

Potential use of Euphorbia hirta and Euphorbia jolkinii extracts as antimicrobial agents agaisnt Pseudomonas aeruginosa, Morganella morganii, and Klebsiella pneumoniae

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I. Introduction

Euphorbia hirta and *Euphorbia jolkinii* are two plants belonging to the Euphorbiaceae family. Euphorbia hirta has been traditionally used in the folk medicine of different cultures,



awakening the interest of the scientific community because of the link of its **bioactivities** to the **compounds** present in this species. Therefore, this study was run considering the need to find novel natural antimicrobial extracts to be used in the industry replacing the synthetic ones by testing both E. hirta and E. jolkinii ethanol extracts.

II. Objective and methodology

Considering the need of society to find **new** antibacterial extracts and the potential use of these plants for this goal, the **antibacterial activity** of ethanol extracts (10 mg/mL) against eight food and nine clinical bacteria have been characterized.

Unsaturated fatty acids Organic acids Euphorbia jolkinii Euphorbia hirta

By a **colorimetric assay**, minimum inhibitory concentration (MIC) and minimum bactericidal concentration (MBC) were determined, being streptomycin (1 mg/mL), ampicillin (10 mg/mL), and methicillin (1 mg/mL), the antibiotics used as controls.



III. Results

Table 1. Antibacterial activity of *E. hirta* and *E. jolkinii* tested in both clinical and food contaminant bacteria

ANTIBACTERIAL ACTIVITY

			· ·	
F hirta	F jalkinii	Amnicillin	Iminonom	Vancomucin

As it is shown in Table 1, results showed an antibacterial activity comparable to or superior to ampicillin when extracts were tested against three clinical bacteria (Pseudomonas aeruginosa, Morganella morganii, and Klebsiella pneumoniae). Thus, the MIC values obtained when *E. hirta* extract was used were: 5, 1.25, and <10 for K. pneumoniae, M. morganii, and P. aeruginosa, respectively. For the E. jolkinii extract, MIC values were 1.25 for the three bacteria, while MIC values for ampicillin were 10, <10, and <10 for K. pneumoniae, M. morganii, and P. *aeruginosa*, respectively.

		E. hirta	E. jolkinii	Ampicillin	Imipenem	Vancomycin
		(10 mg/mL)	(10 mg/mL)	(10 mg/mL)	(1 mg/mL)	(1 mg/mL)
Escherichia coli	MIC	5	2.5	<0.15	<0.0078	ND
	MBC	>10	>10	< 0.15	< 0.0078	ND
Klebsiella pneumoniae	MIC	5	1.25	10	<0.0078	ND
	MBC	>10	>10	>10	< 0.0078	ND
Managanalla managanii	MIC	1.25	1.25	>10	<0.0078	ND
Morganella morganii	MBC	>10	>10	>10	<0.0078	ND
Development in a Lilia	MIC	2.5	0.6	< 0.15	< 0.0078	ND
Proteus mirabilis	MBC	>10	>10	< 0.15	< 0.0078	ND
Providomonan acmusinona	MIC	>10	1.25	>10	0.5	ND
Pseudomonas aeruginosa	MBC	>10	>10	>10	1	ND
Entene co cour fa co alia	MIC	0.6	1.25	< 0.15	ND	< 0.0078
Enterococcus faecalis	MBC	>10	>10	< 0.15	ND	< 0.0078
Listeria monocytogenes	MIC	2.5	1.25	< 0.15	< 0.0078	ND
	MBC	>10	>10	< 0.15	< 0.0078	ND
MDCA	MIC	2.5	0.3	< 0.15	ND	0.25
MRSA	MBC	>10	>10	< 0.15	ND	>0.5
Propionibactarium acres	MIC	2.5	2.5	ND	ND	0.07
Propionibacterium acnes	MBC	>10	>10	ND	ND	5
		Food cont	aminant bacteria			
		E. hirta (10 mg/mL)	E. jolkinii (10 mg/mL)	Streptomycin (1 mg/mL)	Methicillin (1 mg/mL)	Ampicillin (10 mg/mL)
	MIC	5	2.5	0.007	ND	0.15
Enterobacter cloacae	MBC	>10	>10	0.007	ND	0.15
Factorial in a li	MIC	5	10	0.01	ND	0.15
Escherichia coli	MBC	>10	>10	0.01	ND	0.15
Pseudomonas aeruginosa	MIC	>10	>10	0.06	ND	0.63
	MBC	>10	>10	0.06	ND	0.63
Salmonella enterocolitica	MIC	10	10	0.007	ND	0.15
	MBC	>10	>10	0.007	ND	0.15
Vanainia antana ditian	MIC	2.5	2.5	0.007	ND	0.15
Yersinia enterocolitica	MBC	>10	>10	0.007	ND	0.15
D	MIC	5	5	0.007	ND	ND
Bacillus cereus	MBC	>10	>10	0.007	ND	ND

These results open a new potential use of ethanol extracts of both E. hirta and E. jolkinii in the food industry as natural inhibitors of different microorganisms.



Ductitus cercus	MBC	>10	>10	0.007	ND	ND
Listeria monocytogenes	MIC	5	1.25	0.007	ND	0.15
	MBC	>10	>10	0.007	ND	0.15
Staphylococcus aureus	MIC	2.5	0.6	0.007	0.007	0.15
	MBC	>10	>10	0.007	0.007	0.15

IV. Conclusions and future directions

Considering the data provided, both extracts could be used as natural antimicrobial alternatives since they showed high antibacterial activity against three clinical bacteria, with E. jolkinii having the best results.

Myricetin enzymes

Figure 1. Antibacterial activity mechanism of both plant extracts and the compounds likely to have this activity.

> The potential industrial application of both extracts

as natural antimicrobials

should be tested in the future

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