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Analysis of anticancer taxanes in Turkish hazelnut (*Corylus avellana* L.) genotypes by high-performance liquid chromatography

KUTLUTÜRK et al. Taxanes in Turkish Hazelnut Genotypes

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Abstract:

Objectives: This study aimed to investigate the anticancer taxane profiles of edible and non-edible parts of seven Turkish hazelnut (*Corylus avellana*) genotypes. Hazelnut is one of the healthy foods rich in nutrients and antioxidants. Its regular consumption is associated with a reduced risk of coronary heart disease and cancer. Hazelnut has been described as a plant source that produces taxanes which are widely used in many cancers. Turkey is a homeland of hazelnut culture and has its own cultivars. Investigation of anticancer taxane profiles in different parts of Turkish hazelnut genotypes is important to show the potential and value of this plant from the perspective of the pharmaceutical and food industries.

Materials and Methods: In this study, green leafy covers-GLCs and hard shells-HSs (non-edible parts), skinless kernels-SKs, brown skins-BSs, and brown-skinned kernels-BSKs (edible parts) of 'Çakıldak', 'Sivri', 'Tombul', 'Palaz', and 'Kalınkara' as standard and 'Ham', and 'Sivri Yağlı' as local genotypes were used. The five parts of each genotype were ground to powder and eliminated to a size of less than 80 mesh. Each part was extracted by hexane and methanol for 10-deacetylbaccatin III (10-DAB III), baccatin III (BAC III), cephalomannine, and paclitaxel analysis as three replicates. Samples and standards were analyzed by acetonitrile:water gradient method on NOVA Spher 100 Phenyl-Hexyl C18 column in HPLC reverse phase system with 228 nm UV detector and 1.0 mL/min flow rate. Microsoft Office Excel, 2016, and ANOVA Jamovi Version 2.3 were used for statistical and data analysis, consecutively.

Results: Hazelnut parts differed to a very high degree from each other in terms of the highest amount of 10-DAB III (Ham HSs, 9,15 μ g/g), BAC III (Kalınkara BSs, 7.24 μ g/g), cephalomannine ('Sivri Yağlı' BSs, 6.37 μ g/g) and paclitaxel (Ham BSKs, 4.36 μ g/g) they contained. While HSs, BSKs, and BSs were rich in taxanes in all of the analyzed genotypes, SKs, and GLCs remain limited for anticancer taxanes.

Conclusion: This is the first report that revealed the differences in taxane contents of Turkish hazelnuts including previously untested standard and local genotypes and their parts. Significant differences between genotype and hazelnut parts are expected to highlight the health benefits of consuming raw Turkish hazelnut with brown skins and their possible use as a functional food. These results add more information to elucidate the bioactive potential of Turkish hazelnuts and their by-products and provide a promising resource for the food and pharmaceutical industry with an anticancer perspective.

Keywords: Corylus avellana, taxanes, anticancer, functional foods, by-products

1. Introduction

Among nut species, hazelnut (*Corylus avellana* L.), is the most valuable tree nut crop worldwide.¹ Hazelnut species exhibit distribution mainly in Turkey, Italy, Spain, Portugal, France, and some parts of the USA.² For the cultivation of high-quality hazelnut varieties, Turkey has suitable growing conditions.³ With 80 % of cultivated

hazelnut area in the world, Turkey is the leading producer and alone meets about 69 % of global hazelnut production (776 thousand tonnes).⁴

Hazelnut fruit is divided into three parts; these are green leafy cover (GLC), hard shell (HS), and hazelnut kernel which can be consumed raw (with brown skin) or roasted (without brown skin). In addition to their role as a popular snack both in Turkey and worldwide, approximately 90% of hazelnuts are utilized as an ingredient in a variety of processed foods, especially in confectionery and bakery products.⁵ The kernel, a main product of hazelnuts, plays a special role in human nutrition and health due to its particular composition. ⁶⁻⁸ The distinctive nutritional and sensory characteristics of hazelnuts render them an incomparable and optimal ingredient for food products. However, the kernel constitutes less than 50 % of the total nut weight, and substantial by-products (green leafy cover, hard shell, and brown skin) are removed during harvesting and processing.^{1,9} Among these by-products, none has nearly significant commercial value. The hard shell, a majority of the byproducts, is mainly used as a low-value heating source, and the green leafy covers are rarely used as organic fertilizers.³ The use of by-products as readily available in diverse bioactive compounds and functional food ingredients could become a valuable source ^{9,10} Therefore, in recent years, the number of studies that focused on investigating the utilization of hazelnut by-products has increased.¹⁰⁻¹² Studies on hard shells and brown skins mainly investigate the extraction, characterization, and identification of substantial bioactive molecules such as phenolic compounds, dietary fiber, cellulosic compounds, activated carbon, and pigments.^{5,13-15} In addition, researchers found that green leafy cover extracts contained bioactive compounds that have high antioxidant, antibacterial, and free radical scavenger properties.^{3,16} It has been shown that hazelnut is also a source of anticancer taxanes.¹⁷⁻²¹

Paclitaxel, also known as Taxol® is a representative of a class of diterpenes taxanes, which are widely used as a chemotherapeutic agent to stabilize microtubules in many types of cancer. Since the discovery of its antitumoral activity, Taxol[®] has been approved as a chemotherapeutic drug for the treatment of ovarian, breast, lung, bladder, prostate, melanoma, esophageal, and other types of solid tumors.²² It has also been used to treat AIDSrelated Kaposi sarcoma.²³ Taxol® is originally extracted in low yield from the bark of Taxus brevifolia. Chemical and partial synthesis of Taxol® from its precursors, 10-DAB III, BAC III, and cephalomannine are also very expensive and time-consuming.¹⁹ Although it was generally considered a particular metabolite of Taxus spp., paclitaxel extraction was carried out from hazelnuts.¹⁷ For more than two decades hazelnut and its by-products have been studied from the aspect of anticancer taxane contents. ¹⁷ In addition to paclitaxel, 10-DAB III, BAC III, and cephalomannine were isolated from hazelnut hard shells, green leafy covers, and leaves.^{18,19,20,24} As an alternative to taxol, the synthetic analog, taxotereTM (generic name Docetaxel), was acquired by semi-synthesis from baccatin III and 10-deacetylbaccatin III, two taxol precursor molecules. The American Food and Drug Administration (FDA) authorized the use of Paclitaxel and Docetaxel for treating ovarian and breast cancers.²⁵ On the other hand, the natural congener of paclitaxel, cephalomannine has shown anti-tumor activity in several different cell lines.²⁶ The cytotoxic effects of cephalomannine were first demonstrated in human glioblastoma cells.²⁷ Later on, Zhang et al.²⁸ demonstrated that cephalomannine significantly attenuates hepatocellular carcinoma progression in vitro and in vivo and bone metastasis in prostate cancer in vivo.²⁹ Therefore, cephalomannine has recently been evaluated as an anti-cancer drug candidate.²⁸⁻³²

Among Turkish hazelnut (*C. avellana* L.) genotypes ('Tombul', Palaz', 'Çakıldak', 'Sivri', 'Kalınkara', 'Ham' and 'Sivri Yağlı') used in this study, 'Tombul', 'Palaz' and 'Çakıldak' categorized as standard and most grown in Turkey. However, the cultivation of 'Tombul' and 'Palaz' decreased because they start to leaf early in the spring, while the cultivation of the 'Çakıldak' has increased because it leaves late and is, therefore, more resistant to spring frosts. 'Sivri' and 'Kalınkara', which are standard genotypes, are generally preferred as pollinators due to their lower nut quality. Local genotypes, 'Ham' and 'Sivri Yağlı', are available only as pollinators in the orchard. Within the 18 cultivars grown in Turkey, the 'Tombul' is considered the first-quality hazelnut due to its high oil content, distinctive taste, and aroma, and easily and quickly removable brown skin during roasting.³³ Since it, beings, the most consumed cultivar both nationally and internationally, 'Tombul' hazelnut leaves, hard shells, and green leafy covers have been studied for their taxane contents.^{6,19} In addition to 'Tombul', the present work is aimed to investigate the taxane content of other hazels grown in Turkey, in terms of genotype and hazelnut parts for the first time. Thus, taxane contents of 7 different hazelnuts ('Çakıldak', 'Sivri', 'Tombul', 'Palaz', and 'Kalınkara' as standard and 'Ham', 'Sivri Yağlı' local genotypes) parts (GLCs, HSs, SKs, BSs, and BSKs) were analyzed by HPLC. Results were discussed from the perspective of functional food and a promising resource for the pharmaceutical industry.

2. Materials and methods

2.1. Plant materials and chemicals

In this study, green leafy covers-GLCs and hard shells-HSs (non-edible parts), skinless kernels-SKs, brown skins-BSs, and brown-skinned kernels-BSKs (edible parts) of 'Çakıldak', 'Sivri', 'Tombul', 'Palaz', and 'Kalınkara' as standard and 'Ham', and 'Sivri Yağlı' as local hazelnut (*Corylus avellana L.*) genotypes were used. 'Tombul', 'Palaz', and 'Çakıldak' were selected as they were the most grown in Turkey. The others are preferred as pollinators and they have not been studied yet. Hazelnut samples were harvested in August 2017 in Fatsa district of Ordu province (40°59'27.9"N 37°35'01.6"E), Turkey. The samples were kept in cloth bags at room temperature until analyses were carried out.

HPLC grade of acetonitrile (>99.93%), methanol (99.99%), and hexane for organic trace analysis UniSolv® were obtained from Merck (Darmstadt, Germany). All the standard reference compounds; 10-DAB III (>95.0%), BAC III (>95.0%), cephalomannine (>97.0%), and paclitaxel (>95.0%) were acquired from Sigma Aldrich (Germany).

2.2. Taxane extraction

Oil and methanol extractions were adapted from Oguzkan et al.²¹ and performed as follows. Five parts (GLCs, HSs, BSKs, SKs, and BSs,) of 7 different Turkish hazelnuts were used as experimental material. HSs, BSKs SKs, and BSs were separately grounded by the Sinbo SCM 2934 coffee grinder while GLCs were grounded by the Loyka LKD 100 Sample Grinder Mill. GLCs, HSs, BSKs, and SKs were ground to a size of 10 g while BSs were 0.12 g. Then these parts were defatted with hexane in volumetric beakers at room temperature for 1 h using a magnetic stirrer with a heater (25°C, 400 rpm). Grounded GLCs, HSs, BSKs, and SKs were defatted with 20.0 mL hexane (1:20, w/v). Since the amount of ground BSs was small, the oil was removed with 10 mL of hexane (1:10, w/v). Solid-liquid extraction was performed by mixing the grounded samples with 99% methanol on a magnetic stirrer with a heater (25°C, 400 rpm). The defatted GLCs, HSs, BSKs, and SKs of hazelnuts were extracted by 100.0 mL methanol (1:10, w/v) while the defatted BSs were extracted by 10.0 mL methanol (1:10, w/v) at room temperature for 24 h. The resulting slurries were filtered through filter papers into the Erlenmeyer flasks. The solvent was then removed under vacuum at 40°C at 300 rpm using a rotary evaporator (HeidolphTM Hei-VapTM). The residues were dissolved with 3.0 mL (BSs with 2.0 mL) methanol, and samples were stored at - 20°C in a freezer between analyses.

2.3. Sample preparation and HPLC-photo diode array (PDA) analysis of taxane diterpenoids In this study, 10-DAB-III, BAC-III, cephalomannine, and paclitaxel, were used as standards for taxane analysis. All of the samples were filtered and brought to the condition to be analyzed by HPLC. Analyses were performed on Shimadzu CBM-20A/CBM-20A lite brand HPLC. OVA Spher 100 Phenyl-Hexyl C18 column (4.6 mm x 250 mm, 5 μm particle size) was used as a column. Mobile phase acetonitrile:water gradient system was adjusted. PDA was used as a detector.

Identification of taxane diterpenoids in hazelnut extracts was performed according to Oguzkan et al.²¹ The samples were filtered through a 0.20 µm membrane filter into the new amber glass vials (1.5 mL) before being measured in HPLC. The resultant supernatants were pooled and analyzed. The taxane contents in the extracts of hazelnut were quantified by HPLC reverse phase (RP) system (The Shimadzu CBM-20A/CBM-20A lite) having NOVA Spher 100 Phenyl-Hexyl C18 column (4.6 mm x 250 mm, 5 µm particle size). 10-DAB III, BAC III, cephalomannine, and paclitaxel in hazelnut extracts were eluted with a linear gradient of acetonitrile and water (25:75-15 min, 75-25-40 min) at a flow rate of 1.0 mL/min for 55 min. The gradient method was preferred to separate impurities from plant-derived products and proportional changes were made in the gradient method during the analysis period. Acetonitrile and the water phase were preferred because the reverse phase column is very effective in the separation of impurities. Depending on the polarity of the mobile phase, sharp and clear peaks of the sought compounds were obtained with the gradient method.

The injection volume was 20.0μ L and peak values were detected at 228 nm using a UV detector (PDA, Germany). Samples were injected into HPLC at 20.0μ L. Identification of taxanes was carried out by comparison of keeping times with standards. All analysis was performed in triplicate.

Calibration curves for 10-DAB III, BAC III, cephalomannine, and paclitaxel standards were prepared at different concentrations (50 mg/L; 25 mg/L; 10 mg/L; 5 mg/L; 2,5 mg/L and 1 mg/L). Calibration curves were prepared by plotting the peak area against the different standard concentrations. In the study for each hazelnut fraction, the standard solutions were first read and the standard curve was drawn. These different concentrations were analyzed by HPLC and retention times of 10-DAB III, BAC III, cephalomannine, and paclitaxel were

determined. Chromatograms were monitored at 11-12 min for 10-DAB III,16-17 min for BAC III, 25-26 min for cephalomannine, and 26-27 min for paclitaxel and were given in Figure 1.³³

All environmental and personal safety precautions were taken during the experimental and analysis procedures. Protective equipment such as gloves, goggles, and aprons was used during these procedures. In the evaporation process, the pressure setting was controlled from the barometer.

2.4. Statistical analysis

All the tests were performed with three replicates. The results for each taxane of seven genotypes and their parts are presented as the mean \pm SD (Standard Deviation, n=3) by Microsoft Office Excel, 2016 Version16.71. Data

were analyzed by one-way analysis of variance (ANOVA) by using Jamovi Version 2.3. Five parts from seven hazelnuts were performed for each taxane. Statistical significance in mean taxane concentrations between genotypes and their parts was evaluated by one-way ANOVA followed by Tukey and Bonferroni correction as a post *hoc*-test for multiple comparisons in our analysis. These helped ensure the reliability of our statistical inferences. A *p* value less than 0.01 was considered statistically significant. The graphs were generated by Microsoft Office Excel, 2016 Version16.71. The bars of graphics were calculated as \pm standard deviation of the mean for each and total taxanes.

3. Results

Herewith we report anticancer taxane contents of 5 standards and 2 local Turkish hazelnut genotypes both in their non-edible and edible parts for the first time. In addition to the total taxane contents of hazelnut genotypes (Figure 2), 10-DAB III, BAC III, cephalomannine, and paclitaxel contents were given separately for each hazelnut parts (GLC, HS, SK, BS, and BSK) in Figure 3-7.

3.1. Total taxanes patterns in parts of different hazelnut genotypes

The distribution of total taxanes in hazelnut parts (GLC, HS, SK, BS, and BSK) for seven genotypes is given in Figure 2. Although total taxane profiles of hazelnut genotypes were found to be in a wide range (0.21 and 11.51 μ g/g) in tested materials, average taxane contents of the hazelnut parts showed the following trend: BSK (6.08 μ g/g) > HS (5.66 μ g/g) > BS (5.07 μ g/g) > GLC (2.05 μ g/g) > SK (1.19 μ g/g). The genotypes were ranked according to their total taxane contents of all parts as follows; 'Ham' (29.78 μ g/g), Sivri Yağh (29.06 μ g/g), Palaz (24.95 μ g/g), Sivri (18.45 μ g/g), 'Tombul' (15.01 μ g/g), Kalınkara (14.96 μ g/g), 'Çakıldak' (8.09 μ g/g). **3.2. 10-DAB III, BAC III, cephalomannine, paclitaxel, and total taxane contents of hazelnut green leafy cover**

10-DAB III, BAC III, cephalomannine, paclitaxel, and total taxane contents in the green leafy cover of seven genotypes are given in Figure 3. HPLC results showed that extracts of GLCs mainly contained paclitaxel. The total amount of paclitaxel (7.46 μ g/g) comprised 52 % of total GLC's taxanes. Although this compound is substantially represented by Palaz, paclitaxel is found in nearly all tested genotypes. 10-DAB III was the second most abundant taxane and comprised 23 % of total GLC's taxanes. On the other hand, cephalomannine is not detected and/or in small quantities for GLC extracts.

3.3. 10-deacetylbaccatin III, baccatin III, cephalomannine, paclitaxel, and total taxane content of hazelnut hard shell

10-DAB III, BAC III, cephalomannine, paclitaxel, and total taxane contents in hard shells of seven genotypes are illustrated in Figure 4. In the results obtained, the total 10-DAB III (31,06 μ g/g), an important intermediate for Taxol® comprised 78 % of total HS's taxanes. BAC III was the second most abundant taxane and comprised 14 % of total HS's taxane. Hard shells of seven genotypes nearly contained all taxanes in their extracts and are the most advantageous parts for these molecules after BSKs.

3.4. 10-DAB III, BAC III, cephalomannine, paclitaxel, and total taxane contents of hazelnut skinless kernel

10-DAB III, BAC III, cephalomannine, paclitaxel, and total taxane contents in skinless hazelnut kernels of seven genotypes are given in Figure 5. HPLC results showed that extracts of SKs mainly contained BAC III. The total amount of BAC III (5.67 μ g/g) comprised 68 % of total SK's taxanes. Although this compound is substantially represented by Palaz, BAC III is found in all tested genotypes. Cephalomannine was the second most abundant taxane and comprised 31 % of total SK's taxanes. On the other hand, 10-DAB III and paclitaxel are not detected and/or in small quantities for SK extracts.

3.5. 10-DAB III, BAC III, cephalomannine, paclitaxel, and total taxane contents of hazelnut brown skin

10-DAB III, BAC III, cephalomannine, paclitaxel, and total taxane contents in kernel's brown skins of seven genotypes are illustrated in Figure 6. In the results obtained, the taxane compound distribution of BS extracts showed parallelism with SKs. BAC III and cephalomannine contents comprised 68 % and 32 % of total BSs taxanes, respectively. 10-DAB III and paclitaxel were not detected in BS extracts for any tested hazelnut genotypes. However, BS extracts accumulated ~4-fold higher content than SKs for both BAC III and cephalomannine.

3.6. 10-DAB III, BAC III, cephalomannine, paclitaxel, and total taxane content of hazelnut brownskinned kernels

10-DAB III, BAC III, cephalomannine, paclitaxel, and total taxane contents in brown-skinned kernels of seven varieties are given in Figure 7. Our results show that BSK extracts showed significant differences from BS and SK extracts in distribution and quantity for taxane compounds. The BSK extracts represented the hazelnut part with the highest abundance of taxanes. While, BS and SK accumulated only BAC III and cephalomannine, nearly all taxanes were identified in BSK extracts. According to our results, extracts of BSK mainly contained cephalomannine (42%). BAC III, the main compound of BS and SK, was the second most abundant molecule in

BSKs (32%). Despite 10-DAB III and paclitaxel molecules were not identified in both SK and BS, BSK extracts contained these taxanes at 13% and 14%, respectively.

4. Discussion

Hazelnut kernels are a nutritious and health-promoting food because of their high content of sanitary lipids, vitamins, essential amino acids, dietary fibers, and specialized phytochemicals that own respectable biological activities. The health benefits of hazelnuts have taken a new significance after they were determined to produce anticancer taxanes.^{1,34-36} Although antioxidant and phenolic compounds mainly represent phytochemicals in kernels, findings of paclitaxel and other taxanes also in them, generate a distinguished aspect of health-promoting properties of this nut.^{1,17-19,37} Together with kernel bioactivity studies, by-products (green leafy cover, hard shells, and brown skin) have been forwarded to significant research areas for phytochemicals.^{35,36} The health benefits of hazelnuts expanded after they were determined to produce anticancer taxanes. Previous works concentrated on the recovery of paclitaxel from by-products of a few hazelnuts ('Tombul', 'Ventimiglia', 'Genova', 'Alessandria', 'Gasaway'), and little is known about whether non-edible and edible parts contained taxanes in Turkish cultivars.^{17-19,37}

This is the first report to analyze seven Turkish genotypes separately in terms of paclitaxel and its derivatives and evidenced taxane compounds reveal a high variability depending on the non-edible and edible parts of genotypes. This variability was particularly noticeable in terms of the total taxane content of the genotypes. While 'Palaz' was detected as the most advantageous genotype for GLC (2.3 times higher than average), SK (3.7 times higher than average), and BSK (1.8 times higher than average) extracts, 'Ham' (2 times higher than average) and 'Sivri Yağlı' (2.2 times higher than average) had highest total taxanes of their HSs and BSs, respectively. According to the highest taxane contents, parts of 'Sivri', 'Kalınkara', and 'Çakıldak' were ranked as BSK, BS, and HS. Our analysis showed 'Ham's HS has the highest total taxane content in all tested parts of genotypes. While the total taxane amount was significantly high in GLC and SK parts of 'Palaz', there was no difference when compared with 'Sivri' for BSK. Hard shells (HS) of Ham had the highest total taxane content, however, in BS's there was no significant difference between with 'Sivri Yağlı'. Moreover, 'Tombul', the most known hazelnut genotype in paclitaxel recovery studies, had relatively lower amounts of these compounds (Figure 2).

Since Hoffman et al.¹⁷ discovered paclitaxel in *C. avellana*, HSs have become a prominent material for the investigation of paclitaxel and other taxanes. Hoffman and Shahidi¹⁹ showed HSs contain more taxanes than GLC extracts. Oguzkan et al.²¹ analyzed a mix of HS samples collected from different regions of Turkey and they identified paclitaxel and its precursor BAC III. In the same study, green leafy covers were also investigated, and BAC III was to be the main taxane. Our analysis indicated that the highest level of total 10-DAB III content among all tested parts was found in the hard shells of the hazelnuts. According to our data, GLC extracts contain the highest paclitaxel level in all tested parts. Furthermore, our analysis identifies 'Palaz' as the richest genotype in paclitaxel content among the others. Interestingly, the GLC extracts from the 'Tombul' are similar to the findings of a previous study conducted by Hoffman and Shahidi¹⁹, which proposed that the GLC included only BAC III. These findings highlight the differences in the amount of paclitaxel present in various genotypes and their parts. Taxane contents are affected by multiple variables such as extraction methods, genotype, and growing conditions.³⁷ Although, these by-products are readily available natural resources for paclitaxel and its derivatives, because of the production of HSs and GLCs in thousands of tons each year.

Despite the kernels not being analyzed in detail, the current study indicated the presence of taxanes in the skinless kernel, its brown skin, and brown-skinned kernels of Turkish hazelnut genotypes.^{19,37} Among the analyzed hazelnut extracts, the brown-skinned kernel was the most advantageous part for the synthesis of taxane compounds. Together with the diversity present in the genotypes, BSKs contain almost all taxane compounds, and the second most abundant paclitaxel, which is mainly represented by the 'Ham' genotype. Previous research indicated that BSK extracts were unfavorable for 'Tombul' and there is not any knowledge of 'Ham' for the recovery of taxane compounds. The SK extracts were found to have relatively lower taxane production compared to other parts.¹⁹ Previous research emphasized that SK has a general trend showing a lower bioactive potential that is mostly contributed by phenolic compounds¹. According to our results, skins of the kernels were discovered as the most accumulative part of the kernel for BAC III, an essential precursor compound for the semi-synthesis of Taxol® and a potent anticancer agent cephalomannine. This finding has significant implications for the fields of phytochemical and pharmaceutical research of hazelnuts. It highlights the significance that hazelnut skins are to the bioavailability of these beneficial anticancer compounds and, provides the possible relevance of this natural source in the production of pharmaceuticals.³² It is known that taxanes such as paclitaxel, which exhibit certain cytotoxicity, prevent cancer cells from proliferating and inhibit the rate at which middle- and late-stage cells transform, are found in hazelnut shells, green leafy cover, and leaves.⁵ In

addition, recent studies that investigated the activity of cephalomannine in cell viability considered that this compound is a promising natural agent for treating different cancer types.³² It has also been proven that cephalomannine significantly attenuates hepatocellular carcinoma progression.²⁸ Li and Parry³⁸ monitored the antioxidant activities and antiproliferative effects of Turkish and Oregon hazelnut parts and demonstrated that extracts from roasted hazelnut skins suppressed the proliferation of cancer cells and exhibited free radical scavenging properties. Consuming the kernel with brown skin is strongly suggested for preventing the loss of bioactive compounds from the kernel, although there is no unanimous opinion about the kernel and its brown skin relationships in terms of bioactivity.^{1,38,39}

This is the first report that revealed the differences in taxane contents of Turkish standard and local hazelnut genotypes in different edible and non-edible kernel parts. The data reveals that there are significant differences between genotype and hazelnut parts. Previous studies have demonstrated that the synthesis of taxane compounds in these plants can exhibit significant heterogeneity depending on conditions such as land compositions, height, climate, genotype, and tissue type. Turkey with a wide-ranging collection of hazelnut genotypes and a leading position in global production has promising potential for the recovery of these compounds and a rich source of this functional food with an anticancer perspective.

5. Study limitations

In this study, 4 anticancer taxane compounds were analyzed separately in 5 different kernel parts of 7 Turkish hazelnut genotypes and were presented with comprehensive and detailed data for future research. However, in addition to the hazelnut parts analyzed in this study, enriching the results by analyzing leaf and stem samples, in the same way, may contribute to the overall assessment of taxane contents in Turkish hazelnut varieties.

6. Conclusions

Turkey is a leading global hazelnut producer, resulting in being one of the countries with the highest amounts of by-products such as green leafy cover and hard shells. Identification of taxanes in hazelnut by-products which are commonly considered as discarded material and produced thousands of tomes per year is a future alternative for obtaining these compounds from readily available natural resources. Moreover, our findings challenged previous knowledge about the recovery of taxanes from edible parts of hazelnuts. While HSs, BSKs, and BSs were rich in taxanes in all of the analyzed hazelnut genotypes, SKs, and GLCs remain limited for these anticancer taxanes. Surprisingly, brown-skinned kernels of local genotypes were found rich in taxanes than skinless kernels of standard genotypes. Our study has revealed significant findings on the abundance of taxane compounds in Turkish hazelnuts including previously untested genotypes and their parts. These findings are expected to highlight the health benefits of consumption of a certain amount of the kernels of 'Ham', 'Palaz', 'Kalınkara', 'Sivri', and 'Tombul' hazelnuts containing the paclitaxel compound can prevent the irregular proliferation of body cells. Collectively, the data presented in this study can be used not only in breeding projects aiming at choosing cultivars with higher levels of health-promoting compounds but also to expand the pharmaceutical potential of local genotypes with further studies based on using a holistic approach.

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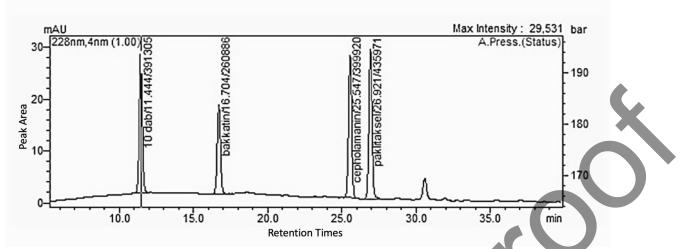


Figure 1. The chromatogram shows the peak areas and retention times of the taxanes. 10-DAB III 11.444 min and peak area 391.305, BAC-III 16.704 min and peak area 260.886, Cephalomannine 25,547 min and peak area 399.920 Paclitaxel 26.921 min and peak area 435.971.

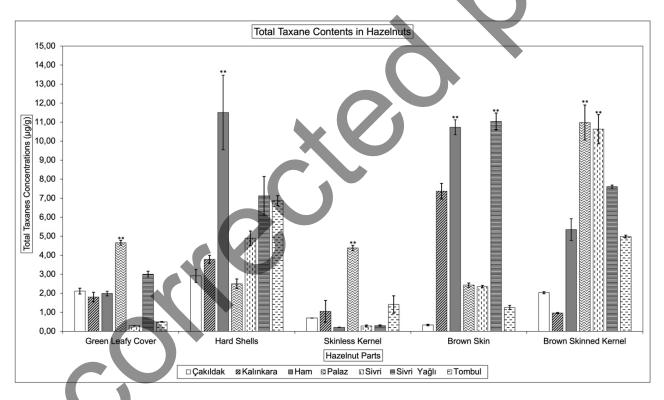


Figure 2. Total taxane contents in GLC, HS, SK, BS, and BSK of seven hazelnut genotypes. Statistical significance between hazelnut genotypes and their parts was determined using a one-way analysis of variance (ANOVA). ** indicate the p-value less than 0.01, demonstrating significant differences in the total taxane concentrations among the hazelnut genotypes within each part. Bars represent the mean ± SD.

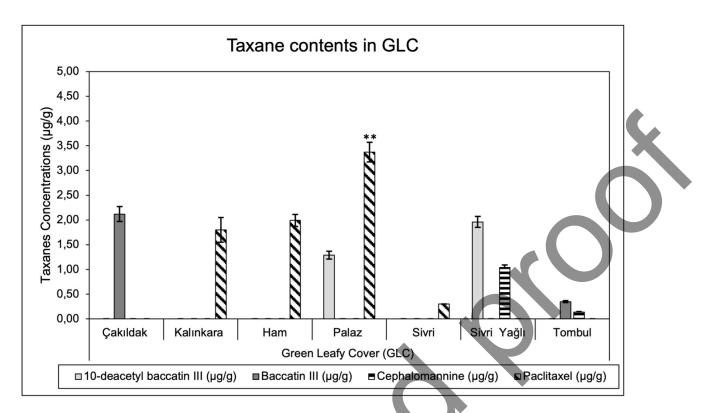
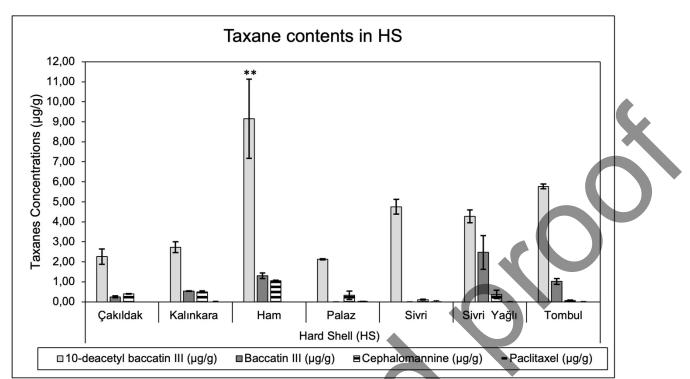
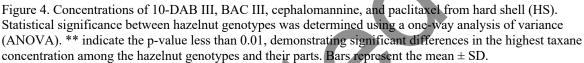


Figure 3. Concentrations of 10-DAB III, BAC III, cephalomannine, and paclitaxel from green leafy cover (GLC). Statistical significance between hazelnut genotypes was determined using a one-way analysis of variance (ANOVA). ** indicate the p-value less than 0.01, demonstrating significant differences in the highest taxane concentration among the hazelnut genotypes and their parts. Bars represent the mean \pm SD.





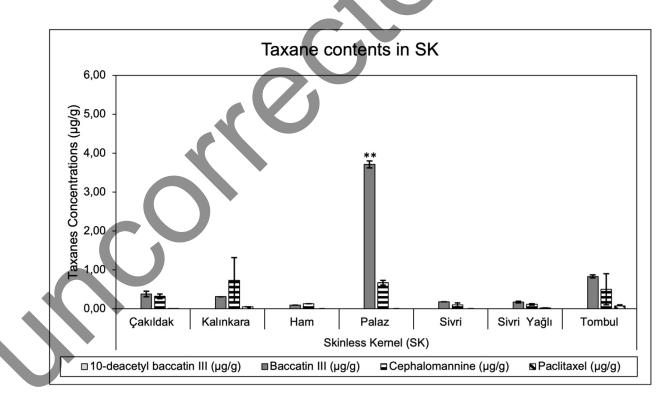


Figure 5. Concentrations of 10-DAB III, BAC III, cephalomannine, and paclitaxel from the skinless kernel (SK). Statistical significance between hazelnut genotypes was determined using a one-way analysis of variance (ANOVA). ** indicate the p-value less than 0.01, demonstrating significant differences in the highest taxane concentration among the hazelnut genotypes and their parts. Bars represent the mean ± SD.

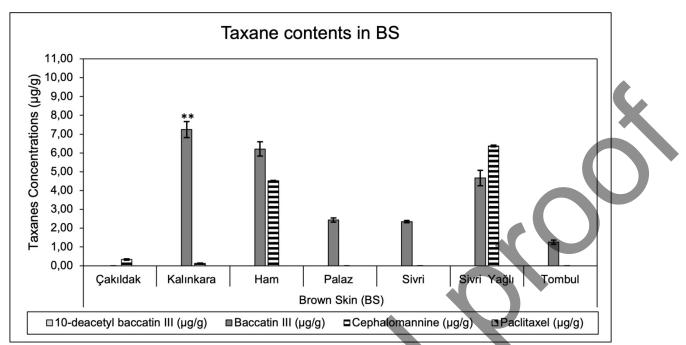


Figure 6. Concentrations of 10-DAB III, BAC III, cephalomannine, and paclitaxel from brown skin (BS). Statistical significance between hazelnut genotypes was determined using a one-way analysis of variance (ANOVA). ** indicate the p-value less than 0.01, demonstrating significant differences in the highest taxane concentration among the hazelnut genotypes and their parts. Bars represent the mean \pm SD.

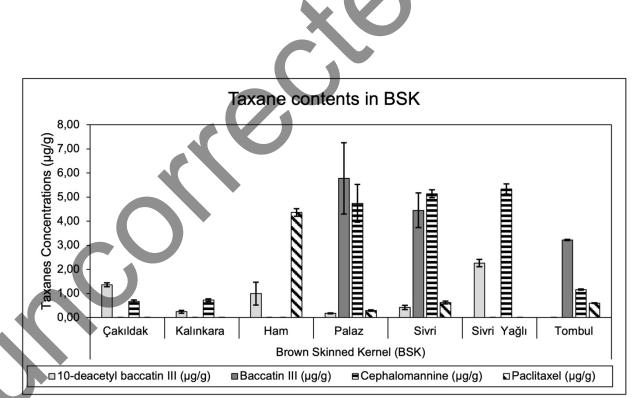


Figure 7. Concentrations of 10-DAB III, BAC III, cephalomannine, and paclitaxel from the brown-skinned kernel (BSK). Statistical significance between hazelnut genotypes was determined using a one-way analysis of variance (ANOVA). Bars represent the mean \pm SD.

Table 1. Concentrations of 10-DAB III, BAC III, cephalomannine, paclitaxel, and total taxanes from GLC, HS, SK, BS, and BSK of seven hazelnut genotypes. (Tombul, Palaz, Çakıldak, Sivri, Kalınkara, Ham and Sivri Yağlı). Abbreviations are as follows: ND, not detected; No peak is detected. All concentrations of taxanes are presented as mean \pm SD (μ g/g dry weight).

Hazelnut parts	Hazelnut varieties	10-deacetyl baccatin III (μg/g)	Baccatin III (µg/g)	Cephalomannine (µg/g)	Paclitaxel (µg/g)	Total (µg/g)
Green Leafy Cover (GLC)	Çakıldak	ND	2,12 ± 0,15	ND	ND	$2,12 \pm 0,15$
	Kalınkara	ND	ND	ND	$1,80 \pm 0,25$	$1,80 \pm 0,25$
	Ham	ND	ND	ND	1,99 ± 0,12	1,99 ± 0,12
	Palaz	$1,\!29 \pm 0,\!08$	ND	ND	3,37 ±0,20	4,66 ± 0,13
	Sivri	ND	ND	ND	$0,30 \pm 0,00$	$0,\!30 \pm 0,\!00$
	Sivri Yağlı	$1,96 \pm 0,11$	ND	$1,04 \pm 0,05$	ND	$3{,}00\pm0{,}16$
	Tombul	ND	$0,\!35\pm0,\!02$	$0,\!14\pm0,\!01$	ND	$0{,}50\pm0{,}02$
Hard Shell (HS)	Çakıldak	$2,\!26\pm0,\!38$	$0,25 \pm 0,05$	$0,\!40\pm0,\!02$	ND	$2{,}91\pm0{,}34$
	Kalınkara	$2{,}73\pm0{,}27$	0,54 ± 0,01	$0,50 \pm 0,05$	0,02 ± 0,02	3,78 ± 0,21
	Ham	$9,15 \pm 1,98$	1,31 ± 0,14	$1,05 \pm 0,03$	ND	$11,51 \pm 1,96$
	Palaz	$2,13 \pm 0,03$	ND	0,33 ± 0,21	$0,04 \pm 0,00$	$2{,}50\pm0{,}25$
	Sivri	$4,\!75\pm0,\!37$	ND	0,11 ± 0,02	$0,03 \pm 0,03$	$4,\!89\pm0,\!38$
	Sivri Yağlı	$4,\!27\pm0,\!32$	$2,\!47 \pm 0,\!84$	$0,37 \pm 0,21$	$0,\!02\pm0,\!01$	$7,\!12\pm1,\!02$
	Tombul	$5{,}77\pm0{,}12$	$1,\!02\pm0,\!15$	$0,07 \pm 0,03$	$0,\!02\pm0,\!00$	$6{,}87 \pm 0{,}26$
Brown Skin (BS)	Çakıldak	ND	ND	0,33 ± 0,04	ND	$0{,}33\pm0{,}04$
	Kalınkara	ND	$7,24 \pm 0,43$	$0,13 \pm 0,01$	ND	$7,\!37 \pm 0,\!41$
	Ham	ND	6,21 ± 0,38	$\textbf{4,52} \pm \textbf{0,02}$	ND	$10{,}73\pm0{,}39$
	Palaz	ND	$2,43 \pm 0,11$	ND	ND	$2,\!43 \pm 0,\!11$
	Sivri	ND	$2,35 \pm 0,06$	ND	ND	$2,\!35\pm0,\!06$
	Sivri Yağlı	ND	$4,67 \pm 0,41$	$6{,}37 \pm 0{,}05$	ND	$11,\!04\pm0,\!44$
	Tombul	ND	$1,\!25\pm0,\!11$	ND	ND	$1,\!25 \pm 0,\!11$
Brown Skinned Kernel Skinless Kernel (SK) (BSK)	Çakıldak	ND	$0{,}38\ \pm 0{,}07$	$0,\!32\pm0,\!06$	0,00	$0{,}70\pm0{,}03$
	Kalınkara	ND	0,31 ± 0,00	$0,73 \pm 0,58$	$0,\!05\pm0,\!00$	$1,05 \pm 0,57$
	Ham	ND	$0,\!09\pm0,\!00$	0,13 ± 0,00	ND	0,21 ± 0,01
	Palaz	ND	3,71 ± 0,09	$0,\!67 \pm 0,\!06$	ND	$4{,}38\pm0{,}14$
	Sivri	ND	$0{,}18\ \pm 0{,}00$	$0,\!10 \pm 0,\!05$	ND	$0{,}28\pm0{,}05$
	Sivri Yağlı	ND	$0,\!17\ \pm 0,\!02$	0,11 ± 0,02	$0,02 \pm 0,00$	$0{,}30\pm0{,}04$
	Tombul	ND	$0,\!83\pm0,\!04$	$0,50 \pm 0,40$	$0,\!07\pm0,\!03$	$1,\!41 \pm 0,\!46$
	Çakıldak	$1,36 \pm 0,08$	ND	$0,\!67 \pm 0,\!06$	ND	$2{,}03\pm0{,}05$
	Kalınkara	$0,\!24 \pm 0,\!06$	ND	$0,72 \pm 0,05$	ND	$0,\!96\pm0,\!02$
	Ham	$0,\!99 \pm 0,\!47$	ND	ND	4,36 ± 0,16	$5{,}35\pm0{,}57$
	Palaz	$0,\!17 \pm 0,\!02$	$5{,}78 \pm 1{,}48$	$4,74 \pm 0,78$	$0,\!29 \pm 0,\!03$	$10,\!98\pm0,\!92$
	Sivri	0,42 ± 0,09	$4,\!45\pm0,\!72$	$5,14 \pm 0,16$	0,62 ± 0,06	$10,\!63 \pm 0,\!77$
	Sivri Yağlı	$2,\!26\pm0,\!15$	ND	$5{,}34\pm0{,}21$	ND	$7{,}68 \pm 0{,}08$
	Tombul	ND	$3,22 \pm 0,02$	$1,16 \pm 0,02$	0,60 ± 0,01	$4,98 \pm 0,07$