

Research Article

Antimicrobial Activity of Flavonoids against Extended-Spectrum β -Lactamase (ES β L)-Producing *Klebsiella pneumoniae*

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Abstract

Purpose: In the present study, six flavonoids (5,7-dimethoxyflavanone-4'-O- β -D-glucopyranoside, 5,7-dimethoxyflavanone-4'-O-[2''-O-(5'''-O-trans-cinnamoyl)- β -D-apiofuranosyl]- β -D-glucopyranoside, naringenin-7-O- β -D-glucopyranoside, 5,7,3'-trihydroxy-flavanone-4'-O- β -D-glucopyranoside, rutin, and nicotiflorin) isolated from *Galium fissurense*, *Viscum album ssp. album* and *Cirsium hypoleucum* were screened against extended-spectrum β -lactamase producing multidrug-resistant (trimetoprim-sulphametoxazole, sulbactam-ampicillin, clavulonate-amoxicillin, ceftriaxon, cefepime, imipenem, ceftazidime, tobramycin, gentamicin, ofloxacin, ciprofloxacin) bacteria *Klebsiella pneumoniae* (ES β Ls).

Methods: We performed susceptibility testing according to the Clinical and Laboratory Standards Institute (CLSI, formerly NCCLS) and used an inhibition endpoint for determination of the minimum inhibition concentrations (MICs).

Results: All the flavonoids showed *in vitro* antimicrobial activity against all the isolated strains of *K. pneumoniae* similar to the control antibacterial (ofloxacin) at the concentrations of 32 - 64 μ g ml⁻¹; another control, ampicillin, had no activity. Since, ES β L-producing strains are known to be resistant to all β -lactam antibiotics, our results fall notably within the concentration range for antimicrobial activity.

Conclusion: To the best of our knowledge, this is the first report of the study of the activity of these flavonoids against (ES β L)-producing *K. pneumoniae* and may throw light to the low-toxicity of flavonoids, and their potentials for developing therapies for infections caused by ES β L-producing bacteria in the future. Further work is under investigation to identify their precise antibacterial mechanism.

Keywords: Antimicrobial activity, ES β Ls, Flavonoids, *Klebsiella pneumoniae*, Ofloxacin, Ampicillin.

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INTRODUCTION

Multiple drug resistance has significantly increased in recent years. The existence of enzymes of extended-spectrum β -lactamases (ES β Ls) producing organism that are resistant to virtually all β -lactam antibiotics have been reported¹. ES β Ls are plasmid-mediated class A enzymes commonly found in the family Enterobacteriaceae, mainly *K. pneumoniae*². These microorganisms are Gram-negative rods that cause bacterial pneumonia and hospital-acquired infections³. The increase in ES β L-producing organisms is sure to create significant therapeutic problems in the future. The available treatment regimens for infections caused by ES β L-producing bacteria are not always effective. Therefore, it is necessary to discover new antimicrobial compounds against ES β L-producing *K. pneumoniae* strains.

Flavonoids occur as aglycones, glycosides and methylated derivatives and are widely distributed in the plant kingdom. They have been reported to possess a variety of biological activities including antiallergic, antidiabetic, antiinflammatory, antiviral, antiproliferative and anticarcinogenic, hepatoprotective, and antioxidant activities. Since these secondary metabolites are synthesized by plants in response to microbial infection, it should not be surprising that they have been found *in vitro* to be effective antimicrobial substances against a wide array of microorganisms⁴⁻⁶. Also, their antimicrobial activities against some Gram-negative and Gram-positive bacteria have been reported in many papers². Therefore, the flavonoids may be promising new class compounds in antimicrobial therapy.

In this study, six flavonoids isolated from three medicinal plants, which are used in Turkish folk medicine, were screened against *K. pneumoniae*. *Galium fissurense* Ehrend. & Schönb.-Tem. (Rubiaceae), which is an endemic plant in Turkey, has long been used in folk medicine for a variety of purposes, especially as a diuretic, astringent, choleric and in the treatment of some stomach, gout and epilepsy⁷. *Viscum album* L. (Loranthaceae) is reported to have various

biological activities such as hypoglycemic, anti-inflammatory, anti-viral, anti-atherogenic, and analgesic activities. On the other hand, we have previously shown that the extracts of *V. album* ssp. *album* have inhibitory effects on IL-1 α ⁸. In folk medicine, the decoction of *Cirsium* sp. (Asteraceae) seeds and roots is used for healing hemorrhoids and the flowers are a good remedy for peptic ulcer. In addition, the stem is also used for treating ailments such as cough and bronchitis in Anatolia⁹. In a previous work of our group, the antimicrobial activities of various *C. hypoleucum* DC. extracts (methanol, n-hexane, chloroform, ethylacetate, n-butanol and remaining water extracts) were tested against various microorganisms including *Herpes simplex* Type-1 (*HSV-1*) and *Parainfluenza viruses* (*PIV*)¹⁰.

In this study, antimicrobial activity results of six flavonoids isolated from three plants used in traditional Turkish medicine against ten extended-spectrum β -lactamase producing multidrug - resistant (trimetoprim - sulphametoxazole, sulbactam-ampicillin, clavulonate-amoxicilin, ceftriaxon, cefepime, imipenem, ceftazidime, tobramicin, gentamicin, ofloxacin, ciprofloxacin) bacteria *K. pneumoniae* (ES β Ls) were presented. *In vitro* broth microdilution testing was performed in accordance with the guidelines of CLSI. Ampicillin and ofloxacin were used as control agents while *K. pneumoniae* RSKK (3) was used as the test microorganism.

EXPERIMENTAL

Test compounds

5,7-dimethoxyflavanone-4'-O- β -D-glucopyranoside (**1**) was isolated from the ethylacetate extract of the leaves and stems of *V. album* ssp. *album* (Loranthaceae) from *Armeniaca vulgaris* Lam. by medium pressure liquid chromatography (MPLC) and 5,7-dimethoxyflavanone-4'-O-[2''-O-(5'''-O-trans-cinnamoyl)- β -D-apiofuranosyl]- β -D-glucopyranoside (**2**) was also isolated from the n-butanol extract of the same plant using several chromatographic methods¹¹.

A novel flavanone glucoside, 5,7,3'-trihydroxyflavanone-4'-O- β -D-glucopyranoside (**3**), was

isolated from the ethanol extract of the herb of *G. fissurense*, in addition to naringenin-7-*O*- β -D-glucopyranoside (**4**)⁷. From the n-butanol extract of *C. hypoleucum* aerial parts (Asteraceae), after multi-stage column chromatographies, two flavonoids, quercetin-3-*O*-rutinoside (rutin) (**5**) and kaempferol-3-*O*-rutinoside (nicotiflorin) (**6**) were previously obtained⁹. The structures of isolated compounds were elucidated by conventional methods of analysis, as well as by different NMR and MS techniques.

Microbiological studies

Preparation of the test materials

The flavonoids were dissolved in dimethylsulphoxide to a concentration of 256 $\mu\text{g ml}^{-1}$ and sterilized by filtration using 0.22 μm Millipore filter (MA 01730, USA) and then used as the stock solutions. Reference antibacterial agents of ampicillin (AMP; Faco), ofloxacin (OFX; Hoechst Marion Roussel) were obtained from their respective manufacturers and dissolved in phosphate buffer solution (ampicillin; pH: 8.0, 0.1 mol/L), and in water (ofloxacin). The stock solutions of these agents were prepared in medium according to as CLSI (formerly National Committee for Clinical Laboratory Standards) recommendations (CLSI; formerly NCCLS)¹².

Microorganisms and Inoculum preparation

Isolated strains of ten *K. pneumoniae* that are resistant to trimetoprim-sulphamethoxazole (SXT; Oxoid; 25 μg ; ≤ 10 mm), sulbactam-ampicillin (SAM; Oxoid; 20 μg ; ≤ 11 mm), clavulonate-amoxicilin (AMC; Oxoid; 20 μg ; ≤ 13 mm), ceftriaxon (CRO; Oxoid; 30 μg ; ≤ 25 mm), cefepime (CPM; Oxoid; 30 μg ; ≤ 14 mm), imipenem (IMP; Oxoid; 10 μg ; ≤ 13 mm), ceftazidime (CAZ; Oxoid; 30 μg ; ≤ 14 mm), tobramycin (TOB; Oxoid; 10 μg ; ≤ 12 mm), gentamicin (GM; Oxoid; 10 μg ; ≤ 12 mm), ofloxacin (OFX; Oxoid 5 μg ; ≤ 12 mm), ciprofloxacin (CIP; Oxoid 5 μg ; ≤ 13 mm) in disc diffusion test were used for the determination of antibacterial activity (minimum inhibition concentration, MIC). *K. pneumoniae* RSKK 574 (Refik Saydam Central Hygiene Institute-Culture Collection,

The Ministry of Health of Republic of Türkiye, Ankara) was used as the control strain¹².

Mueller Hinton Broth (MHB; Oxoid) and Mueller Hinton Agar (MHA; Oxoid) were applied for growing and diluting of the bacteria suspensions. The microorganism suspensions used for inoculation were prepared at 10^7 cfu (colony forming units)/ml by diluting fresh cultures at McFarland 0.5 density (10^8 cfu ml^{-1}). Suspensions of all bacteria were added in each well of the diluted test compounds density of 10^5 cfu ml^{-1} ^{12,13}.

Confirmatory test for ES β L-producing *K. pneumoniae* isolates

The double-disc synergy and agar diffusion tests were used as screening tools to detect ES β L-producing strains. In the double-disc synergy test, the antibiotic discs (Oxoid) used were cefotaxime (30 μg) and ceftazidime (30 μg) placed on Mueller-Hinton agar adjacent to a co-amoxiclav disc (20 μg amoxicillin plus 10 μg clavulanate). The procedures and interpretation of the double-disc synergy test were described previously¹⁴. The agar diffusion test was performed according to NCCLS guidelines¹². A ≤ 5 mm increase in a zone diameter for either ceftazidime/clavulanic acid (30 μg / 10 μg) or cefotaxime/clavulanic acid (30 μg / 10 μg) versus its zone when tested alone was taken as being indicative of ES β L-production¹⁵.

Antimicrobial activity evaluation

The microdilution method was employed for antibacterial tests. Media were placed into each 96 wells of the microplates. Sample solutions at 256 $\mu\text{g ml}^{-1}$ were added into first rows of microplates and two-fold dilutions of the compounds (128-0.0312 $\mu\text{g ml}^{-1}$) were made by dispensing the solutions into the remaining wells. 10 μl culture suspensions were inoculated into all the wells. The sealed microplates were incubated at 35°C for 18h. The lowest concentration of the flavonoids that completely inhibit macroscopic growth was determined and the MICs were recorded¹⁶.

RESULTS

Antimicrobial effects of six flavonoids isolated from three Turkish plants on ten isolated strains of ES β L-containing *K. pneumoniae* are presented in Table 1. In our previous studies, we reported that two new flavonoids, 5,7,3'-trihydroxy-flavanone-4'-O- β -D-glucopyranoside (**1**) and 5,7-dimethoxyflavanone-4'-O-[2''-O-(5'''-O-*trans*-

of all *K. pneumoniae* at 32 and 64 $\mu\text{g ml}^{-1}$ concentrations, which are close to the effective concentrations exhibited by the control agents. Notably, these compounds possessed quite remarkable antimicrobial activities against isolates, *Kp*₃, *Kp*₅, *Kp*₆ and *Kp*₁₀, similar ampicillin and ofloxacin (32 $\mu\text{g ml}^{-1}$). Isolates *Kp*₁, *Kp*₂, *Kp*₄, *Kp*₇₋₉ were inhibited at a concentration of 64 $\mu\text{g ml}^{-1}$ by the tested

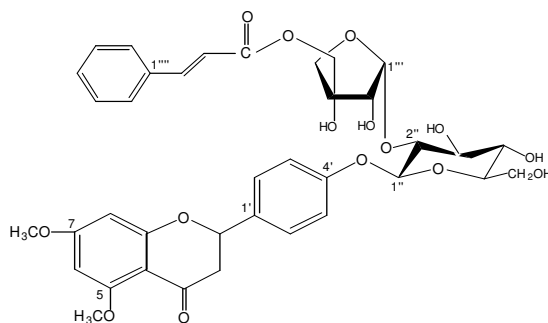
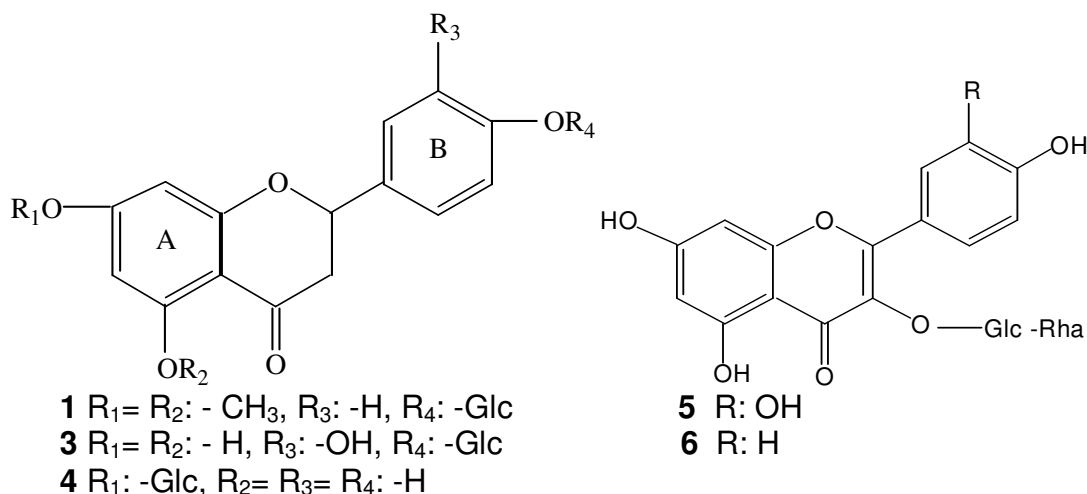


Fig 1: Structures of investigated compounds

cinnamoyl)- β -D-apiofuranosyl]- β -D-glucopyranoside (**2**), were isolated for the first time from *V. album ssp. album* and *G. fissurensis*, respectively^{7,11}. When compared with control agents (ampicillin and ofloxacin), all of the tested flavonoids (**1-6**) showed remarkable activities against isolated strains

flavonoids, the observed activities being twice the dose of ampicillin and ofloxacin (32 $\mu\text{g ml}^{-1}$). On the other hand, the activity at 8 $\mu\text{g ml}^{-1}$ concentration of the flavonoids (**1-6**) against *K. pneumoniae* RSKK 574 seems less active when compared with ampicillin (2 $\mu\text{g ml}^{-1}$) and ofloxacin (<0.12 $\mu\text{g ml}^{-1}$).

Table 1: Antimicrobial activity as MICs ($\mu\text{g ml}^{-1}$) of flavonoids and the references against tested isolated strains of *Klebsiella pneumoniae* ($Kp_{n: 1-10}$)

Compound	Microorganisms										K. <i>pneumoniae</i> RSKK 574
	Kp_1	Kp_2	Kp_3	Kp_4	Kp_5	Kp_6	Kp_7	Kp_8	Kp_9	Kp_{10}	
	R (SXT; AMC; SAM; CRO; CPM; IMP; TOB)	R (SXT; AMC; CRO; IMP; TOB)	R (AMC ; SAM; CRO; CPM)	R (AMC; CRO; CAZ; CPM; GM)	R (AMC ; CRO; CPM; GM)	R (AMC; SAM; CRO; CPM)	R (SXT; AMC; SAM; CRO; CPM; OFX)	R (SXT; AMC; SAM; CRO; CPM; OFX; CIP)	R (SXT; AMC; SAM; CRO; CPM; OFX; CIP)	R (AMC; SAM; CRO; OFX)	
5,7-dimethoxyflavanone-4'-O- β -D-glucopyranoside (1)	64	64	32	64	32	32	64	64	64	32	8
5,7-dimethoxyflavanone-4'-O-[2''-O-(5'''-O-trans-cinnamoyl)- β -D-apiofuranosyl]- β -D-glucopyranoside (2)	64	64	32	64	32	32	64	64	64	32	8
5,7,3'-trihydroxyflavanone-4'-O- β -D-glucopyranoside (3)	64	64	32	64	32	32	64	64	64	32	8
naringenin-7-O- β -D-glucopyranoside (4)	64	64	32	64	32	32	64	64	64	32	8
Quercetin-3-O-rutinoside (rutin) (5)	64	64	32	64	32	32	64	64	64	32	8
Kaempferol-3-O-rutinoside (nicotiflorin) (6)	64	64	32	64	32	32	64	64	64	32	8
Ampicillin	-	-	-	-	-	-	-	-	-	-	2
Ofloxacin	32	32	32	32	32	32	32	32	32	32	<0.12

R: Resistant, SXT: Trimetoprim-Sulphametoxazole, SAM: Sulbactam-Ampicillin, AMC: Clavulonate-Amoxicillin, CRO: Ceftriaxon, CPM: Cefepime, IMP: Imipenem, TOB: Tobramicin, CAZ: Ceftazidime, GM: Gentamicin, CIP: Ciprofloxacin.

DISCUSSION

Plasmid-mediated ES β LS pose a worldwide resistance problem. These isolates are usually resistant to all aminoglycosides, third-

and fourth-generation cephalosporins and monobactams. Such isolates are involved frequently in outbreaks of infection, particularly in high-risk areas, such as

intensive care or neonatal units. Since the discovery of ES β L-producing coliforms in the mid-1980s, over 100 types of different enzymes have been described and have become a worldwide problem. Thus, the members of Enterobacteriaceae producing ES β L are a clinical threat and have been associated with increasing mortality in patients with severe infection. ES β Ls are mutant enzymes, which derived from TEM or SHV (class A) enzymes. They confer variable levels of resistance to cefotaxime, ceftazidime and other broad-spectrum cephalosporins and to monobactams such as aztreonam. Thus, the choice of effective and safe drugs to be used against ES β L is shrinking rapidly. Therefore, in recent years, researches are increasingly turning their attention to folk medicine for new leads to develop better drugs against clinical ES β L-producing *K. pneumoniae* isolates^{2,17-19}

In this study, structure-activity relationships do not appear to play a role here as all six flavonoids exhibited similar MICs (32–64 $\mu\text{g ml}^{-1}$) against all of the *K. pneumoniae* strains. We observed that active flavonoids **3–6** have an obligatory C–4 keto group and hydroxyl group substitutions at C–5 and have at least one hydroxyl group on ring B. These results are in consonance with the findings of Xu and Lee¹⁹. On the other hand, although the other active flavonoids **1** and **2** have methoxyl group substitutions at C-5 and C-7, these compounds also exhibited antimicrobial activity against *K. pneumoniae* at similar MICs values to those of ampicillin and ofloxacin. Alcaraz *et al* and Xu and Lee demonstrated that substitution with methoxyl groups drastically decreased the antibacterial of flavonoids^{20,21}. Additionally, it is reported that rutin (**5**) from *Linum capitatum* Kit. flowers inhibited the growth of *P. aeruginosa*, but did not show any activity against *E. coli* different from other Gram-negative bacteria²¹. According to the other report, the same compound tested by the disc-diffusion method showed no activity against *K. pneumoniae* ATCC 13883²⁰. These inconsistencies may be due to variations within each assay. Furthermore, it was not stated whether the

test flavonoids were obtained from a commercial or natural source.

To the best of our knowledge, this is the first report on the inhibitory activity of 5,7-dimethoxyflavanone-4'-O- β -D-glucopyranoside (**1**), 5,7-dimethoxyflavanone-4'-O-[2''-O-(5'''-O-*trans*-cinnamoyl)- β -D-apiofuranosyl]- β -D-glucopyranoside(**2**), 5,7,3'-trihydroxyflavanone-4'-O- β -D-glucopyranoside (**3**), naringenin-7-O- β -D-glucopyranoside (**4**), and nicotiflorin (**6**) against ES β L-producing *K. pneumoniae* isolates.

CONCLUSION

All the flavonoids (**1-6**) showed *in vitro* antimicrobial activity against all the isolated strains of *K. pneumoniae*, similar to that produced by the control antibacterial (ofloxacin) at the concentration of 32-64 $\mu\text{g ml}^{-1}$; on the other hand, another control, ampicillin, had no activity. To the best of our knowledge, this is the first report of the evaluation of the activity of these flavonoids against (ES β L)-producing *K. pneumoniae*. On the basis of these data presented, these flavonoids may be considered potential therapeutic compounds for infections that may be caused by ES β L-producing bacteria in the future. Therefore, further work is under way to identify their precise antibacterial mechanism. Additionally, the antimicrobial activity of some antibacterials in combination with these flavonoids against extended-spectrum β -Lactamase (ES β L)-producing *K. pneumoniae* may also need to be evaluated for the treatment of infections caused by ES β L-producing bacteria.

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