

EVALUATION OF ANTIOXIDANT POTENTIAL OF *ASIMINA TRILOBA* (L.) DUNAL FLOWERS AND FRUITS

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ABSTRACT

This study aimed to test the antioxidant parameters of the flower and fruit ethanol extracts of 15 genotypes of *Asimina triloba* (L.) Dunal. (AzTr-01 – AzTr-15) collected in the M.M. Gryshko National Botanical Garden (Kyiv, Ukraine). Antioxidant activity of the extracts of the investigated genotypes was measured with free radical scavenging activity (FRSA) by the DPPH method and molybdenum reducing power (MRP) expressed as mg TE (Trolox equivalent) per g of DW (dry weight). The total polyphenol content (TPC) was expressed as mg GAE (gallic acid equivalent) per g of DW, total flavonoid content (TFC) was expressed as mg QE (quercetin equivalent) per g of DW, and total phenolic acid content (TPAC) expressed as mg CAE (caffeic acid equivalent) per g of DW. It was determined that the flower and fruit extracts of *A. triloba* demonstrated 5.45-77 and 2.33-7.21 mg TE/g of FRSA, respectively. Depending on a genotype, MRP of the flower and fruit extracts was 85.13-114.65 and 98.19-261.33 mg TE/g, respectively. TPC of the flower extracts ranged from 40.63 to 55.47 mg GAE/g and fruit extracts from 22.59 to 38.13 mg GAE/g. The flower extracts demonstrated 22.54-36.18 mg QE/g of TFC and 9.31-16.24 mg CAE/g of TPAC. The fruit extracts exhibited 15.07-30.55 mg QE/g and 3.11-6.58 mg CAE/g. These results indicated that various plant parts of *A. triloba* genotypes are appropriate for usage in food and pharmaceutical products as a raw material with a high content of biologically active compounds.

Keywords: Pawpaw, genotypes, antioxidant activity, phenolic compounds

INTRODUCTION

Recently, numerous studies have highlighted the growing interest in the search for new plant species, especially neglected, non-traditional, and underutilized plant species that are an important source of biologically active compounds due to their beneficial role (Klymenko *et al.*, 2019a, b; Szot *et al.*, 2020; Levon and Klymenko, 2021; Zhurba *et al.*, 2021; Mendelova *et al.*, 2022; Ľorbová *et al.*, 2023). *Asimina triloba* (L.) Dunal (pawpaw, paw paw, paw-paw, common pawpaw) native to eastern North America and Canada belongs to the family Annonaceae Juss. (Layne, 1996; Brannan and Coyle, 2021; Tułowicki, 2021). *Asimina triloba* is extensively cultivated in 26 states, in addition, this species is widely distributed in Asia, including in Korea, China, and Japan (Brannan *et al.*, 2015; Nam *et al.*, 2017; Brannan and Coyle, 2021).

The fruits of *Asimina triloba* are a valuable source of vitamins (particularly riboflavin and niacin) and mineral elements such as potassium, magnesium, iron, copper, and manganese (Galli *et al.*, 2007), as well as high polyphenolic content (Kobayashi *et al.*, 2008; Harris and Brannan, 2009; Brindza *et al.*, 2019), and essential amino acids, fatty acids (Wood and Peterson, 1999), annonacin (Tran *et al.*, 2021), and polysaccharides (Arifkhodzhaev *et al.*, 1985) are studied by many authors.

Extracts from different parts of the plant (roots, twigs, leaves, fruit, and seeds) have potent anticancer properties due to the presence of acetogenins (Gu *et al.*, 1999; Brannan and Farag, 2009; Coothankandaswamy *et al.*, 2010; Levine *et al.*, 2015; Avula *et al.*, 2018; Nam *et al.*, 2018). The pawpaw extracts can be natural products that may be used for the prevention and treatment of gastric and cervical cancers (Nam *et al.*, 2021).

Bullatacin, asimicin, and trilobacin represent the most potent and major bioactive acetogenins that are found in extract twigs from the paw paw tree (Hui *et al.*, 1989; Zhao *et al.*, 1992).

Fruits of *Asimina triloba* are very fragrant and may be used in cosmetics and skin products (Layne, 1996; Brannan *et al.*, 2012).

Biologically active compounds in *A. triloba* are presented in many plant parts, such as roots, bark, twigs, leaves, flowers, and seeds (Hui *et al.*, 1989; Zhao *et al.*, 1992, 1994; Alali *et al.*, 1999; Goodrich *et al.*, 2006; Cuendet *et al.*, 2008; Farag, 2009; Pande and Akoh, 2010; Nam *et al.*, 2021). Acetogenins, located in roots,

twigs, flowers, and seeds of *A. triloba*, and essential oil obtained from leaves (Alali *et al.*, 1999; Farag, 2009), are strong inhibitors of cancer cells (Ratnayake *et al.*, 1992; Woo *et al.*, 1995; Ko *et al.*, 2011; Sica *et al.*, 2016). The extracts from fruit, leaf, bark, and twig are used as insecticidal against insects (Rupprecht *et al.*, 1986; Ratnayake *et al.*, 1992; Zhao *et al.*, 1994; Gu *et al.*, 1999).

The stem bark extracts of *Asimina triloba* (Annonaceae) have three further acetogenins: asimicin, asimicin, and asimicin, and potent antitumor and pesticidal effects (Zhao *et al.*, 1994). Three new biologically active annonaceous acetogenins were isolated from the ethanol extract of *Asimina triloba* seeds: asimilobin, cis-murisolinone, and trans-murisolinone (Woo *et al.*, 1995). Asimicin was the first acetogenin isolated from the seeds and stem bark of *Asimina triloba* (Rupprecht *et al.*, 1986).

Considering the above review, it should be noted that the investigation of different parts of plants for the presence of antioxidant agents is very important for the development of new pharmaceutical and food products. In this regard, the study of the antioxidant activity of non-traditional fruit plants is essential for searching for new plant raw materials rich in phenolics, flavonoids, or phenolic acids.

This study aimed to evaluate antioxidant parameters of *A. triloba* flowers and fruits, and to compare the less studied flowers of *A. triloba* genotypes in the context of biologically active compounds.

MATERIAL AND METHODS

Biological material

The flowers (Figure 1) and fruits (Figure 2) of 15 genotypes of *Asimina triloba* (AzTr-01 – AzTr-15) collected in the M.M. Gryshko National Botanical Garden were the objects of these investigations. Flowers were collected during the period of full flowering. The fruits were harvested at full ripeness.

The research was conducted in 2023 in the NBG, which is located on the right bank of the Dnipro River in the southeastern part of Kyiv on the low Pechersky slopes of the Kyiv Highlands (197 m above sea level) in the Zvirynets tract (50°27'N.; 30°31' E.). The climate is moderately continental, the winter is mild, the summer is warm. The average monthly temperature in January is -3.8°C, in July +21.5°C.

The average annual temperature is +7.7 °C. The average annual precipitation is about 640 mm; fall throughout the year.



Figure 1 Flowers of *Asimina triloba* (L.) Dunal.



Figure 2 Fruits of *Asimina triloba* (L.) Dunal.

Chemicals

All the chemicals used were of analytical grade and were purchased from Sigma-Aldrich (St. Louis, MO, USA) and CentralChem (Slovakia).

Preparation of the sample extracts

Each 0.25 g sample was extracted with 20 mL of 80% ethanol for 2 hours using a GFL 3005 laboratory shaker (GFL, Burgwedel, Germany). Following extraction, the samples were centrifuged at 4605 RCF for 10 minutes (Rotofix 32 A, Hettich, Germany). The resulting supernatant was subsequently utilized to assess FRSA (free radical scavenging activity) via the DPPH assay, MRP (antioxidant capacity) using the phosphomolybdenum method, as well as to determine other antioxidant parameters, including total polyphenol, flavonoid, and phenolic acid contents.

Free radical scavenging activity – DPPH method

The antiradical activity, or free radical scavenging activity (FRSA), of the samples was determined according to the method of [Sánchez-Moreno et al. \(1998\)](#) using 2,2-diphenyl-1-picrylhydrazyl (DPPH). For the assay, 0.4 mL of each sample was mixed with 3.6 mL of a DPPH solution (prepared by dissolving 0.025 g DPPH in 100 mL ethanol). After incubation, the absorbance was measured at 515 nm using a Jenway 6405 UV/Vis spectrophotometer (England). Trolox (6-hydroxy-2,5,7,8-tetramethylchroman-2-carboxylic acid) was used as the calibration standard within a concentration range of 10–100 mg/L ($R^2 = 0.989$). Antioxidant activity was expressed in terms of Trolox equivalents per gram of dry matter (mg/g DM).

Molybdenum reducing power

The molybdenum reducing power (MRP) of the samples was assessed using the method described by [Prieto et al. \(1999\)](#), with minor modifications. A mixture containing 1 mL of the sample, 2.8 mL of 0.1 M monopotassium phosphate, 6 mL of 1 M sulfuric acid, 0.4 mL of 0.1 M ammonium heptamolybdate, and 0.8 mL of distilled water was incubated at 90 °C for 120 minutes, followed by cooling to room temperature. Absorbance at 700 nm was measured using a Jenway 6405 UV/Vis spectrophotometer (England). Trolox (concentrations ranging from 10–1000 mg/L; $R^2 = 0.998$) was used as the reference standard, and the results were expressed as Trolox equivalents (mg/g DM).

Total polyphenol content

The total polyphenol content (TPC) was measured by the method of [Singleton and Rossi \(1965\)](#) using the Folin-Ciocalteu reagent. A quantity of 0.1 mL of each sample was mixed with 0.1 mL of the Folin-Ciocalteu reagent, 1 mL of 20% (w/v) sodium carbonate, and 8.8 mL of distilled water. After 30 min in darkness, the absorbance at 700 nm was measured with the spectrophotometer Jenway (6405 UV/Vis, England). Gallic acid (25–300 mg/L; $R^2 = 0.998$) was used as the standard. The results were expressed in mg/g DW gallic acid equivalent.

Total flavonoid content

The total flavonoid content (TFC) was measured using a modified procedure based on the method of [Shafii et al. \(2017\)](#). A 0.5 mL sample was combined with 0.1 mL of a 10% (w/v) ethanolic aluminum chloride solution, 0.1 mL of 1 M potassium acetate, and 4.3 mL of distilled water. The mixture was allowed to stand in the dark for 30 minutes, after which the absorbance was recorded at 415 nm with a Jenway 6405 UV/Vis spectrophotometer (England). Quercetin, with concentrations

ranging from 1–400 mg/L ($R^2 = 0.9977$), was used as the standard. Results were expressed as quercetin equivalents (mg/g DW).

Total phenolic acid content

The total phenolic acid content (TPAC) was quantified using the method described by [Árvay et al. \(2017\)](#). A 0.5 mL aliquot of the sample was combined with 0.5 mL of 0.5 M hydrochloric acid, 0.5 mL of Arnov's reagent, 0.5 mL of 1 M sodium hydroxide, and 0.5 mL of water. After mixing, the absorbance was measured at 490 nm using a Jenway 6405 UV/Vis spectrophotometer (England). Caffeic acid, with a concentration range of 1–200 mg/L ($R^2 = 0.999$), was used as the standard. The results were expressed as caffeic acid equivalents (mg/g DW).

Statistical analysis

The results are presented as the mean values of three replicates $\pm$ standard deviation (SD). The data were analyzed using the ANOVA test, and mean differences were assessed using the Tukey-Kramer test ($p < 0.05$).

RESULTS AND DISCUSSION

As far as we are aware, this is the first report that investigates the antioxidant potential of *Asimina triloba* flowers and contrasts it with the activity observed in its fruits.

Many authors studied flowers of various plant species as a part of the human diet for their beneficial health effects ([Takahashi et al., 2020](#); [Janarny et al., 2021](#)). More than 180 species of flowers have been used worldwide as dietary supplements in the form of tea, wine, food ingredients for cakes, smoothies, or even vegetables ([Lu et al., 2016](#)).

The flowers are good sources of phenolic acids, flavanols, flavones, flavanones, and anthocyanins ([Zheng et al., 2019](#)) as well as carotenoids, alkaloids, and lignans ([Lu et al., 2016](#)). Research has indicated that the antioxidant activity of certain edible flowers is either comparable to or exceeds that of some vegetables and fruits, based on experimental findings ([Cavauiolo et al., 2013](#)). Flower extracts of some plant species have high antioxidant activity ([Pires et al., 2018](#)), anti-inflammatory properties ([Lee et al., 2018](#)), anticancer properties ([Solomon et al., 2016](#)), anti-diabetic properties ([Nambirajan et al., 2018](#)), anti-Alzheimer's effect ([Bekir et al., 2016](#)), hepatoprotective effects ([Singhal and Gupta, 2012](#)), cardioprotective ([Ittagi et al., 2014](#)), gastroprotective ([Guedes et al., 2002](#)), antimicrobial activity ([Najda et al., 2021](#)).

DPPH is a fast, straightforward, and cost-effective technique for assessing the antioxidant capacity of foods. It utilizes the free radical 2,2-Diphenyl-1-picrylhydrazyl, commonly employed to evaluate the potential of compounds to function as free radical scavengers or hydrogen donors, and to measure their antioxidant activity ([Shekhar and Anju, 2014](#)).

TEAC (Trolox equivalent antioxidant capacity) by DPPH and phosphomolybdenum method was used in this study to evaluate the antioxidant potential of the investigated plant extracts.

We determined that the mean antioxidant activity of the ethanol extracts of the flower and fruits of *Asimina triloba* measured in the DPPH method was 6.59 and 4.96 mg TE/g DW (Figure 3). This parameter for flowers and fruits varieties was from 5.45 (AzTr-11) to 7.77 (AzTr-09) mg TE/g DW and from 2.33 (AzTr-07) to 7.21 (AzTr-06) mg TE/g DW, respectively.

The ethanol extracts of *Asimina triloba* flowers exhibited molybdenum reducing power of extracts from 85.13 (AzTr-04) to 114.65 (AzTr-09) mg TE/g DW, fruits from 98.19 (AzTr-04) to 261.33 (AzTr-06) mg TE/g DW (Figure 4).

Comparison of the antioxidant activity values of fruits *A. triloba* with other species by the DPPH method showed some similarities or differences.

[Horčínová Sedláčková et al. \(2022\)](#) demonstrated in the berries of *M. aquifolium* the antioxidant activity determined by the Trolox method in the methanolic extracts (76.2 mg TE.g⁻¹ DW), in the ethanolic extracts (54.3 mg TE.g⁻¹ DW), in acetone extracts (63.4 mg TE.g⁻¹ DW) and aqueous extracts (35.5 mg TE.g⁻¹ DW). The results for antioxidant activity (μMol Trolox/g) of *Cornus mas* berries determined by the DPPH method varied from 5.94 to 16.56, ABTS method varied from 13.560 to 33.96, FRAP method varied from 8.45 to 22.49 ([Klymenko et al., 2019b](#)).

Polyphenols, as key phytochemicals found in different plant parts, are extensively studied for their use in both traditional and modern medicine. They play a significant role in the development of new medicinal products and are recognized for their wide range of health benefits ([Yang et al., 2000](#); [Bose et al., 2008](#); [Visioli et al., 2011](#); [Khurana et al., 2013](#); [Mrduljaš et al., 2017](#); [García-Conesa and Larossa, 2020](#)). Moreover, these compounds exhibit low toxicity to the human body and are essential for maintaining the oxidative stability of foods. Their inclusion in food and pharmaceutical products has significantly enhanced their quality ([Rasouli et al., 2017](#)).

The total content of polyphenol compounds in our study was for *Asimina triloba* flowers of 40.63 (AzTr-04) to 55.47 (AzTr-06) mg GAE/g DW, for fruits from 22.59 (AzTr-04) to 38.13 (AzTr-06) mg GAE /g DW (Figure 5).

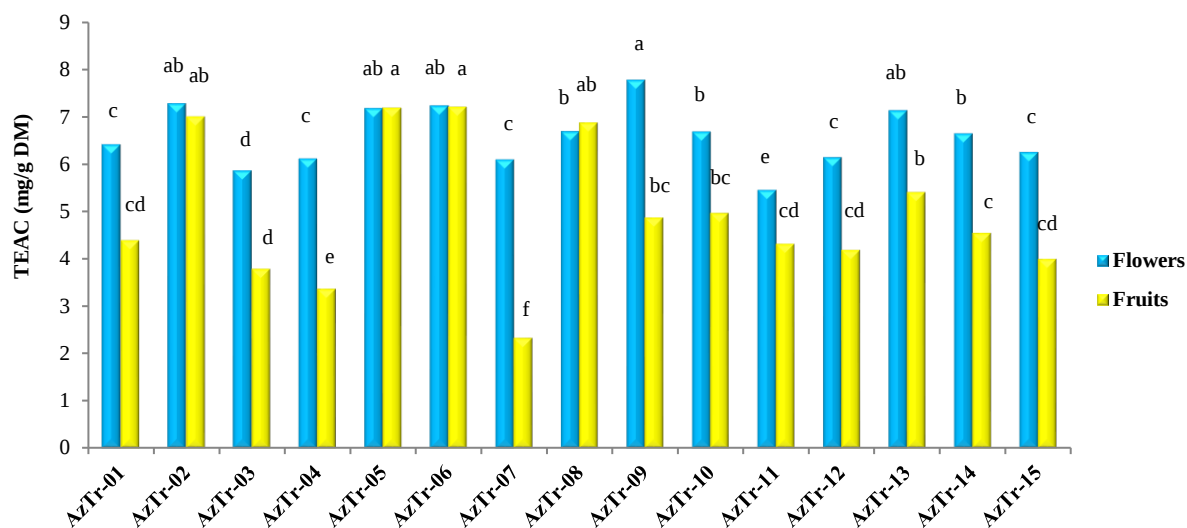


Figure 3 Free radical scavenging activity of *Asimina triloba* (L.) Dunal. flowers and fruits evaluated by the DPPH method (different superscripts in each column indicate the significant differences in the mean at $p < 0.05$); TEAC – Trolox equivalent antioxidant capacity

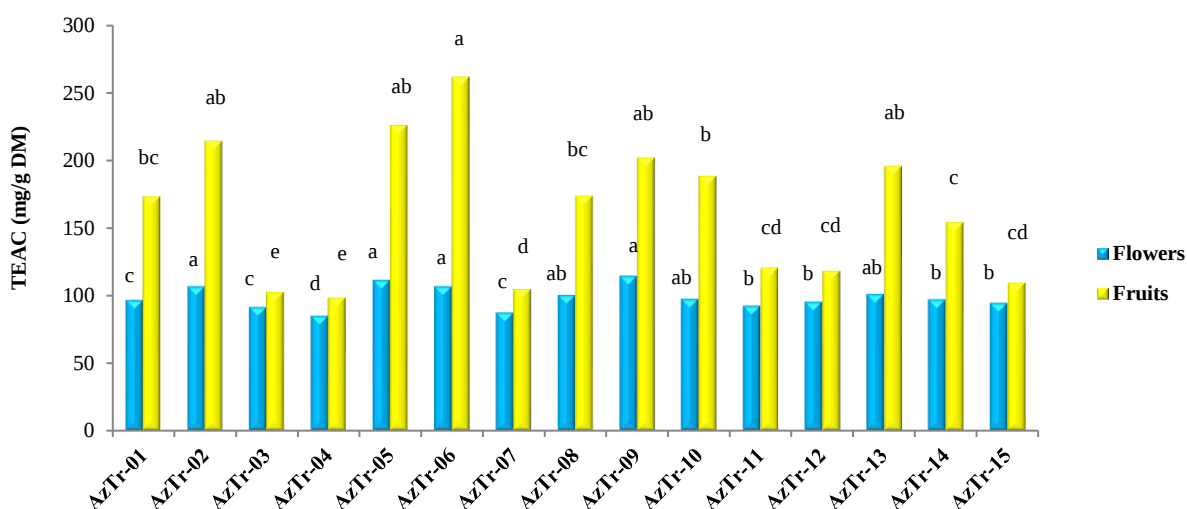


Figure 4 Molybdenum reducing power of *Asimina triloba* (L.) Dunal. flowers and fruit extracts (different superscripts in each column indicate the significant differences in the mean at $p < 0.05$); TEAC – Trolox equivalent antioxidant capacity

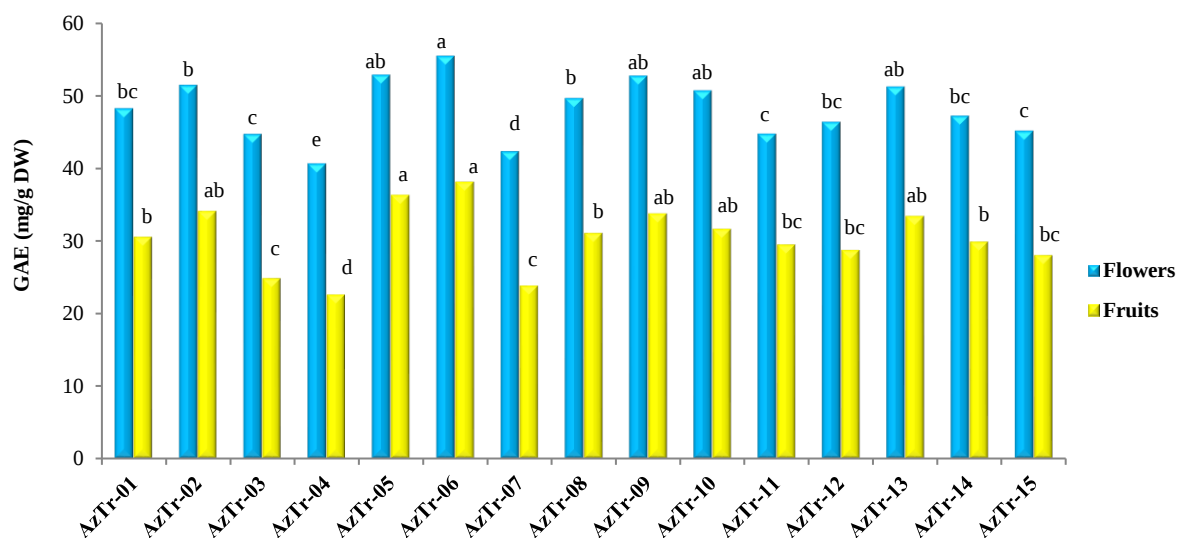


Figure 5 Total polyphenol content in *Asimina triloba* (L.) Dunal. flowers and fruits (different superscripts in each column indicate the significant differences in the mean at $p < 0.05$); GAE – gallic acid equivalent

Regarding the total phenolic content in *Crataegus oxyacantha*, the values for flowers varied between 2364.1 and 3179.6 mg GAE/100 g DW, while for fruits, they ranged from 1912.1 to 4339.0 mg GAE/100 g DW (Issaadi et al., 2020). Edwards et al. (2012) reported the total phenols quantified in the flowers of different *Crataegus* species: in *C. azarolus* values ranged from 46 to 1014 mg/100 g DW. As regards TFC, the average content in flowers was almost two-fold higher than in fruits (660.7 and 367.1 mg QE/100 g DW, respectively) (Issaadi et al., 2020).

Flavonoids are a group of bioactive compounds that are extensively found in foodstuffs of plant origin (Prasad et al., 2009). Flavonoids are classified into subgroups based on their chemical structure: flavanones, flavones, flavonols, flavan-3-ols, anthocyanins, and isoflavones. The growing body of scientific

evidence indicates that flavonoids play a beneficial role in the prevention of many diseases, thus a diet rich in flavonoids has beneficial effects on human health (Majewska and Czczot, 2009; Shabrova et al., 2011; Hui et al., 2013; Kozłowska and Szostak-Węgierek, 2014; Ballard and Maróstica Junior, 2019; Mondal and Rahaman, 2020).

The total content of flavonoids in the investigated ethanolic flower extracts of *Asimina triloba* was from 22.54 (AzTr-07) to 36.18 (AzTr-06) mg QE/g DW, for fruits from 15.07 (AzTr-04) to 30.55 (AzTr-06) mg QE/g DW (Figure 6).

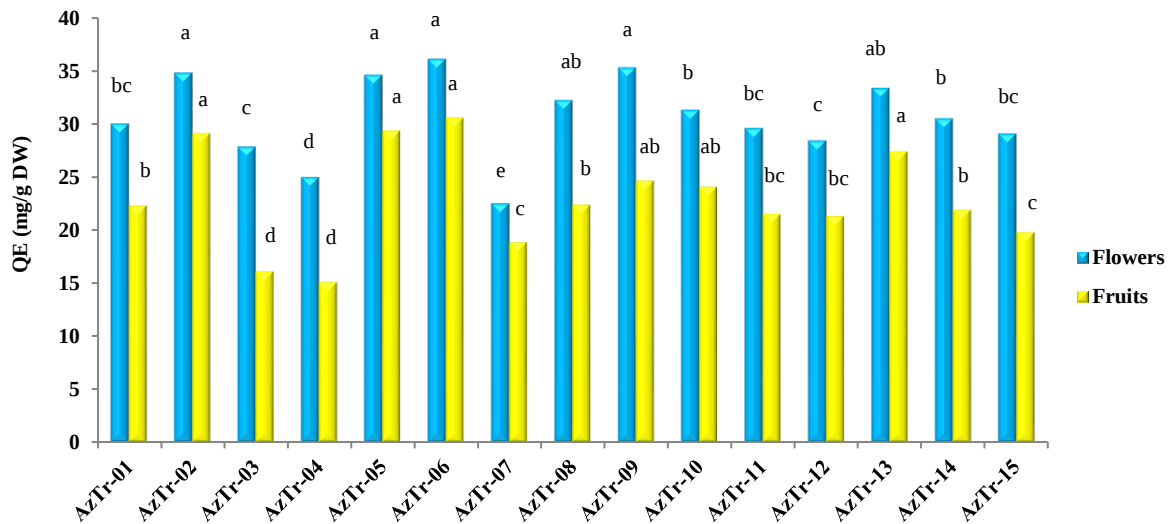


Figure 6 Total flavonoid content in *Asimina triloba* (L.) Dunal. flowers and fruit (different superscripts in each column indicate the significant differences in the mean at $p < 0.05$); QE – quercetin equivalent

The mean concentrations of flavonoids and phenolics in these five cultivars were 1.59 and 7.86 mg/g DW, respectively (Zhou et al., 2011).

The average daily intake of flavonoids for Greek and Spanish participants is 93 mg and 126.1 mg per person, respectively (Dilis and Trichopoulou, 2010; Zamora-Ros et al., 2010). In Western Europe, the daily flavonoid intake per person generally falls within the range of 100 to 1000 mg (Majewska and Czczot, 2009; Wilczyńska and Retel, 2011). In Far Eastern countries like Japan, individuals may consume as much as 2 g of flavonoids daily due to a high intake of legumes, soy, and tea (Wang et al., 2009).

Phenolic acids, as secondary metabolites, are widely distributed across the plant kingdom. These compounds contribute to the distinct taste, flavor, and health benefits of fruits and vegetables (Tomas-Barberan and Espin, 2001), enhancing

their overall quality for consumers. In recent years, there has been growing recognition of the antioxidant properties of phenolic compounds and their potential application as natural antioxidants in processed foods. Numerous studies in agriculture, biology, chemistry, and medicine have demonstrated the roles and effects of phenolic acids on human health (Cotelle, 2001; Hwang et al., 2006; Madlener et al., 2007; Ghasemzadeh and Ghasemzadeh, 2011; Godos et al., 2021).

The determination of the total phenolic acid content showed that in fruit extracts of *Asimina triloba* genotypes, they accumulated from 9.31 (AzTr-07) to 16.24 (AzTr-06) mg CAE/g DW, in flowers from 3.11 (AzTr-03) to 6.58 (AzTr-06) mg CAE/g DW (Figure 7).

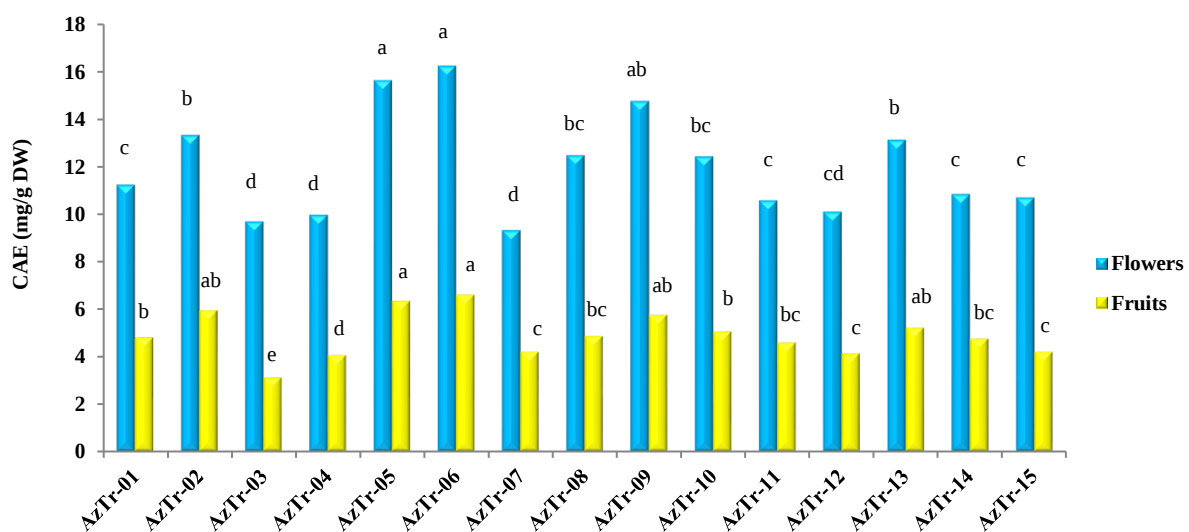


Figure 7 The total phenolic acid content in *Asimina triloba* (L.) Dunal. flowers and fruits (different superscripts in each column indicate the significant differences in the mean at $p < 0.05$); CAE – caffeic acid equivalent

We can compare our results with previous studies of *A. triloba* fruits. The TPC ranged from 22.13 to 37.36 mg GAE/g, TFC ranged from 15.10 to 32.03 mg QE/g, and phenolic acids content ranged from 0.23 to 0.76 mg CAE/g. All the tested

samples exhibited DPPH radical scavenging activities with values from 2.84 to 7.04 mg TE/g. Antioxidant activity by the molybdenum-reducing antioxidant power method ranged from 97.25 to 275.41 mg TE/g of DW (Brindza et al., 2019).

In their study, Kobayashi et al. (2008) measured the phenolic compound levels in *A. triloba* fruits at different ripening stages. The highest concentration was found in the puree of unripe fruits (123.06 mg GAE/100g FW), and the lowest in the puree of ripe fruits (98.42 mg GAE/100g FW).

Research on the phenolic profile is also valuable in lesser-known and underutilized species, which are significant for their potential in preventing various diseases.

Klymenko et al. (2019b) determined antioxidant activity in *Chaenomeles japonica* fruits by using DPPH method in the interval 10.08 to 21.42 μ Mol Trolox/g, those obtained by ABTS method varied from 31.13 to 70.87 μ Mol Trolox/g, and those obtained by the FRAP method varied from 14.97 to 42.51 μ Mol Trolox/g. The results for total phenolic content varied from 162.46 to 408.09 mg GAE/100 g. The results for the total phenolic content of *Cornus mas* fruits by the same method varied from 91.34 to 289.79 mg GAE/100g (Klymenko et al., 2019a). Horčinová

Sedláčková et al. (2021) demonstrated in the fruits of *Chaenomeles speciosa* various contents of total polyphenols profile in methanolic extract (13 214.2 mg GAE/100g DW), ethanolic extract (782.8 mg/100g DW), acetone extract (685.8 mg GAE/100g DW), and in aqueous extract (591.6 mg GAE/100g DW). Fruits contained a high amount of flavonoids 480–11 680 mg GAE/100g DW, in aqueous and methanolic extracts, respectively.

There can be considerable variation in bioactive constituents and antioxidant activities among species, varieties, cultivars, and depending on the agro-ecological conditions in which they are grown (Zhou et al., 2011; Mlynarczyk, 2020; Issaadi et al., 2020).

The results of bivariate correlation analysis for Pearson correlation coefficients are presented in Table 1.

Table 1 Results of bivariate correlation analysis for Pearson correlation coefficients

Content	Polyphenols	Phenolic acids	Flavonoids	DPPH	MRP
Flowers					
Polyphenols	1				
Phenolic acids	0.646*	1			
Flavonoids	0.516*	0.756*	1		
DPPH	0.933**	0.804**	0.803**	1	
MRP	0.847**	0.910**	0.922**	0.888**	1
Fruits					
Polyphenols	1				
Phenolic acids	0.662*	1			
Flavonoids	0.571*	0.733*	1		
DPPH	0.918**	0.810**	0.842**	1	
MRP	0.950**	0.942**	0.837**	0.861**	1

Legend: *Correlation significance at 0.05 level. **Correlation significance at 0.001 level.

The lowest correlation values between polyphenols and flavonoids were found in flowers and fruits ($r = 0.516$, $r = 0.571$, respectively) as well as between polyphenols and phenolic acids ($r = 0.646$, $r = 0.662$, respectively), but still, it can be considered moderate.

The highest value was obtained for the correlation between polyphenols and DPPH AOA ($r = 0.966$ in flowers) as well as between polyphenols and MRP AOA ($r = 0.950$ in fruits).

A strong correlation was observed between the DPPH and MRP antioxidant activities ($r = 0.888$ and $r = 0.861$, respectively, with significance at the 1% level), which is expected as both methods assess the same property, albeit using different approaches.

Furthermore, significant correlations were found between phenolic acids, flavonoids, and both AOA measurements, clearly indicating that phenolic compounds play a major role in the antioxidant activity of the analyzed flowers.

These findings indicate that the phenolic compounds in pawpaw flowers and fruits significantly influence their antioxidant capacity, as has been observed in other fruit and vegetable species (Kobayashi et al., 2008; Klymenko et al., 2019b; Brindza et al., 2019).

CONCLUSION

This study demonstrated the high antioxidant potential of raw plant material, namely flowers and fruits, of fifteen *A. triloba* genotypes. The values of antioxidant activity depended on the investigated plant extracts and genotypes of *A. triloba*. The highest values of total polyphenol, flavonoid, and phenolic acid content, and antioxidant activities were determined in all the flower ethanolic extracts, except the flower extract AzTr-08 which demonstrated a slightly lower average value of FRSA compared with fruit extract. Our results suggest that strong antioxidant activity was correlated with total polyphenol, phenolic acid, and flavonoid content by both the DPPH method ($r = 0.803$ – 0.933) and molybdenum-reducing power of extracts ($r = 0.837$ – 0.950). A strong correlation was found also between FRSA and MRP of flower and fruit extracts ($r = 0.888$ and $r = 0.861$). Therefore, *A. triloba* genotypes can be regarded to be valuable sources of the plant raw material with a high content of antioxidants of a polyphenol nature. The data obtained can be useful for further investigations and the development of new functional food products.

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Conflicts of Interest: The authors declare no conflict of interest

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