**Full Length Research Paper**

**Antibacterial effect of Jordanian propolis and isolated flavonoids against human pathogenic bacteria**

Rula M. Darwish¹*, Ra’ed J. Abu Fares², Musa H. Abu Zarga³ and Ibrahim K. Nazer²

¹Pharmaceutics and Pharmaceutical Technology Department, Faculty of Pharmacy, University of Jordan, Amman, Jordan.
²Plant Protection Department, Faculty of Agriculture, University of Jordan, Amman, Jordan.
³Chemistry Department, Faculty of Science, University of Jordan, Amman, Jordan.

Accepted 11 August, 2010

Propolis is a natural product widely consumed in folk medicine. The present study was carried out to investigate the antibacterial activity of Jordanian propolis, collected from two locations with two different dominant floras (Type I: Pine trees and Type II: Oak trees). Zones of inhibition and minimum inhibitory concentrations (MICs) were determined on methicillin resistant *Staphylococcus aureus* (MRSA), multidrug resistant *Escherichia coli* and standard strains of both bacteria. Propolis Type I and Type II showed antibacterial activity against MRSA (MIC 4.69 and 18.75 µg ml⁻¹, respectively). Crude propolis from Type I showed higher antibacterial activity than Type II against the tested bacteria. Three pure phenolic compounds (three flavonoids) namely, pinobanksin-3-O-acetate, pinocembrin and chrysin, were isolated from fractions I-2 and I-4, and screened *in vitro* for antibacterial activity. Pinobanksin-3-O-acetate and pinocembrin exhibited antibacterial activity especially against MRSA, while chrysin was only active against standard *S. aureus*. This is the first report that shows *in vitro* antibacterial activity of isolated flavonoids from Jordanian propolis against standard and resistant strains of *E. coli* and MRSA. Overall, results of this study highlight the important role of propolis botanical source on the antibacterial activity of such natural material which might affect its medical applications.

**Key words:** Antibacterial activity, human pathogens, flavonoids, propolis, methicillin resistant *Staphylococcus aureus*, *Escherichia coli*.

**INTRODUCTION**

Propolis is a natural product widely consumed in folk medicine. Its medicinal and antimicrobial properties have been widely investigated (Kujumgiev et al., 1999; Hegazi and Abd El Hady, 2002; Prytyzk et al., 2003; Onlen et al., 2007; Darwish et al., 2009). Different researchers (Sforcin et al., 2000; Trusheva et al., 2006; Katicio and Nazime 2006; Yaghoubi. et al., 2007) have reported that propolis antibacterial activity is attributed to a number of phenolic compounds, mainly flavonoids, phenolic acids and their esters. Some prenylated ρ-coumaric acids were isolated from propolis in several countries (Kosalec et al., 2004). The antibacterial activity of volatile compounds and diterpenes from Brazilian propolis was identified by Bankova et al. (2000). Propolis and some of its cinnamic acid derivatives and flavonoids were responsible for uncoupling the energy transducing cytoplasmic membrane inhibiting bacterial motility, which might contribute to the antibacterial action (Bankova et al., 2000).

It is reported that pathogenic isolates have a relatively large potential for developing antibiotic resistance (Fluit et al., 2000; Sahm et al., 2001; Schwaber et al., 2006). The increase in antibiotic resistant bacteria is largely due to the widespread use of antibiotics in medicine, in animal care and in agriculture. The problem is compounded by the lack of new antibiotics to attack bacteria in different

---

Abbreviations: MICs, Minimum inhibitory concentrations; MRSA, methicillin resistant *Staphylococcus aureus*; TLC, thin layer chromatography; MDR, multi drug resistance; CC, column chromatography.
ways to circumvent the resistant genes. Therefore, finding antimicrobial agents which are effective or might enhance the antibiotic efficacy against resistant bacteria would be an advantage. *Staphylococcus aureus* and *Escherichia coli* are of the major causes of hospital-acquired infections (Karlowsky et al., 2004; Fluit et al., 2000; Huang et al., 2006). These organisms occur naturally in and on human body. However, certain strains can lead to infections and are becoming resistant to antibiotics.

Numerous researchers have reported the biological activities of propolis collected in different parts of the world. However, no work has been conducted on the antibacterial activity of Jordanian propolis or its components. Thus, the objectives of this research were to investigate the antibacterial activity of the crude propolis, separated propolis fractions and promising constituents from two different floras (pine and oak trees) against resistant and standard strains of two medically important bacteria *S. aureus* and *E. coli*.

### MATERIALS AND METHODS

**Extraction of propolis**

Two Jordanian propolis samples were collected by honey bees (*Apis mellifera*) from two locations with two different dominant floras: University of Jordan main campus, located in Amman in which common pine trees with the majority of *Pinus halepenses* were the source of Type I propolis and Al-Hashmeah, located about 12 km west of Amman and is characterized by intensive oak trees (*Quercus coccifera*) which were the source of Type II propolis. Collected propolis was stored at -20°C. A specimen was retained and extracted once with chloroform for one day and twice with methanol for two days each. The extracts were evaporated under reduced pressure. The residual material was partitioned between hexane and 10% aqueous methanol. The crude aqueous methanol extract of Type I and Type II propolis were chromatographed on a silica gel column as described for fraction I-2 and I-4 was tested by the NCCLS broth microdilution method as described by the National Committee for Clinical Laboratory Standards (NCCLS, 2000a). The isolates were grown overnight on a nonselective medium at 35°C. Bacterial inocula were prepared by suspending the freshly grown bacteria in sterile normal saline and adjusted to a 0.5 McFarland standard. The suspension was inoculated onto 9 cm plates of Miller-Hinton agar (Merck). Cups of 9 mm were filled with saturated solutions of the crude propolis or the fractions. The plates were incubated for 24 h at 37°C. Control cups containing the solvent alone were also included. The absence of growth was interpreted as antibacterial activity. The antibiotics clindamycin (2 units/disc), penicillin G (10 units/disc), cephalixin (30 µg/disc) and nalidixic acid (30 µg/disc) were used as standards to compare the activity of propolis in inhibiting the growth of bacteria. The antibacterial activity was determined by comparing zones of inhibition (mm) and complete inhibition (including the diameter of the disc) produced by the propolis or its fractions with those in the controls and standards. Each experiment was carried out three times and was correlated against the controls.

**Identification of the isolated compounds**

Mass spectra for the flavones were obtained using a Varian MAT 112 spectrometer. 1H-NMR spectra were recorded on a Bruker DPX-300 MHz spectrometer with TMS as internal standard. 13C-NMR spectra were recorded at 75.5 MHz. Silica gel DF (Fluka) was used for CC and silica gel glass plates G-UV254 (Macherey-Nagel) were used for TLC. All chemical procedures were carried out at the Chemistry Department, University of Jordan. Identified structures of the isolated compounds are presented in Figure 4.

**Antimicrobial assay**

Two resistant bacterial strains, methicillin resistant *S. aureus* (MRSA) and resistant *E. coli* were isolated from hospitalized patients at the Jordan University Hospital and approved by biochemical tests. Two standard bacterial strains *S. aureus* ATCC 6538P and *E. coli* ATCC 8739 were also used in the study. The susceptibilities of the isolates to the crude propolis extracts and the fractions were determined by the agar diffusion method on solid media with Mueller-Hinton agar plates (Merck, Germany) according to the recommendations of the National Committee for Clinical Laboratory Standards (NCCLS, 2000a). The isolates were grown overnight on a nonselective medium at 35°C. Bacterial inocula were prepared by suspending the freshly grown bacteria in sterile normal saline and adjusted to a 0.5 McFarland standard. The suspension was inoculated onto 9 cm plates of Miller-Hinton agar (Merck). Cups of 9 mm were filled with saturated solutions of the crude propolis or the fractions. The plates were incubated for 24 h at 37°C. Control cups containing the solvent alone were also included. The absence of growth was interpreted as antibacterial activity. The antibiotics clindamycin (2 units/disc), penicillin G (10 units/disc), cephalixin (30 µg/disc) and nalidixic acid (30 µg/disc) were used as standards to compare the activity of propolis in inhibiting the growth of bacteria. The antibacterial activity was determined by comparing zones of inhibition (mm) and complete inhibition (including the diameter of the disc) produced by the propolis or its fractions with those in the controls and standards. Each experiment was carried out three times and was correlated against the controls.

**Determination of the minimum inhibitory concentration (MIC)**

The MICs of the propolis and the fractions were conventionally determined in triplicate for each strain by the macrodilution broth method as described by the NCCLS (National Committee for Clinical Laboratory Standards, 2000b). Serial dilutions of each propolis and fractions were prepared in macrodilution tubes. Bacterial suspensions were adjusted to the 0.5 McFarland standards (approximately 1 to 2 × 10^8 cfu/ml). Final inocula were adjusted to the 10^4 cfu/ml. A constant amount of bacteria were added to all tubes and they were incubated at 37°C for 18 – 24 h. Each tube was examined for growth, comparing each tube to the control. The MIC was defined as the lowest concentration of propolis at which there was no visible growth of the organism. MICs of the antibiotics were determined in the same way. A positive growth control was included where bacterial suspension was added to a tube filled with nutrient broth without crude propolis. An uninoculated tube of nutrient broth was also added to serve as negative growth control.

**Broth microdilution method for determining MIC of isolated flavonoids**

The susceptibility of each bacterial isolate to the three compounds isolated from fraction I-2 and I-4 was tested by the NCCLS broth microdilution reference method (NCCLS, 2000b) with some modifications. MIC test was performed in 96 flat bottom microtiter plates (TPP, Switzerland). Each test well was filled with a volume of 100 µl nutrient broth. Extracts and fractions under investigation (100 µl) were added to the first column of test-well and mixed. A series of
randomized design (CRD) arrangement and analyzed according to laboratory conditions using a split plot design with complete treatments of gram positive and negative bacteria were carried out. Statistical analysis was performed using the Student’s t-test. Plates 1a, 1b, 2a and b. Type I crude propolis and fraction I-4 produced the highest antibacterial activity with zones of inhibition of 24.67 and 25.33 mm, respectively on standard S. aureus (Figure 1 and Plate 1a) and 17.00 and 16.67 mm, respectively on MRSA (Figure 2 and Plate 1b). The activity of fractions I-4 indicates that the active components are concentrated in this fraction. This is in agreement with reports of several researchers which indicate that each propolis sample contain 80 - 100 chemical compounds with different concentrations (Bankova et al., 2000; Kosalec et al., 2004; Trushera et al.; 2004; Park et al., 2005; Yaghoubi et al., 2007).

Crude propolis Type II against standard S. aureus did not differ significantly from fraction II-4 and was less effective compared to Type I crude propolis. Type II and fraction II-4 exerted zone of inhibition of 18.67 and 18.00 mm, respectively (Figure 1 and Plate 2a). Their effect on MRSA was less, with inhibition zones of 14.67 and 14.33 mm, respectively (Figure 2 and Plate 2b).

The antibacterial activity of antibiotics used (penicillin G 2 units/disc, cephalexin 10 units/disc, clindamycin 30 µg/disc and nalidixic acid 30 µg /disc) was comparable with that of crude propolis Type I with zones of inhibition of 23, 23, 24 and 18 mm, respectively against standard S. aureus (Table 1). Fractions I-2 and I-4 of Type I gave inhibition zones of 22 and 25.33 mm against standard S. aureus which are comparable with that obtained by the antibiotics used (Figure 1 and Table 1) indicating that the inhibitory effect of the propolis is mainly attributed to these fractions. It is worth mentioning that the antibiotics used have no effect on MRSA (Table 1), while crude propolis Type I, Type II and their fractions inhibited the growth of this strain. Crude propolis Type I, Type II, fractions I-2 and I-4 and fractions II-2 and II-4 gave inhibition zones of 17.00, 14.67, 13.67, 18, 11.67 and 14.33 mm, respectively, against MRSA (Figure 2, Plates 1b and 2b).

The present results on standard S. aureus are in agreement with those obtained by several authors like Kujumgiev et al. (1999) who found that the inhibition zones obtained by propolis from Mongolia, Albania, Egypt, Brazil and the antibiotic streptomycin were 24, 21.8, 24.3, 21.8 and 28 mm, respectively. These results are also comparable with results obtained by Prytz et al. (2003) who measured the inhibition zone for Bulgarian propolis as 20 mm. They are also comparable with results obtained by Stepanovic et al. (2003) who found that the inhibition zone of propolis form different geographical areas of Serbia ranged from 18 - 23 mm.

Results reveal that there is antibacterial activity of

<table>
<thead>
<tr>
<th>Treatment</th>
<th>Zone of inhibition (mm) ±SD</th>
<th>Standard</th>
<th>Resistant</th>
</tr>
</thead>
<tbody>
<tr>
<td>S. aureus</td>
<td>E. coli</td>
<td>MRSA</td>
<td>E.coli</td>
</tr>
<tr>
<td>Penicillin G</td>
<td>23 ± 2.5</td>
<td>23 ± 1.92</td>
<td>N</td>
</tr>
<tr>
<td>Cephalexin</td>
<td>23 ± 1.00</td>
<td>22 ± 1.51</td>
<td>N</td>
</tr>
<tr>
<td>Clindamycin</td>
<td>24 ± 0.67</td>
<td>24 ± 1.75</td>
<td>N</td>
</tr>
<tr>
<td>Nalidixic acid</td>
<td>18 ± 1.33</td>
<td>22 ± 2.00</td>
<td>N</td>
</tr>
<tr>
<td>Methanol</td>
<td>N</td>
<td>N</td>
<td>N</td>
</tr>
</tbody>
</table>

*Average of three replicates; N: no zone of inhibition.

**Table 1. Means zone of inhibition by four antibacterial agents on standard and resistant (MRSA) S. aureus and E. coli.**

- Statistical analysis
- Treatments of gram positive and negative bacteria were carried out under laboratory conditions using a split plot design with complete randomized design (CRD) arrangement and analyzed according to LSD test SAS (1990). Three replicates of Petri dishes were used for each treatment, mean zone of inhibition and standard deviations were calculated and presented in Table 1.

**RESULTS AND DISCUSSION**

The antibacterial activity of crude propolis and different fractions from Type I and II against standard S. aureus and MRSA was tested as presented in Figures 1 and 2 and Plates 1a, 1b, 2a and b. Type I crude propolis and fraction I-4 produced the highest antibacterial activity with zones of inhibition of 24.67 and 25.33 mm, respectively on standard S. aureus (Figure 1 and Plate 1a) and 17.00 and 16.67 mm, respectively on MRSA (Figure 2 and Plate 1b). The activity of fractions I-4 indicates that the active components are concentrated in this fraction. This is in agreement with reports of several researchers which
Figure 1. Mean zone of inhibition by various propolis treatments against standard *S. aureus* ATCC 6538P.

Figure 2. Mean zone of inhibition by various propolis treatments against resistant *S. aureus* MRSA.
propolis Types I and II against the different studied bacteria (including gram positive and negative bacteria) and that Type I propolis has better antibacterial activity than Type II. This indicates that propolis from the different


regions in Jordan exhibits antibacterial effect. This again supports the commonly reported statement in literature, that sensitivity of microbes and differences in chemical composition of propolis are greatly affected by variations in geographical origins (Bankova et al., 2000; Abd El Hady and Hegazi, 2002; Kartal et al., 2003; Trusheva et al., 2006).

Results in Table 2 emphasize the effect of different flora on the antibacterial activity. MICs of Type I and II propolis against standard *S. aureus* were 0.585 and 4.69 mg ml\(^{-1}\), respectively. The MIC of Type I propolis is similar to what was reported by Sforcin et al. (2000) on propolis collected from Brazil. Our MIC results from pine propolis (Type I) on standard *S. aureus* (0.585 mg ml\(^{-1}\)) are better than those reported by Hegazi and Abd El Hady (2002) in Egypt which were 2.2 and 2.6 mg ml\(^{-1}\), respectively. However, Moreno et al. (1999) reported that propolis collected from Argentina had lower MIC value of 0.04 mg ml\(^{-1}\) against the same strain. Again, this difference was related to the different constituents of propolis collected from different geographical regions (Bankova et al., 2000; Abd El Hady and Hegazi, 2002). It is worth adding that Jordanian crude propolis Type I and II were also active against MRSA with 4.69 and 18.75 mg ml\(^{-1}\), respectively (Table 2). This bioactivity on resistant bacterial strains opens the door for practical application.

Several researchers (Kujumgiev et al., 1999; Moreno et al., 1999; Sforcin et al., 2000; Stepanović et al., 2003; Gonzalez et al., 2005) reported that there was no effect of propolis from different geographical regions on standard *E. coli*. Our results however, show that there is some antibacterial effect of propolis on gram negative bacteria, but it is rather limited with a zone of inhibition of 17.33 and 15.67mm for crude propolis Type I and fraction I-4 treatments, respectively (Figure 3). Meanwhile, propolis Type II has no effect. This again might reflect the fact that chemical composition of propolis differs greatly from one region to another (Burdoc, 1998; Bankova et al., 2000; Abd El Hady and Hegazi, 2002; Prytyzk et al., 2003; Stepanovic et al., 2003).

The MIC value against standard *E. coli* was 4.69 mg ml\(^{-1}\) for Type I propolis, while Type II propolis showed no activity (Table 2). This MIC value is better than that reported by Sforcin et al. (2000) of 8 mg ml\(^{-1}\) on the same strain. However, it was less than the MIC obtained by Hegazi and Abd El Hady (2002) who reported an MIC value of 1.6 mg ml\(^{-1}\) against the same bacteria. The variation might reflect the difference in the composition of the propolis, since the bacterial strain used is the same. The low sensitivity of *E. coli* shown in Table 2 and Figure 3 is in agreement with the findings by many researchers where this bacterium showed either very low sensitivity or total lack of sensitivity against propolis (Marcucci, 1995; Kujumgiev et al., 1999; Gonzalez et al., 2005). This emphasizes the fact that, gram negative bacteria are less sensitive than gram positive strains, which is in agreement with several previous reports (Burdoc, 1998; Moreno et al., 1999; Sforcin et al., 2000; Abd El Hady and Hegazi, 2002; Gonzalez et al., 2005). The most possible explanation for the low sensitivity of gram negative bacteria is that their outer membrane inhibits and/or retards the penetration of propolis (Tegos et al., 2002). Another possible reason is their possession of multi drug resistance (MDR) pumps, which extrude amphipathic toxins across the outer membrane (Tegos et al., 2002).

Results in Table 2 show an MIC value of 18.75 mg ml\(^{-1}\) for Type I propolis against resistant *E. coli*, while Type II propolis showed no activity on this strain. The results showed that crude propolis of both Types I and II and their fractions had no (or slight) antibacterial activity on resistant *E. coli*. However, they had activity on resistant *S. aureus* (MRSA). It is worth mentioning that the antibiotics used had no effect on the same strains (Table 2).

Overall results indicate that the antibacterial activity of the crude propolis, fraction I-2 and I-4 are superior to other fractions, indicating that the bioactive components are concentrated in these fractions. Our preliminary studies indicate that the compounds in these fractions are flavonoids. This is in agreement with the findings in the literature, which indicate that the active compounds present in propolis include mainly flavonoids, phenolic acids and esters (Marcucci, 1995; Burdoc, 1998). Major compounds from the active fractions (I-2 and I-4) were isolated by a combination of silica gel column chromatography (CC) and TLC. Fraction I-2 afforded the flavans 1 (Jung et al., 1990) and 2 (Fang et al., 1988) while fraction 4 afforded flavan 3 (Chen et al., 2003) as the major constituents (Figure 4). The three compounds were identified by spectroscopic methods including \(^1\)H-NMR,

### Table 2. Minimal inhibitory concentration (MIC)* of crude propolis against standard and resistant strain of *S. aureus* and *E. coli*.

<table>
<thead>
<tr>
<th>Microorganism</th>
<th>Crude propolis (mg ml(^{-1}))</th>
<th>Antibacterial (µg ml(^{-1}))</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>Type I</td>
<td>Type II</td>
</tr>
<tr>
<td><em>Staphylococcus aureus</em> ATCC 6538P</td>
<td>0.585</td>
<td>4.69</td>
</tr>
<tr>
<td><em>Escherichia coli</em> ATCC 8739</td>
<td>4.69</td>
<td>ND</td>
</tr>
<tr>
<td>MRSA</td>
<td>4.69</td>
<td>18.75</td>
</tr>
<tr>
<td>Resistant <em>Escherichia coli</em></td>
<td>18.75</td>
<td>ND</td>
</tr>
</tbody>
</table>

*Average of three replicates; ND: not detected.
Figure 3. Mean zone of inhibition by crude and seven pine propolis fractions Type I against standard *E. coli*.

Figure 4. The chemical structure of the isolated compounds.
Table 3. Mean zone of inhibition by pure compounds isolated from fractions I-2 and I-4 against standard and resistant strains of *S. aureus* and *E. coli*.

<table>
<thead>
<tr>
<th>Bacteria</th>
<th>Zone of inhibition( mm)</th>
<th>Blank (solvent)</th>
<th>Pinobanksin-3-O-acetate</th>
<th>Pinocembrin</th>
<th>Chrysin</th>
</tr>
</thead>
<tbody>
<tr>
<td>Standard <em>S. aureus</em></td>
<td>N</td>
<td>13</td>
<td>13</td>
<td>10</td>
<td></td>
</tr>
<tr>
<td>MRSA</td>
<td>N</td>
<td>22</td>
<td>21</td>
<td>N</td>
<td></td>
</tr>
<tr>
<td>Standard <em>E. coli</em></td>
<td>N</td>
<td>10</td>
<td>N</td>
<td>N</td>
<td></td>
</tr>
<tr>
<td>Resistant <em>E. coli</em></td>
<td>N</td>
<td>12</td>
<td>N</td>
<td>N</td>
<td></td>
</tr>
</tbody>
</table>

Diameter of filter paper disc is 6 mm; *Average of three replicates; N: no zone of inhibition.

Table 4. Minimal inhibitory concentration (MIC) of pure compounds isolated from fractions I-2 and I-4 against standard and resistant strains of *Staphylococcus aureus* and *Escherichia coli*.

<table>
<thead>
<tr>
<th>Bacteria</th>
<th>MIC (µg ml(^{-1}))</th>
<th>Pinobanksin-3-O-acetate</th>
<th>Pinocembrin</th>
<th>Chrysin</th>
</tr>
</thead>
<tbody>
<tr>
<td>Standard <em>S. aureus</em></td>
<td>250</td>
<td>300</td>
<td>250</td>
<td></td>
</tr>
<tr>
<td>MRSA</td>
<td>250</td>
<td>ND</td>
<td>ND</td>
<td></td>
</tr>
<tr>
<td>Standard <em>E. coli</em></td>
<td>500</td>
<td>ND</td>
<td>ND</td>
<td></td>
</tr>
<tr>
<td>Resistant <em>E. coli</em></td>
<td>&gt;500</td>
<td>ND</td>
<td>ND</td>
<td></td>
</tr>
</tbody>
</table>

*Average of three replicates; ND: not detected.

\(^{13}\)C-NMR (with DEPT) and MS.

Screening of the antibacterial activity of the pure compounds isolated from fractions I-2 and I-4 shown in Table 3 reveal that, pinobanksin-3-O-acetate and pinocembrin which were isolated from fraction I-2 have good activity against both strains of *S. aureus*. Their effect was more promising against the resistant strain with inhibition zones of about 22 mm compared to 13 mm against standard strain (Table 3). Chrysin which was isolated from fraction I-4 inhibited the growth of standard *S. aureus* only (zone of inhibition was 10 mm). The inhibition of growth of standard and resistant *E. coli* was noticed only with pinobanksin-3-O-acetate with an inhibition zone of 10 and 12 mm, respectively (Table 3). The promising antibacterial effect of the isolated compounds may be due to the fact that ring B in all of the three flavonoids is not oxygenated; such a structural feature is uncommon in flavonoids. MICs of pinobanksin-3-O-acetate, pinocembrin and chrysin against standard *S. aureus* were similar (about 250 µg ml\(^{-1}\)) (Table 4). These MICs were high when compared with antibiotics MICs. Clindamycin, penicillin G, cephalaxin and nalidixic acid MICs were 7.02, 4, 32 and 8 µg ml\(^{-1}\), respectively (Table 2). This is in agreement with the results obtained by Koo et al. (2002) where the MICs for flavonoids isolated from propolis were greater than 500 µg ml\(^{-1}\) (Table 2). This is due to the fact that ring B in all of the three flavonoids is not oxygenated; such a structural feature is uncommon in flavonoids. MICs of pinobanksin-3-O-acetate, pinocembrin and chrysin against standard *S. aureus* were similar (about 250 µg ml\(^{-1}\)) (Table 4). These MICs were high when compared with antibiotics MICs. Clindamycin, penicillin G, cephalaxin and nalidixic acid MICs were 7.02, 4, 32 and 8 µg ml\(^{-1}\), respectively (Table 2). This is in agreement with the results obtained by Koo et al. (2002) where the MICs for flavonoids isolated from propolis were greater than 500 µg ml\(^{-1}\) (Table 2). These MICs were high when compared with antibiotics MICs. Clindamycin, penicillin G, cephalaxin and nalidixic acid MICs were 7.02, 4, 32 and 8 µg ml\(^{-1}\), respectively (Table 2). This is in agreement with the results obtained by Koo et al. (2002) where the MICs for flavonoids isolated from propolis were greater than 500 µg ml\(^{-1}\) (Table 2). These MICs were high when compared with antibiotics MICs. Clindamycin, penicillin G, cephalaxin and nalidixic acid MICs were 7.02, 4, 32 and 8 µg ml\(^{-1}\), respectively (Table 2). This is in agreement with the results obtained by Koo et al. (2002) where the MICs for flavonoids isolated from propolis were greater than 500 µg ml\(^{-1}\) (Table 2). These MICs were high when compared with antibiotics MICs. Clindamycin, penicillin G, cephalaxin and nalidixic acid MICs were 7.02, 4, 32 and 8 µg ml\(^{-1}\), respectively (Table 2). This is in agreement with the results obtained by Koo et al. (2002) where the MICs for flavonoids isolated from propolis were greater than 500 µg ml\(^{-1}\) (Table 2). These MICs were high when compared with antibiotics MICs. Clindamycin, penicillin G, cephalaxin and nalidixic acid MICs were 7.02, 4, 32 and 8 µg ml\(^{-1}\), respectively (Table 2). This is in agreement with the results obtained by Koo et al. (2002) where the MICs for flavonoids isolated from propolis were greater than 500 µg ml\(^{-1}\) (Table 2). These MICs were high when compared with antibiotics MICs. Clindamycin, penicillin G, cephalaxin and nalidixic acid MICs were 7.02, 4, 32 and 8 µg ml\(^{-1}\), respectively (Table 2). This is in agreement with the results obtained by Koo et al. (2002) where the MICs for flavonoids isolated from propolis were greater than 500 µg ml\(^{-1}\) (Table 2).

The antibacterial activity of the pure compounds was less than expected suggesting that the antibacterial activity of propolis could involve a synergistic interaction between its components. This is in agreement with the study by Hegazi and Abd El Hady (2002) and would explain the effect of crude propolis on bacteria (Kujumgiev et al., 1999).

In conclusion, propolis samples could be used for the standard and resistant strain-infection treatment such as an alternative therapy. The resistant strains are increasing rapidly all over the world; these urge further studies to be done with other microorganisms.

ACKNOWLEDGEMENTS

The authors are grateful to the Deanship of Scientific Research, University of Jordan, and the Higher Council for Science and Technology, Amman, Jordan for the financial support of this work. Sincere thanks are extended to Al-Hikma Pharmaceuticals for their partnership and supply of some of the bacteria and antibiotics used in this work. Our gratitude is extended to Prof. Asem Shehabi from Jordan University Hospital for supplying the resistant strains of bacteria used in this study and to Dr. Mayadah Shehadeh and Bassam Alheyari for their help in writing the manuscript and arranging the graphs.

REFERENCES

Abd El Hady FK, Hegazi AG (2002). Egyptian Propolis: Chemical


