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Abstract

Background and objectives: Microbial resistance to antibiotics is one of the most common problems in the health care system. Therefore, many efforts have been performed to find new compounds as antimicrobial compounds. This study carried out to investigate the in-vitro antibacterial effect of methanolic extract of peppermint on standard Staphylococcus aureus, Bacillus cereus, Escherichia coli and Pseudomonas aeruginosa strains.

Methods: In this experimental laboratory study, after collecting and performing pharmacognosy evaluations, methanolic extract of the peppermint plant was prepared and its antimicrobial effects on several bacteria were determined at concentrations of 20 to 400 mg/ml using the agar well diffusion method, minimum inhibitory concentration (MIC), minimum bactericidal concentration (MBC) methods. The test was repeated five times for each bacterium and the collected data were analyzed using SPSS software.

Results: The methanolic extract of peppermint had antimicrobial effects against standard strains of Staphylococcus aureus, Bacillus cereus, Escherichia coli and Pseudomonas aeruginosa. The highest effect was observed in S. aureus and the least in P. aeruginosa. MIC and MBC of methanol extract on S. aureus, B. cereus, E. coli, and P. aeruginosa were 6.5-15.5, 15.5-25, 50-100, and 100-200 mg/ml, respectively.

Conclusion: The results of this study showed that peppermint extract can be considered as an antibacterial medicinal herb and that if the concentration of the extract increases, its antibacterial properties will also increase. Thus, it can be used as an alternative to the usual chemical drugs in the treatment of infections after evaluating their effects in vitro.

Keywords: Antibacterial effect, Methanolic extract, Peppermint, In vitro.
Introduction

The resistance of microorganisms to antibiotics raises problems that have become one of the major concerns of human society and the health care system (1). With the emergence of antibiotic-resistant, the efficacy of existing drugs has decreased and this has increasingly enhanced the failure of antimicrobial therapies (2). Besides, the widespread use of industrial-origin drugs as well as improper use of these drugs can cause many side effects, sometimes resulting in more serious toxic effects than the disease itself (3). One way to prevent the emergence of antibiotic resistance is to use compounds that work similar to antimicrobial drugs but differ from existing industrial sources (4,5). In fact, due to the increasing incidence of antibiotic resistance, there is a pressing need for new antibacterial drugs, and of the potential sources, such as plants (6). The use of medicinal herbs has been one of the earliest human achievements to treat most diseases in different countries, because the plants make compounds that are related to the antimicrobial properties, including alkaloids, flavonoids, isoflavonoids, tannins, glycosides, and phenolic compounds. In other words, secondary metabolites in plants have antimicrobial properties (7). Mentha includes more than 25 species, but the most common species used is Mentha piperita Lamiaceae (Peppermint). It is one of the most popular aromatic plants, which have traditionally been used in folk medicine. Peppermint is a combination of blue mint and ornamental or spicy mint (8). It has dark green and fragrant leaves and grows 60 to 90 cm. Peppermint is a perennial herbaceous plant whose leaves, are elliptic, transverse, serrate, slightly lanceolate, 4–7cm long and 2–3 cm wide. Peppermint has high menthol content and its oil also contains menthone and carboxyl esters. For this reason, it seems that it can have antimicrobial properties (9). Today, in different countries of the world, more than one thousand tons of oils are produced from this plant, which shows its importance in different parts of the world. Much research has been done on the medicinal properties of peppermint (10). Hence, the objective of this study was to assess the antimicrobial properties of Mentha piperita Lamiaceae on standard bacteria, including *S. aureus*, *B. cereus*, *E. coli* and *P. aeruginosa* in vitro.

Materials and Methods

Preparation of the peppermint plant

Plant samples were collected from natural areas around Tabriz city, Iran. After collection and transfer, the samples were cleaned and dried in a large, well-ventilated area away from sunlight. First, the samples were completely dried, aerial organs such as stems were wounded, and leaves attached to roots were removed, then they were prepared for grinding.

Extraction of methanol extract

Soxhlet method was used for extraction so that 60 g of dried plant powder with 300 ml of methanol as solvent was placed in Soxhlet extractor for 8 hours. The solvent evaporated slowly and was concentrated by using a rotary apparatus at 40° C. Next, 5% Dimethyl sulfoxide (DMSO) was used to prepare the extracts. The concentrations of the extracts were 20, 30, 50, and 400 mg/ml for using in Minimum Inhibitory Concentration (MIC) and Disc diffusion experiments.
Preparing the standard microorganisms

The microorganisms studied in this study were: *Staphylococcus aureus* ATCC 25923, *Bacillus cereus* ATCC 1247, *Escherichia coli* ATCC 25922, and *Pseudomonas aeruginosa* ATCC 27853. All strains were prepared from the microbial collection at the University of Tehran. The culture was performed on the Muller Hinton agar medium (Merck, Germany) to obtaining colonies, then, to prepare microbial suspension from fresh culture, 4-5 colonies were transferred to Mueller Hinton Broth medium (Merck, Germany) and microbial turbidity prepared according to McFarland standard tube No. 0.5 (1.5×10^6 CFU/ml). To reach the bacterial concentration of 1.5×10^6 CFU/ml, microbial suspension diluted to 0.5 McFarland tube equivalents to 0.01 and next, it was checked by spectrophotometer.

Investigation of the antimicrobial effect of methanol extract

To investigate the antimicrobial effect of peppermint methanol extract, concentrations of 20, 30, 50, and 400 mg/ml of methanol extract were prepared in 5% DMSO solvent. In this study, the agar well diffusion and dilution test were used to investigate the antimicrobial effect of methanol extract.

Determination of the antimicrobial effect by agar well diffusion

In the agar well diffusion method, 500 µl of the microbial suspension was transferred to the Muller Hinton agar medium and it was cultured by sterile swabs in three directions. Then, wells were made at 6 mm in diameter and 2.5 cm apart on the culture medium surface (the bottom of the wells filled with the medium again). Finally, 100µl of 20, 30, 50, and 400 mg/ml concentrations of the methanol extract were loaded into each well. The 5% DMSO was used as negative control and chloramphenicol was used as a positive control. Plates were then incubated at 37 °C for 24h and microbial cultures were measured for the presence or absence of growth zone in millimeters.

Determination of MIC and MBC

The minimum inhibitory concentration (MIC) and minimum bactericidal concentration (MBC) of methanol extract were determined by dilution test. In this method, for the determination of MIC, the methanol extract was prepared in serial dilutions of 6.25, 12.5, 25, 50, 100, and 200 mg/ml from Mueller Hinton broth. Then, 1 ml of the active bacterial suspension (1.5×10^6 CFU/ml) was added to each dilution; positive control (medium containing bacterial and without extract) and negative control (medium without bacterial) were used. Finally, the tubes were incubated at 37 °C for 24h. After incubation, the tubes were examined for turbidity due to bacterial growth; and the last dilution in which no turbidity was observed (non-growth) was considered as MIC.

Samples were obtained from all tubes with no growth. Minimum bactericidal concentration (MBC) was determined by plate culture. Plates were incubated for 24 hours at 37° C; the tube with the lowest concentration of extract that the bacterial growth was visible on its plate was considered as MBC of that substance. Each experiment was repeated 5 times to reduce the error of the experiment. Statistical analysis was performed using SPSS software. ANOVA and Chi-square tests were used to investigate the significant differences in the results. The difference between groups was determined.

Results

According to Table 1, the antibacterial activity of methanol extract of peppermint in quantitative and qualitative methods showed that this extract has a significant inhibitory effect on *S. aureus* and *B. cereus* bacteria. The inhibitory effect was enhanced by increasing the concentration of methanolic extract. This study proved that the inhibitory effects of peppermint methanolic extract on gram-positive bacteria were significantly higher compared to gram-negative bacteria. MIC and MBC values of methanol extract of peppermint against the tested bacteria revealed that, like the well diffusion method, peppermint extract on gram-positive bacteria has a higher bactericidal effect than gram-negative bacteria (Table 2). These results indicated that there was a significant difference between the tested bacteria in the sensitivity of peppermint (p <0.05).

Table 1: Mean diameter of non-growth zone of methanolic extract of peppermint against selected bacteria in millimeters (mean ±standard deviation)

<table>
<thead>
<tr>
<th>Concentration of Extract (mg/ml)</th>
<th>Bacteria Strain</th>
<th>20</th>
<th>30</th>
<th>50</th>
<th>400</th>
<th>Negative control</th>
<th>Positive control</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td><em>S. aureus</em></td>
<td>9.8 ± 0.83</td>
<td>15.2 ± 0.83</td>
<td>20.6 ± 1.15</td>
<td>25.6 ± 1.14</td>
<td>--</td>
<td>23</td>
</tr>
<tr>
<td></td>
<td><em>B. cereus</em></td>
<td>8.4 ± 1.14</td>
<td>14.4 ± 1.14</td>
<td>18.4 ± 0.57</td>
<td>24.4 ± 1.14</td>
<td>--</td>
<td>21</td>
</tr>
<tr>
<td></td>
<td><em>E. coli</em></td>
<td>0</td>
<td>11 ± 0.70</td>
<td>15.2 ± 0.83</td>
<td>19.8 ± 1.15</td>
<td>--</td>
<td>25</td>
</tr>
<tr>
<td></td>
<td><em>P. aeruginosa</em></td>
<td>0</td>
<td>0</td>
<td>9.8 ± 1.30</td>
<td>12.4 ± 1.67</td>
<td>--</td>
<td>24</td>
</tr>
</tbody>
</table>

Table 2. MIC and MBC values of methanol extract of peppermint (mg/ml)

<table>
<thead>
<tr>
<th>Concentration of Extract</th>
<th>MIC</th>
<th>MBC</th>
</tr>
</thead>
<tbody>
<tr>
<td><em>S. aureus</em></td>
<td>6.25</td>
<td>12.5</td>
</tr>
<tr>
<td><em>B. cereus</em></td>
<td>12.5</td>
<td>25</td>
</tr>
<tr>
<td><em>E. coli</em></td>
<td>50</td>
<td>100</td>
</tr>
<tr>
<td><em>P. aeruginosa</em></td>
<td>100</td>
<td>200</td>
</tr>
</tbody>
</table>

Discussion

Nowadays, bacteria resistance has enhanced to a variety of antibiotics, therefore, many efforts have been made to obtain and use new compounds. Some of these compounds come from plants. Plants have played an important role in maintaining health and improving the quality of human life for thousands of years (11). *Mentha Piperita* that is the most popular...
In-vitro antibacterial effects of methanolic extract of peppermint

Mahmoudi S. et al.

member of this family, commonly known as
Peppermint, has long been used as an
aromatic and appetizing plant. Medicinal
properties of peppermint include
antispasmodic, anti-vomiting, and anti-
bloating (12). Peppermint is one of the most
widely consumed medicinal plants with an
annual consumption of about 7,000 tons
worldwide (13). Tylor VE in 1993 introduced
peppermint as a rich source of metabolites
that have major nutritional uses and have also
antispasmodic and antibacterial features (14).
Iscan et al. in 2002, found that
lipopolysaccharides in the external membrane
of gram-negative bacteria are responsible for
their higher resistance to antibacterial agents
(15). Dorman HJD et al. stated in their studies
that some essences such as that of peppermint
are highly efficient on E.coli (16). In 2005,
Moreira MR et al. surveyed the antimicrobial
effects of different plants on various species
of E.coli; and the essence of peppermint
showed significant antimicrobial effects (17).
Antibacterial activity of peppermint oil
against E. coli and S. aureus and antioxidant
activity were surveyed by Rasouli et al. in
2008 (18). Another research by Kaur et al. in
2010 reported that the peppermint leaf extract
has antibacterial activity because of tannins
and flavonoids. Further study also confirmed
the antimicrobial effect in the essence of
peppermint (19). Singah et al. Concluded in
2011 that peppermint extract has a greater
effect on gram-positive bacteria than gram-
negative bacteria; the diameter of non-growth
on negative gram-negative bacteria such as E.
coli and Klebsiella pneumonia is 12.4 and 5.1
mm, respectively, while in the case of gram-
positive bacteria such as Streptococcus
pyogenes and S.aureus are 17.2 and 13.1 mm,
respectively (20). In 2012, Pramila DM
examined the antifungal effect of the Mentha
Piperita in his research and was able to
confirm it (21). Another research done in
2015 approved that previous studies have
verified that thymol, carvacrol, menthol, and
parasiman are the most important components
in the antimicrobial activity of peppermint
essence. Furthermore, it has been confirmed
that as the concentration of these compounds
in peppermint essence becomes greater, its
antimicrobial activity will be more (22).
Moreover, Zaidi et al. studied the antimicrobial activity of essential oil from
Mentha piperita against 4 fungal and 11
bacterial clinical isolates; it showed the
maximum activity against S. aureus,
producing a zone of inhibition of 19.2 ± 0.07
mm (23).

Conclusion

One of the reasons for the difference MIC in
different studies is the differences in the
composition of the extracts. The composition
of the extracts from a species can vary based
on the region's geography, harvest season,
plant age, growth stage, and the method of
drying and extraction. In general, the plant
extract has the highest antimicrobial activity
during flowering or immediately after
flowering. Further, the extracts obtained from
different parts of a particular plant have
different antimicrobial activity. Additionally,
the sensitivity of different bacteria to different
extracts is different. It is evident that the
extract derived from the peppermint plant has
antimicrobial activity against Staphylococcus
aureus, Bacillus cereus, and Escherichia coli,
and it can be used as alternative medicine
after further investigations on laboratory
animals and its side effects.
References


18. Rasooli I, Gachkar L, Yadegarinia D, Bagher Rezaei M, Alipoor Astaneh S. Antibacterial and antioxidative characterisation of essential oils


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