorganisms. A total of 27 fresh clinical isolates identified by conventional procedures (National Committee for Clinical Laboratory Standards, 1995) and ATCC standard strains were used (Table 1).

Minimum inhibitory concentration (MIC). MICs were determined by a broth microdilution assay in microtitre

Table 1. MICs and MBCs (mg/L) of *Rhus coriaria* L. leaf extracts against 27 strains of standard and clinically isolated microorganisms

Standard strains	MIC	MBC
<i>Escherichia coli</i> ATCC 25922	312	625
<i>Escherichia coli</i> ATCC 30213	156	312
<i>Pseudomonas aeruginosa</i> NCTC 10662	625	1250
<i>Staphylococcus aureus</i> ATCC 29213	156	312
<i>Enterococcus faecalis</i> ATCC 29212	156	312
<i>Bacillus subtilis</i> ATCC 6603	78	156
<i>Candida albicans</i> ATCC 3183	625	1250
Clinical strains		
<i>Escherichia coli</i> 66	625	625
<i>Escherichia coli</i> 61	625	1250
<i>Escherichia coli</i> 64	312	312
<i>Escherichia coli</i> 71	312	625
<i>Pseudomonas aeruginosa</i> 052	312	625
<i>Pseudomonas aeruginosa</i> 020	625	625
<i>Klebsiella pneumoniae</i> 62	312	625
<i>Klebsiella pneumoniae</i> 006	312	625
<i>Klebsiella pneumoniae</i> 028	312	625
<i>Klebsiella pneumoniae</i> 017	625	625
<i>Klebsiella pneumoniae</i> 100	312	312
<i>Enterobacter cloacae</i> 02	312	625
<i>Enterobacter cloacae</i> 201	312	625
<i>Enterobacter agglomerans</i> 021	625	625
<i>Enterobacter agglomerans</i> 010	312	625
<i>Enterobacter agglomerans</i> 099	312	625
<i>Citrobacter freundii</i> 081	312	625
<i>Staphylococcus aureus</i> 16	156	312
<i>Staphylococcus aureus</i> 22	312	312
<i>Staphylococcus aureus</i> 050	156	312

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plates according to the NCCLS reference methods (National Committee for Clinical Laboratory Standards, 1995; Murray *et al.*, 1995) using Mueller–Hinton broth at pH 7.3.

A total of 12 concentrations were performed from 25 000 to 12.25 mg/L.

A suspension of microorganism (1 µL) was added to each well containing the drug and to the control well without drug. The final concentration of each microorganism in each well was 5×10^5 CFU/mL.

The present investigation was carried out using 6 bacterial and 1 mycotic standard strains and 20 clinical isolates of Gram-positive and Gram-negative bacteria.

The MIC is defined as the lowest concentration of drug that inhibits the visible growth after 18–24 h.

Minimum bactericidal concentration (MBC). MBCs were determined by plating 0.01 mL samples from clear 1 mL tubes onto plant extract-free agar plates.

The MBC was the concentration at which there was a 99.9% reduction of the original inoculum (Amsterdam, 1991).

RESULTS AND DISCUSSION

The results of the antimicrobial activity (MICs and MBCs) of *Rhus coriaria* L. leaf extract against the microorganisms tested are shown in Table 1.

The inhibitory action of this plant extract was high against both Gram-negative and Gram-positive bacteria.

Regarding Gram-negative bacteria, *Rhus coriaria* L. leaf methanol extract was most active against *Escherichia coli* ATCC 30213 (MIC = 156 mg/L); for all other Gram-negative strains MIC values ranged from 312 mg/L to 625 mg/L.

The extract also showed very good activity against Gram-positive microorganisms inhibiting *Bacillus subtilis* strain at 78 mg/L and *Staphylococcus aureus* strains (both standard and clinical isolates) at 156 and 312 mg/L. *Escherichia faecalis* was also inhibited at 156 mg/L. *Candida albicans* ATCC 3183 was susceptible at 625 mg/L.

The significant antimicrobial activity of *Rhus coriaria* L. leaf methanol extract was evidenced by MBCs results.

In all cases, MBC values are generally the same or twice the MIC values for all microorganisms.

The present study has shown that *Rhus coriaria* L. leaf extract is active against both Gram-negative and Gram-positive bacteria.

The extract has excellent bactericidal activity with MBCs being equal to twice the MICs.

The results suggest further investigations into the antimicrobial activity of *Rhus coriaria* L. leaf extract, in particular against *Staphylococcus aureus* and enteric Gram-positive and Gram-negative bacteria (which were frequently resistant to drugs commonly used in therapy) should be carried out.

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