Antioxidant activities, total phenolic and flavonoid contents of honey collected from different botanical origins

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Abstract: In this study, it is aimed to determine the presence of antioxidant capacity, total phenolic and flavonoid contents in six different types (multiflora, pine, chestnut, sunflower, acacia, citrus) and eventually 65 samples of honey from different parts of Turkey. Pollen analysis of all honey samples in the laboratory was carried out to determine the purity (> 65-70%) of the plant source. Total phenolic content determined in honey samples was found the highest value in pine honey 166.46 ± 5.80 (mgGAE / 100 g honey) while the lowest value was found in flower honey with (106.04 ± 9.55). The level of flavonoid contents of the groups was lowest on the flower and citrus honey (1.3 ± 0.2 and 1.6 ± 0.1) and the highest value were on chestnut and pine (2.7±0.4 and 2.8 ± 0.2) were detected. Comparing the radical scavenger activity in honey groups, the activity of chestnut honey was the highest (100.54 ± 22.72). The results of this study show that the phytochemical structure and biological activity of honey are completely different from each other depending on the plant source. On the other hand, it is possible to say that the antioxidant, phenolic, and flavonoid values are high, which is a good indicator of the quality and naturalness of honey.

Keywords: Antioxidant, flavonoid, honey, honey bee, phenolic acid.

Introduction

For centuries, bees and bee products have been used for therapeutical purposes. Today, they maintain their actuality in the field of supportive therapy and show a fast development under the name of apitherapy. Honey, which is one of the apitherapy products and one of the oldest nutritional sources of mankind, is a sweet product which bees collect from flower nectars or from secretions of some insects living on plants and store in honeycombs by exposing them to enzymatic change (3, 6, 22). The formation and composition of honey produced naturally differ significantly according to regions and sources of the plant (23). The composition of plant nectar differs according to the geography the plant is grown in, soil fertility, rainfall, light, altitude and a great number of other environmental factors (14, 24). In other words, the quality and biochemical characteristics of honey differ according to the source of nectar (8, 13, 22). Studies conducted show that the antioxidant activity of natural foods is higher than food products with synthetic structure (18). For this reason, daily intake of antioxidant food is important and required for free radicals which form as a metabolism product in the organism. In addition to being a food product and source of energy, honey is also important for human health as a natural antioxidant source due to various phytochemicals it includes (3, 12, 17).
Antioxidant compositions of honey are enzymatic (glucose oxidase, catalase, peroxidase) or non-enzymatic (phenolic acids, flavonoids, ascorbic acid, tocopherol, carotenoids) (18, 19). Phenolic acids contribute significantly to the antioxidant activity of honey (1, 17). Flavonoids have a significant antioxidant and anti-inflammatory function by providing free-radical scavenging, inhibiting cyclooxygenase and lipoxygenase enzymes, chelating transition metals such as iron and copper, protecting α-tocopherol at LDL and providing oxidizable ascorbic acid regeneration (16). The rates of these compositions in honey differ significantly depending on the source flora honey is collected from and the structure of the geographical areas (8, 12, 23). For this reason, every honey has a different apitherapic value. The present study researches the biological activities of honey obtained from different regions and plant sources of Turkey.

**Materials and Methods**

**Sampling:** The study was conducted with a total of 65 honey samples from different regions of Turkey (11 sample meadow honey from the provinces of Erzurum and Sivas, 11 sample pine honey from Muğla, 11 sample chestnut honey from Sinop, 11 sample Acacia honey from Trabzon, 10 sample from citrus tree honey from Antalya and 11 sample sunflower honey from Samsun). Pollen analyses of all samples were made in the laboratory and purity degree of plant sources was found (>65-70%).

**Biochemical analysis:** Calculation of the total antioxidant effect in the samples was made according to the free radical scavenging effects of the prepared extracts on DPPH (1,1-diphenyl-2-picyrylhydrazyl) in honey by using an indirect method. As a result of DPPH radical scavenging activity, IC50 values were assessed as mg/ml. The measurement was made according to Meda and Dimins method modified at 520 nm with a spectrophotometer (9, 21). AA [%] = (Abs cont-Abs sample)/ Abs cont.X100. Total phenolic concentration level was read at 750 nm in spectrophotometer with Folin-Ciocalteau method and the total phenol amount was calculated as equivalent to mg gallic acid in 100 gram extract (4, 9). The Dowd method was used in total flavonoid analysis. In this method, honey solutions prepared based on quercetin (mgQE/100gr) concentration in mg in a kilogram of honey were read at 415 nm with a spectrophotometer (9, 21).

**Statistical analysis:** The data obtained from the study were evaluated with the variance analysis (ANOVA) technique in factorial order and the differences between the means were determined by Duncan multiple comparison test. Statistical evaluations were made using the SPSS statistical program (26).

**Results**

Average values of total phenolic, flavonoid levels and antioxidant activity found in honey samples of groups are given in (Table 1, Figure1).

![Figure 1](image-url) Means of total phenolic, flavonoid and antioxidant analysis results in honey samples

| Samples        | Total flavonoid (Min-Max) | Total phenolic (Min-Max) | Antioxidant, DPPH-sc₅ₒ (Min-Max) |
|----------------|---------------------------|--------------------------|----------------------------------|
| Citrus tree    | 1.6 ± 0.1, 1.2 - 1.9      | 113.77 ± 4.27, 100.21-120.62 | 152.65 ± 4.96, 140.40-163.91    |
| Chestnut       | 2.7 ± 0.4, 1.4 - 4.0      | 138.27 ± 19.31, 97.64-206.22 | 140.54 ± 22.72, 100.54-206.38   |
| Sunflower      | 1.8 ± 0.5, 1.7 - 2.0      | 127.27 ± 13.54, 88.03-127.33 | 167.33 ± 9.69, 135.40-198.81    |
| Pine           | 2.1 ± 0.2, 2.1 - 3.6      | 166.46 ± 5.80, 149.37-186.11 | 145.48±16.49, 93.61-206.38      |
| Meadow         | 1.3 ± 0.2, 0.5 - 1.9      | 106.04 ± 9.55, 68.85-130.94  | 270.86±51.22, 136.17-506.38     |
| Acacia         | 2.1± 0.4, 1.3 - 3.5       | 143.71 ± 16.99, 71.16-183.06 | 108.91 ± 54.00, 69.34-142.55    |
The highest total phenolic composition was found in pine honey with 166.46 ± 5.80 (mgGAE/100g honey), while the lowest value was found in flower honey with 106.04 ± 9.55. While the intergroup flavonoid composition lowest value was found in flower and citrus honey (1.30 ± 0.2 and 1.6 ± 0.1), the highest value was found in chestnut and pine honey (2.7 ± 0.4 and 2.8 ± 0.2). Radical scavenging activity comparison of honey groups showed that chestnut honey had the highest activity (100.54 ± 22.72) As a result of the analysis of variance, there was a significant difference between the honey samples (P <0.01).

Discussion and Conclusion

Phenolic acids and flavonoids, which are components of honey, are important due to their antimicrobial, antioxidant, anticancer and antioxidative effects and their being associated with human and animal health and on the other hand due to being a criterion for honey’s being refined (1, 5, 6, 22). The concentration of these compounds which significantly affects honey’s therapeutic activity differs according to plant flora which constitutes the sources of honey (11).

In the present study, it was found that total phenolic composition differed between 106 - 166 mgGAE/100 g in all honey samples. While the lowest concentration was found in flower honey, the highest concentration was found in pine honey (Figure1). Similarities and differences are seen between our study results and the results of the studies previously conducted. While total phenolic acid concentration was reported as 32.59-114.75 mgGAE/100 gr and an average of 74.38 ± 20.54 in South African honey samples (21), phenolic acid concentration was reported as 36.26-102.80 mgGAE/100g and an average of 65.31±19.50 in a study conducted with Nigeria honey (5). Vit et al., (28) reported in their study that phenolic matter amount differed between 38.15 and 182.10 mgGAE/100 g in Venezuela honey. Total phenolic amount in Yemeni honey obtained from different regions and different plants differed between 56.32 and 246.21 mg/100g (1). In a study conducted on the honey in the Indian Bengal region, it was reported that gallic acid rates per 100 g honey differed between 9.9 ± 0.6 and 44.7 ± 2 mg (7). In a study conducted with different honey from Slovenia, the total phenolic composition was found as 241.4 mg/kg in fir tree honey, as 233.9 mg/kg in chestnut honey and as 44.8 mg/kg in acacia honey (4). In studies conducted on acacia honey, total phenolic compositions differ significantly among countries. While Iranian acacia honey total phenolic content was found as 22.9-65.5 mg gallic acid/g (15), it was found as 627 ± 44.03 in Germany acacia honey (2). In a study conducted on the honey of Black Sea flora, average phenolic content was found as 0.224 mg as gallic acid equivalent for 1 g honey sample (11).

In the study, while the lowest average total flavonoid (mgQE/100g honey) value of honey was found in flower honey with an average of 1.3 ± 0.2 the highest value was found in chestnut honey with an average of 2.7 ± 0.4. In a study conducted on Indian honey, quercetin (mg/100g ) amount was found to be between 5.12 ± 0.23 and 19.4 ± 1.38 (Das et al., 2013). In Brazilian multiflora and citrus honey, quercetin (mg/100g) composition was reported as 1.96 ± 1.53 and 0.17 ± 0.15, respectively (20). In a study conducted on pine honey in our country, total flavonoid level was found as 22.80 ± 2.45 on average in QE/kg honey equivalent (25). In the present study, this value was found as 2.8 ± 0.2 in the same honey type.

In the study, total antioxidant activity was found as 100.54 ± 22.72 in chestnut honey and as 270.86 ± 51.22 in multiflora flower honey. When studies conducted in many countries were examined in terms of their similarities and differences of these values, it was found that radical scavenger DPPH average IC50 = 23.92 ± 1.12 mg/mL in Indian honey (7), the same value was found as 10.0 ± 1.8 and 10.7 ± 2.2, respectively in Slovenian chestnut and flower honey (4); average antioxidant activity was found as 31.96 ± 18.07 in chestnut honey in Brazilian multifloral honey and as 15.22 ± 10.75 in citrus honey (20), while antioxidative activity was reported as 35-122 in China unifloral honey (10). Ertürk et al. (11) found IC50 values as between 29.388 and 458.450 mg/mL as a result of the DPPH radical scavenging activity test in Black Sea flora honey in our country.

Some studies show a correlation between the phenolic compound level and antioxidant activity in some types of honey; however, since flavonoids have hydroxyl, they are oxidized very quickly. For this reason, despite structural similarities, the difference between antioxidant activities depends on hydroxylation and methylation degree (1). In our study, no correlation was found between groups. It is possible to say that this is because in addition to flavonoid and phenolic acid, the presence of Vitamin E and C and carotenoids may have influenced total antioxidant activity (27).

Consequently, our observations that phytochemical structure and biological activity of honey differ completely depending on the origins of the plant.
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Conflict of Interest
The authors declared that there is no conflict of interest.

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