CHEMICAL CHARACTERIZATION AND ANTIMICROBIAL ACTIVITY OF SOME ESSENTIAL OILS AFTER THEIR INDUSTRIAL LARGE-SCALE DISTILLATION

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ABSTRACT

Essential oils (EOs) are complex mixtures of several components that show a wide range of biological activities. The objective of the present work was to investigate the composition of the essential oils from industrial large-scale distillation, and evaluate their antimicrobial activity on typical and clinic isolates strains. The qualitative compositions of essential oils of several aromatic plants were determined by GC-FID after their industrial large-scale distillation (Calendula Co., Nova Lubovna, Slovakia). GC-FID analysis revealed that essential oils from large-scale show different qualitative and quantitative content in comparison with literature data. Eleven essential oils, namely, Rosmarinus officinalis L., Thymus vulgaris L., Mentha × piperita L., Matricaria recutita L., Hyssopus officinalis L., Salvia officinalis L., Pimpinella anisum L., Juniperus communis L., Abies alba L., Pinus silvestris L., and Coriandrum sativum L. were tested for their antimicrobial activities using disk diffusion method. Thymus vulgaris L. essential oil shows the most potent antimicrobial activity. On other hand, the essential oil of Hyssopus officinalis L., Mentha × piperita L., Rosmarinus officinalis L., and Coriandrum sativum L. The essential oil show medium antimicrobial activity. There was no antimicrobial activity observed for Matricaria recutita L.

Keywords: essential oil content, chemical analysis, constituents, large-scale distillation, antimicrobial activity

INTRODUCTION

Over the past few decades, research has focused on the health effects of phytochemicals and plant-derived extracts (Newman and Cragg, 2007). In recent years, the varied therapeutic potential of EOs attracted the attention of researchers for their antimicrobial activity (Hercules et. al., 2017; Inouye et. al., 2001; Rota et al., 2004). Modern pharmaceutical science is showing high degree of towards the herbal products as compared to the synthetic products. Herbal medicine is considered to have better affordability, acceptability and compatibility with the human physiology and minimal side effects (Al-Asmari et. al., 2014). The use of herbal products for the medicines, perfumery, cosmetics and food industry and agriculture are increasing among common people, patients and physicians as evident form an increased market of herbal products (Bernal et. al., 2013; Musthaha et. al., 2010). In recent years, large improvement in breeding methods, cultivation, harvesting and processing techniques make produce EOs in large scale possible. Calendula, Co. Company in Nova Lubovna, Slovakia is considered as one of the leader regards to large production of EOs. This company has been producing medicinal EOs and extracts since 1999 (Salamon, 2014). The composition of the aromatic and volatile oils are influenced by ecological, genotype and technological such as; cultivation, types of collection, storage of crude material and processing technique (Mancini et. al., 2015). Interestingly, EOs yield and composition from miniature research distillation equipment are differed from large commercial distilleries for aromatic plant of the same species (Mitchell and Crowe, 1996). As the effect of large scale production on the quantitative and qualitative content are known, this study attempt was made to characterize the chemical composition of the eleven aromatic plants raw-material after industrial large-scale distillation and investigate the effect of large scale production on their antimicrobial activities.

MATERIAL AND METHODS

The raw materials of different aromatic plant species: Juniperus communis L., Coriander (Coriandrum sativum L.), Pine (Pinus silvestris L.), Fir (Abies alba L.), Anise (Pimpinella anisum L.), Sage (Salvia officinalis L.), Hyssop (Hyssopus officinalis L.), Chamomile (Matricaria recutita L.), Peppermint ( Mentha × piperita L.), Thyme (Thymus vulgaris L.), Rosemary (Rosmarinus officinalis L.) for large-scale essential oil isolation were donated by established growers in Slovakia (Agrokarpaty, Ltd., Pivnica, Biox, Ltd., Michalovce and Mipros, Ltd., Nitra) and imported from cooperation companies in Poland (Herbar, Ltd., Milejow), Czech Republic (Pvni jilovska, Co., Jilove u Prahy) and Ukraine (Sumyfiofarmacia, Ltd., Sumy). The raw stock presents different fresh and dry plant parts (fruits, herbs, needles, flowers and seeds), which are chopped after distillation process.

Isolation of essential oils

EOs were obtained by the large-scale distillation apparatus (Oravec et. al., 1988) specifically designed for aromatic and medicinal plants. There are two types of distillation apparatus; type ‘ HV-3000 ‘ (height: 5.250 mm, width: 2.180 mm, with container for 200 - 250 kgs of dried plant material and for 400 - 500 kgs of fresh plant material) and type ‘ HV-300 ‘ (height: 3.400 mm, width: 1.300 mm, with container for 40 - 50 kgs of dry plant material and 100 - 120 kgs of fresh plant material). Both types were developed and technologically upgraded from 2000 to 2007 (Bucko and Salamon, 2007), and proved to be very successful by Calendula Co., in Nova Lubovna, Slovakia (Fig. 1). The first type is used for larger quantities of plant raw materials during their harvest and transport from cultivation fields. The second type for lower quantities of dry plant parts during a winter time (outside a vegetation season). The mass yield of essential oil is higher in this case. The containers of EOs were stored under N₂ at +4 °C in a dark space. Each branches of EOs is determined on their chemical composition in regard to requirements of customers.

GC-FID analyses

The analysis of EOs was carried out using a gas chromatograph Varian 3090, connected to MS Saturn 2100T integrator. The following operating conditions were used: capillary column: RX-5MS, 30 m x 0.250 mm i.d., film thickness: 0.25 μm, carrier gas: He, adjusted to a flux of 1.5 ml/min, injection and FID-detector temperatures: 220 °C respectively 250 °C, a capacity of sample injection: 2 μl, MS-detector with automatic injector type 1177.
Components were identified by their GC retention times, and the resulting values were comparable to those of literature. Oil component standards for comparison were supplied by Extrasythese, Merck, Fulkra, Sigma a Roth.

Antimicrobial activity

Antimicrobial activity of EO’s was determined using disk diffusion method. Sterile filter paper discs (6 mm in diameter) impregnated with 10 μL of essential oil was placed on the dish plate previously inoculated with a microbial suspension. Bacterium and yeast inocula 100 μL in physiological solution were adjusted to the equivalent of 0.5 McFarland standard, and evenly spread on Muller-Hinton agar surface (incubated at 37±2 °C for 24 h), yeasts – on SDA agar (incubated at 35±2 °C for 48 h). The diameters of the inhibition zones were measured in millimeters including diameter of disc (Rhos et al., 2005). Each antimicrobial assay was performed at least three times.

As test culture, the following microorganisms from the ATTC (American Type Culture Collection, USA) collection were used: Candida albicans ATCC 885-653; Staphylococcus aureus ATCC 25923; Escherichia coli ATCC 25922. Enterococcus faecalis ATCC 29212, Streptococcus pyogenes ATCC 19615, Pseudomonas aeruginosa ATCC 27853. Also we used clinical strains of bacteria and yeast, which were isolated from sputum of different patients suffering from pneumonia, obstructive bronchitis, bronchial asthma, chronic obstructive disease and oral cavity patients with periodontal disease.

As a positive control were used: gentamicin (10 mg/disk) for Gram-negative bacteria, ampicilin (10 mg/disk) for Gram-positive bacteria, nystatin (100 UI) for Candida. As negative control were used DMSO.

For the results of experiment, we used statistical software Microsoft Office-Excel (2013) with the calculation of averages, error, and standard deviation.

RESULTS AND DISCUSSION

The large-scale technology of essential oil distillation in the Slovak Republic consists of a main distilling apparatus, a steam condenser, and an additional apparatus (Fig. 1). The shape of the main distilling apparatus is a funnel. It is thermal-isolated and made from stainless steel. An inside screw-plate is driven by an electric engine, which is installed on the apparatus. This screw-plate works as an exceptional stirrer. With regard to this system, the container has a mixing apparatus which is not a usual feature for many other types of commercial equipment. This is extremely useful for a complete distillation procedure and high yield of essential oils. The source of steam flow is a boiler (heated by oil, gas or electricity), and the flow is controlled mechanically, according to the plant mass and cooling needs. The length of distillation depends on medicinal plant species, which are used to isolate essential oils. Essential oil collector has a volume of 75 liter. (Bucko and Salamon, 2007)

In general, many factors could contribute to the differences between yield and composition of essential oils for industrial stills, for example, steam distribution, differences in size of the distillation unit, materials (such as glass versus aluminum), and temperature of condensing units (Mitchell and Crowe, 1996).

Chemical composition of studied essential oils

The EOs have received substantial attraction due to their significant biological usefulness (Miladi et al., 2013). The chemical composition of essential oil can also vary according to the geographical localization (Jaafari et al., 2007). In 1996, Mitchell and colleagues investigated whether a mini-still produces oil yield and composition similar to a larger commercial distillery. The result of their study revealed that was a different qualitative and quantitative content between mini-still produces and larger commercial production (Mitchell and Crowe, 1996). Table 1 shows the chemical composition of studied EOs. Highly active component α-pinene found in high percent in composition of Juniperus communis L. and Pinus sylvestris L. with 52 and 38 %, respectively. α-pinene has been shown to have anti-inflammatory, antimicrobial and exhibits activity as an acetylcholinesterase inhibitor, aiding memory (Russo, 2011; Nissen et al., 2010). Whereas, (−)-α-pinene is a positive modulator of GABAA receptors. It acts at the benzodiazepine binding site (Yang et al., 2016). Interestingly, Pimpinella anisum L. essential oil shows trans- cis-anetholes as one major component with 89.3 ±1.5%. This find is useful for herbal medicinal field. Anetholes has wide range of biological activity (De M et. al., 2002; Bone and Mills, 2013).

Many searches revealed that the cytotoxic activity of Thymus vulgaris essential oil towards cancer cells was mainly due to thymol. Table1 show that the two major contents for this plant species are thymol and p-cymene with 32 and 40 %, respectively. The effect of large distillation can be recognized clearly compared with literature, for example the reported analysis for Thymus vulgaris major content thymol was found to be 49.1, 41.33 and 59.15 % in three different studies (Nikolić et. al., 2014; Miladi et. al., 2013; Ramadan et. al., 2015). This variation in the quantitative content of EO is in agreement with previous study (Mitchell and Crowe, 1996). Rosemary essential oil, for instance, contains mainly α-pinene, p-cymene, α-terpinol, camphor, camphene, cineole and β-pinene as major content. The larger commercial distilleries were nearly identical to mini-stills produced oil in regard to compare between the results, which we obtained and reported. (Santoyo et. al., 2005), but oil constituents differed between the two stills, which may have been resulted from dissimilar harvest and distillation practices (Mitchell and Crowe, 1996).
### Table 1 Chemical composition of studied essential oils of aromatic plants after a large-scale distillation and its qualitative-quantitative characteristics [%, ±t × se. ]

| Essential oils          | α-Pi | Bo  | c/tA | Men | p-Cy | LnL | BoA | α-To | Bon | Cph | Met | Mrn | l-Me | Thy | Lim | Cam | BoS | Ch  | ThJ | PiC | i-PiC | Cin | β-Pi | SaN |
|-------------------------|------|-----|------|-----|------|-----|-----|------|-----|-----|-----|-----|------|-----|-----|-----|-----|-----|-----|-----|-----|-----|-----|-----|-----|
| *Coriandrum silvestris* | 17.0 | -   | -    | 2.0 | -    | 1.0 | 2.0 | -    | 2.0 | 19.0| -   | -   | -    | -   | -   | -   | -   | -   | -   | -   | -   | -   | -   | -   |
| *Pinus alba*            | 2.0  | -   | -    | 3.0 | -    | -   | 2.0 | -    | 2.0 | 2.0 | -   | -   | -    | -   | -   | -   | -   | -   | -   | -   | -   | -   | -   | -   |
| *Abies communis*        | 5.0  | -   | -    | 1.0 | -    | 1.0 | 2.0 | -    | 2.0 | 2.0 | -   | -   | -    | -   | -   | -   | -   | -   | -   | -   | -   | -   | -   | -   |
| *Juniperus silvestrii*  | 2.0  | -   | -    | 2.0 | -    | 1.0 | 2.0 | -    | 2.0 | 2.0 | -   | -   | -    | -   | -   | -   | -   | -   | -   | -   | -   | -   | -   | -   |
| *Artemisia dracunculus* | 8.0  | -   | -    | 1.0 | -    | 1.0 | 2.0 | -    | 2.0 | 2.0 | -   | -   | -    | -   | -   | -   | -   | -   | -   | -   | -   | -   | -   | -   |

Antimicrobial activity of essential oils

Presently, there are a considerable number of studies on the antibacterial effect of EOs upon microorganisms; different plant chemotypes are also known to have different degrees of antimicrobial effect (Koščiva et al., 2006; Kačičnová et al., 2017). Up to our knowledge there is no study has been done on the antimicrobial activities of EOs prepared on industrial large scale. Both Table 2 and 3 shows the antimicrobial activities for eleven EOs namely, Rosmarinus officinalis L, Thymus vulgaris L, Mentha × piperita L, Matricaria recutita L, Hyssopus officinalis L, Salvia officinalis L., Pimpinella anisum L, Juniperus communis L, Abies alba L., Pinus silvestris L, and Coriandrum sativum L. Tested on typical and clinical strain, respectively. A broad-spectrum antimicrobial activity of Thymus vulgaris EO founds against Gram-positive, Gram-negative bacteria and microfungal of Candida genus (Table 2 and 3).

Table 2 Antimicrobial activities of the essential oils against typical strains

<table>
<thead>
<tr>
<th>№</th>
<th>Essential oils</th>
<th>Staphylococcus aureus ATCC 25923</th>
<th>Escherichia coli ATCC 25922</th>
<th>Enterococcus faecalis ATTC 29212</th>
<th>Streptococcus pyogenes ATCC 19615</th>
<th>Pseudomonas aeruginosa ATCC 27853</th>
<th>Candida albicans ATCC 885-653</th>
</tr>
</thead>
<tbody>
<tr>
<td>1</td>
<td>Thymus vulgaris L</td>
<td>33.00±0.25</td>
<td>12.66±0.33</td>
<td>27.26±0.60</td>
<td>12.70±0.44</td>
<td>-</td>
<td>71.00±0.58</td>
</tr>
<tr>
<td>2</td>
<td>Hyssopus officinalis L</td>
<td>17.00±0.20</td>
<td>14.00±0.56</td>
<td>8.33±0.33</td>
<td>11.00±0.57</td>
<td>-</td>
<td>11.50±0.20</td>
</tr>
<tr>
<td>3</td>
<td>Mentha × piperita L</td>
<td>12.00±0.10</td>
<td>8.33±0.33</td>
<td>9.66±0.17</td>
<td>10.33±0.33</td>
<td>-</td>
<td>15.00±0.57</td>
</tr>
<tr>
<td>4</td>
<td>Rosmarinus officinalis L</td>
<td>8.33±0.33</td>
<td>8.56±0.29</td>
<td>7.83±0.44</td>
<td>8.33±0.33</td>
<td>-</td>
<td>12.33±0.33</td>
</tr>
<tr>
<td>5</td>
<td>Coriandrum sativum L</td>
<td>11.00±0.20</td>
<td>15.50±0.29</td>
<td>15.17±0.46</td>
<td>11.00±0.25</td>
<td>-</td>
<td>-</td>
</tr>
<tr>
<td>6</td>
<td>Abies alba L</td>
<td>7.33±0.33</td>
<td>10.00±0.15</td>
<td>-</td>
<td>-</td>
<td>-</td>
<td>30.00±1.25</td>
</tr>
<tr>
<td>7</td>
<td>Salvia officinalis L</td>
<td>9.83±0.44</td>
<td>7.83±0.44</td>
<td>8.33±0.33</td>
<td>-</td>
<td>-</td>
<td>10.17±0.17</td>
</tr>
<tr>
<td>8</td>
<td>Pimpinella anisum L</td>
<td>9.83±0.44</td>
<td>12.16±0.44</td>
<td>-</td>
<td>-</td>
<td>-</td>
<td>-</td>
</tr>
<tr>
<td>9</td>
<td>Juniperus communis L</td>
<td>7.50±0.10</td>
<td>-</td>
<td>-</td>
<td>-</td>
<td>-</td>
<td>15.00±0.25</td>
</tr>
<tr>
<td>10</td>
<td>Pinus silvestris L</td>
<td>-</td>
<td>-</td>
<td>-</td>
<td>-</td>
<td>-</td>
<td>8.00±0.10</td>
</tr>
<tr>
<td>11</td>
<td>Matricaria recutita L</td>
<td>-</td>
<td>-</td>
<td>-</td>
<td>-</td>
<td>-</td>
<td>-</td>
</tr>
</tbody>
</table>

Zones inhibition, mm (× ± s); «» − no inhibition.

Thyme EO shows strong antimicrobial effect on Staphylococcus aureus and Candida albicans. Regard to clinical strains, thyme oil shows potent effect on each of Staphylococcus aureus, Candida albicans and Escherichia coli with inhibition zone 50.00±0.25 mm, 38.00±0.80 mm and 25.33±0.88 mm, respectively. Our results are in agreement with those obtained by Rota et al. (2004), they reported that Satureja montana and Thymus vulgaris EOs showed the high potent inhibitory effect against the gram-positive Listeria monocytogenes and Staphylococcus aureus, and the gram-negative Salmonella enteritidis, Escherichia coli O157:H7, Yersinia entercolitica, and Shigella flexneri. Furthermore, the result presented by Sakkas and Papadopoulos (2017) shows that Thymus oil has a broad spectrum of antimicrobial activity against Campylobacter jejuni, Salmonella enteritidis, Pseudomonas aeruginosa, Klebsiella pneumonia. In addition, these same oils exhibited moderate effects against multi-resistant clinical isolates namely; Acinetobacter baumannii, Escherichia coli, Klebsiella pneumoniae, and Pseudomonas aeruginosa. Table 2. Several previous studies reported that the essential oils of Thymus vulgaris, and their main phenolic constituents, such as thymol show remarkable antimicrobial, antibacterial and antifungal activities (Raut et al., 2013; Laranjo et al., 2017; Rota et al. 2004). Interestingly, thymol is one of the major content of Thymus vulgaris EO.

Table 3 Antimicrobial activities of the essential oils against clinical strains.

<table>
<thead>
<tr>
<th>№</th>
<th>Essential oils</th>
<th>Staphylococcus aureus</th>
<th>Escherichia coli</th>
<th>Streptococcus pyogenes</th>
<th>Pseudomonas aeruginosa</th>
<th>Candida albicans</th>
</tr>
</thead>
<tbody>
<tr>
<td>1</td>
<td>Thymus vulgaris L</td>
<td>50.00±0.25</td>
<td>25.33±0.88</td>
<td>11.33±0.33</td>
<td>-</td>
<td>38.00±0.80</td>
</tr>
<tr>
<td>2</td>
<td>Hyssopus officinalis L</td>
<td>20.00±0.10</td>
<td>10.66±0.88</td>
<td>11.33±0.33</td>
<td>-</td>
<td>12.00±0.80</td>
</tr>
<tr>
<td>3</td>
<td>Mentha × piperita L</td>
<td>10.00±0.15</td>
<td>9.67±0.33</td>
<td>10.00±0.25</td>
<td>-</td>
<td>12.33±0.88</td>
</tr>
<tr>
<td>4</td>
<td>Rosmarinus officinalis L</td>
<td>8.00±0.10</td>
<td>8.33±0.33</td>
<td>13.00±0.50</td>
<td>-</td>
<td>-</td>
</tr>
<tr>
<td>5</td>
<td>Coriandrum sativum L</td>
<td>10.00±0.20</td>
<td>11.00±0.58</td>
<td>13.33±0.33</td>
<td>-</td>
<td>-</td>
</tr>
<tr>
<td>6</td>
<td>Abies alba L</td>
<td>-</td>
<td>-</td>
<td>-</td>
<td>-</td>
<td>20.00±1.15</td>
</tr>
<tr>
<td>7</td>
<td>Salvia officinalis L</td>
<td>12.00±0.10</td>
<td>9.67±0.33</td>
<td>8.00±0.57</td>
<td>-</td>
<td>11.00±0.35</td>
</tr>
<tr>
<td>8</td>
<td>Pimpinella anisum L</td>
<td>-</td>
<td>-</td>
<td>-</td>
<td>-</td>
<td>21.33±0.88</td>
</tr>
<tr>
<td>9</td>
<td>Juniperus communis L</td>
<td>7.00±0.01</td>
<td>-</td>
<td>-</td>
<td>-</td>
<td>10.00±0.58</td>
</tr>
<tr>
<td>10</td>
<td>Pinus silvestris L</td>
<td>-</td>
<td>-</td>
<td>-</td>
<td>-</td>
<td>-</td>
</tr>
<tr>
<td>11</td>
<td>Matricaria recutita L</td>
<td>-</td>
<td>-</td>
<td>-</td>
<td>-</td>
<td>-</td>
</tr>
</tbody>
</table>

Zones inhibition, mm (× ± s); «» − no inhibition.

EOs from Juniperus communis L, Pinus silvestris L, Matricaria recutita L. and Pimpinella anisum L. had weak antimicrobial and antymycotic activity. In 2017, Reddy and colleagues investigated the chemical composition and the antimicrobial activity for Mentha × piperita EO which cultivated in Saudi Arabia. The major content is 36.02% of menthol and 24.56 % of menthon which similar to the result reported in Table 1. It is worth mentioning, the high similarity in the content of EOS was found but biological activity was significantly different (Reddy et al., 2017). In the same manner, Semencic and colleagues (2017) show a higher activity of thyme EO against typical strains of Escherichia coli and Pseudomonas aeruginosa in comparison to thyme EO that reported in this study. Whereas, The Thymus vulgaris EO produced in large scale shows more expressed inhibiting effect toward Staphylococcus aureus. Our results are concordant with Sharma et al. (2002) investigation regard to the anti-microbial activity of Coriandrum sativum L. EO, produced by mini steam distillation, toward Staphylococcus aureus and Candida albicans clinical strains. The work of Inouye et al. (2001) establishes the antibacterial activity of the essential oil samples obtained from various countries. It ascertainment high antibacterial activity of Cinnamon oil and Thyme oil and moderate anti-microbial activity of Tea tree oil and Peppermint oil, which in fact fits with our data. Thereby, our study results shows that essential oils produced in large-scale distillation conditions had an antibacterial activity. According to literature data and our results there was some difference in antimicrobial activity.

CONCLUSION

Our results confirmed higher activity of EOs on Gram-positive bacteria and yeasts than against Gram-negative. They indicated that the highest activity was observed using EO from Thymus vulgaris L. The maximum antymycotic activity was shown by the EOs from Thymus vulgaris L. and Abies alba L. Where each of Hyssopus officinalis L, Mentha × piperita L, and to Coriandrum sativum L. EOs shows moderate antimicrobial activity. On other hand, the effect of EOs toward antibacterial clinical strains shows promising result and this open more area for further investigation. However their antymycotic activity against clinical strains of microfungal fungi was weaker than against their typical strains. The obtained results have proved the need of further studies of the impact of essential oils upon bacterial isolates, including those with multiple resistances to medical preparations. The fact that phytomaterials much less often than antibiotics provoked the formation of resistance in bacteria and microfungal fungi.
References


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