Determination of Flavonoid and Proanthocyanidin Profile of Hungarian Sour Cherry

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Abstract: Hungarian sour cherries (SC) are excellent source of anthocyanins (concentrations (100–300 mg in 100 g fresh fruit) and melatonin (0.15 mg in 100 g fresh fruit), but other flavonoid derivatives also can be isolated by aqueous alcoholic extraction. We have developed a new process for extracting non-extractable procyanidines bound to the membrane, proteins, and fibers. These compounds were separated with UHPLC-MS methods, and the structure of individual components were identified on the basis of their mass fragmentation spectra. The antioxidant capacity of soluble and non-soluble antioxidants were measured with ferric reducing antioxidant power (FRAP), 1,1-diphenyl-2-picrylhydrazyl radical scavenging activity (DPPH), trolox equivalent antioxidant capacity (TEAC) assays, and compared to the new measurement methods of water-soluble antioxidant capacity (ACW), lipid-soluble antioxidant capacity (ACL). Furthermore, total phenolic content (TPC) and total procyanidin content (PAC) were determined. As a result of our investigation, we found that the solvent combination, where in the first step is water–ethanol (1:1), then 100% ethanol were suitable for the extraction of the extractable antioxidants. However, the chemiluminescence method that is based on the elimination of the superoxide radical is more accurate than other colorimetric methods which measure antioxidant capacity.

Keywords: sour cherry; anthocyanins; extractable polyphenols; non-extractable polyphenols

1. Introduction

Sour cherry (Prunus cerasus L.) belongs to the family of Rosaceae, subfamily Prunoideae, to the genus Prunus, subgenus Cerasus. This is the hybrid that is produced by crosses between sweet cherry (Prunus avium (L.) L.) and European dwarf cherry (Prunus fruticosa) [1]. According to the FAOSTAT, world production of sour cherries in 2016 was 1,378,216 tons. The largest producers of cherries are Russia, Poland, Turkey, and the USA. Hungary is only the eighth, with nearly 70 thousand tons. Furthermore, the Hungarian cultivation of sour cherry has several centuries of history. Because of the geographic and climatic conditions of Hungary and the long breeding work, the Hungarian cultivar assortment and their varieties with outstanding nutritional parameters were developed, that are unique in the world. The popularity of the Hungarian varieties is shown by the fact that a Hungarian cultivar ‘Újfehértői fürtős’ was introduced into the United States in 1984 (in Michigan,
Utah, and Wisconsin), that is marketed under the name Balaton and is regarded as a super food [2]. In recent years, clinical trials have shown the positive physiological effects of various components that accumulate in the cherry. Different classes of flavonoids have been shown to increase the expression of the gene encoding the γ-glutamylcysteine synthetase catalytic subunit, a protein reported to be the rate-limiting step in GSH synthesis [3,4]. Cy3G has cytoprotective effects, so it protects cells such as aortic endothelial cells (EC) by reducing hydrogen peroxide (H₂O₂)-induced oxidative stress in vitro and in vivo [5–7]. Youdim et al. (2000) presented for the first time that vascular endothelial cells can incorporate anthocyanins (mainly cyanidin-3-glucoside) into the membrane and cytosol, and the incorporation of anthocyanins by the EC significantly enhanced their resistance to the damaging effects of reactive oxygen species (ROS) [8].

Moreover, flavonoids have anti-oxidative, anti-inflammatory, anti-mutagenic, and anti-carcinogenic properties coupled with their capacity to modulate key cellular enzyme function. Lipoxigenase, xanthine oxidase, and NAPDH oxidase enzymes in EC are activated during inflammatory processes. The activity of the NAPDH oxidase enzyme can cause endothelial dysfunction in two ways. The activity of the NAPDH oxidase enzyme produces oxygen-containing free radicals. The first is that the resulting superoxide anion (O₂ •−) directly affects the function of the endothelial nitric oxide synthase (eNOS) enzymes by utilizing the cofactor needed for NO synthesis. It follows that the functioning of the eNOS enzyme is disturbed. The second is that O₂ •− reacts with NO and connects to peroxinite. Peroxynitrite damages membrane proteins therefore it causes the disruption of mitochondrial electron transport chain [9].

Anthocyanins and some flavone and flavan-3-ol compounds may contribute to the prevention of hypertension [10]. Furthermore, it was demonstrated that the dietary anthocyanin Cy3G acts as a natural activator of eNOS in EC [11]. It has also been known that many extracellular agents (free fatty acids, H₂O₂, TNF-α) contribute to insulin resistance [12].

Furthermore, Guo et al. (2008) investigated the effect of Cy3G on H₂O₂- and TNF-α-induced insulin resistance on the 3T3-L1 adipocyte cell culture. Based on their measurements, it has been demonstrated that Cy3G protects adipocytes by inhibiting the kinase activity of the c-Jun NH2 terminal kinases, so the phosphorylation of the insulin receptor protein (IRS1) occurs via tyrosine [13].

The most significant antioxidant compounds in SC are phenol carboxylic acids (hydroxycinnamic acid, chlorogenic acid, neochlorogenic acid, p-coumaroylquinic acid) [14,15], flavanols (catechin, epicatechin, epigallocatechin, gallatechin), and derivatives [16,17]. The accumulation of melatonin is also significant [18,19]. Polyphenol content in sour cherries and its health effect have been widely studied, for the extractable polyphenols only. Antioxidant capacity is usually measured in food extracts with different combination of organic solvents (e.g., methanol, ethanol, acetone) and water, but these usually do not result complete extraction of antioxidant compounds. This is mainly problem in the case of phenolics compounds, because these compounds are extracted by organic solvents, which probably leaves behind significant other phenolics existing in bound form. Usually, just the amount of extractable fractions were analyzed, the non-extractable compounds (in the solid residue) were ignored and associated with cell wall matrix. Nevertheless, unextractable phenolics from plant foods and their role in health benefits have become increasingly important [14–16].

These non-extractable phenolic compounds are considered to contribute more beneficial effects (gastrointestinal health, cancer, cardiovascular disease) because after gastrointestinal digestion they remain undegraded, and are absorbed into blood plasma after being released by intestinal microflora fermentation [17,18].

Phenolics can be classified as soluble and insoluble-bound form. Thus, the polyphenols that can be extracted from foods with aqueous-organic solvent, called extractable polyphenols (EPP). However, a significant fraction of polyphenols remains in the residue after the extraction; the so-called non-extractable polyphenols (NEPP) [19].

The solid residue contains macromolecules (e.g., high-molecular-weight proanthocyanidins) and single phenolic compounds (e.g., phenolic acids, associated with macromolecules) mainly
polysaccharide constituents of dietary fibre and protein. NEPP are generally not included in polyphenol analysis, however, the NEPP content may be much higher than the EPP fraction. With regards to their chemical nature, NEPP mainly include polyphenols such as proanthocyanidins, other flavonoids, phenolic acids, and hydrolysable tannins. Accordingly, the NEPP usually were divided into two groups, hydrolysable tannins and non-extractable proanthocyanidins (NEPA) [18,20].

Acid and alkaline hydrolysis are the most common chemical methods used to extract the NEPP and recently, many other new methods such as enzymatic hydrolysis has been employed for better release of NEPP from cell wall matrices [14,21–23].

Plants contain various chemical compounds and the antioxidant effectiveness is determined by many factors (the heterogeneity and heterophasic nature of the system, the type of lipid substrate, including its physicochemical state and degree of unsaturation, the types of initiators (notably transition metals), other components, and their possible interaction).

Because of these, for assessment of antioxidant potential of endogenous compounds, single assay methods are not sufficient.

Several in vitro methods exist to measure the total antioxidant capacity. The different antioxidant assays differ in terms of assay principle and experimental conditions. Depending on what kind of reaction is involved, these assays can be classified into two groups: assays based on hydrogen atom transfer (HAT) reactions and assays based on electron transfer (ET) [24].

The most widely used procedures are FRAP (ferric reducing antioxidant power), ABTS (2,2’-Azino-bis(3-ethylbenzothiazoline-6-sulfonic acid)) or TEAC (trolox equivalent antioxidant capacity), DPPH (2,2-diphenyl-1-picrylhydrazyl), and oxygen radical absorbance capacity (ORAC). The FRAP, TEAC, and DPPH methods belong to ET methods. Unfortunately, however, they have some disadvantages and limitations. The main disadvantage of FRAP method is that the measured reducing capacity does not necessarily reflect antioxidant activity. Since the method does not include an oxidisable substrate, no information is provided on the protective properties of antioxidants [25]. DPPH assay is limited because DPPH radicals interact with other radicals (alkyl), and the time response curve to reach the steady state is not linear with different ratios of antioxidant/DPPH [26,27]. The TEAC assay also has several limitations. The ability of an antioxidant to scavenge the artificial ABTS radical may not reflect the antioxidant activity due to other mechanisms effective in complex food lipids or physiologically relevant substrates, including metal chelation and effects of antioxidant partitioning among phases of different polarities [25]. The biological and physiological functions of antioxidants are wide-ranging. The identification and quantitative determination of antioxidants, characterization of antioxidant capacity, and the evaluation of interactions between different food matrices can only be done by examining the entire antioxidant system. In order for it to be compare or examine foods with the same or different matrices, an extraction process have to develop that can efficiently obtaining the active compounds, where the componunds retain their chemical composition. Since it is important to idicificate the antioxidant compounds, and to determine the antioxidant capacity of the components as accurately as possible. The choice of the right method is also a goal because oxidative stress is well known and studied, but there is antioxidative stress too.

In our opinion, PLC technique (HAT method) can be the best choice to determine the antioxidant capacity, since this method is based on the photo-induced autoxidation inhibition of luminol by antioxidants, mediated from the radical O$_2^•−$, which can be found in human body, and is suitable to measure the radical scavenging properties of single antioxidants as well as more complex systems in the nanomolar range.

This study aims at investigating the differences between extraction processes. The first was commonly used combination of methanol, acetone, and water by Saura-Calixto and Goñi [28], and the second was an ethanol extraction that is used by the food and pharmaceutical industry too. During the experiment, not only were the extractable fractions analyzed, but also non-extractable compounds in the solid residue.
2. Results and Discussion

2.1. Extractable Antioxidant Compounds of Sour Cherry

Extracts (1/A + 1/B and 2/A + 2/B), were obtained by the two extraction methods, and analyzed with UHPLC-MS (Table 1). It can be seen that the solution combination 1 and 2 also extracted flavonoids and phenolic components. Among the flavonoids, anthocyanin-glucosides [29] and flavanol-O-glycosides [30] occur in large quantities in the sour cherry. The combination of solvent 1 has proved to be more effective mainly in the extraction of procyanidin C isomer.
Table 1. Polyphenolic compounds identified by LC-MS in Prunus cerasus L. extracts.

| No | RT [min] | Compound                                    | Chemical Formula | Exact Mass (m/z) | Fragment Ions (relative abundance, %) | Reference |
|----|----------|---------------------------------------------|------------------|-----------------|----------------------------------------|-----------|
| 1  | 10.37    | Neochlorogenic acid                         | C16H18O9         | 355.10251       | 163.0391 (100); 145.0286 (11); 135.0443 (13) | [31]      |
| 2  | 12.83    | Coumaroylquinic acid isomer ¹               | C16H18O8         | 337.09293       | 191.0553 (52); 163.0388 (100); 119.0487 (48) | [32]      |
| 3  | 12.84    | Procyanidin B isomer ¹                       | C30H26O12        | 577.13531       | 407.0771 (58); 289.0721 (60); 125.0229 (100) | [31,33]   |
| 4  | 13.45    | Coumaroylquinic acid isomer ²               | C16H18O8         | 337.09317       | 191.0555 (10); 163.0388 (100); 119.0487 (44) | [32]      |
| 5  | 13.92    | Procyanidin C isomer 1                      | C45H38O18        | 865.19922       | 407.0771 (23); 289.0729 (30); 125.0229 (100) | [34]      |
| 6  | 14.12    | Catechin ²                                  | C15H14O6         | 289.07196       | 245.0819 (34); 151.0025 (63); 109.0280 (100) | [31,33]   |
| 7  | 14.77    | Procyanidin C isomer ²                      | C45H38O18        | 865.19910       | 407.0774 (25); 289.0722 (26); 125.0230 (100) | [34]      |
| 8  | 14.94    | Chlorogenic acid ²                          | C16H18O9         | 355.10291       | 163.0390 (100); 145.0286 (11); 135.0443 (12) | [31,33]   |
| 9  | 15.21    | Feruloylquinic acid isomer ¹                | C17H20O9         | 367.10304       | 193.0498 (100); 173.0445 (7); 134.0360 (65) | [32]      |
| 10 | 15.77    | Procyanidin B isomer ²                      | C30H26O12        | 577.13519       | 407.0768 (62); 289.0719 (64); 125.0230 (100) | [31,33]   |
| 11 | 16.42    | Cyanidin-3-O-sophoroside                    | C27H30O16        | 611.16071       | 287.0552 (100); 213.0545 (4); 137.0226 (3) | [35]      |
| 12 | 16.51    | Feruloylquinic acid isomer ²                | C17H20O9         | 367.10304       | 193.0497 (100); 173.0444 (85); 134.0362 (62) | [35]      |
| 13 | 17.23    | Cyanidin-3-O-sophoroside                    | C27H30O16        | 611.16122       | 287.0552 (100); 213.0548 (3); 137.0229 (5) | [35]      |
| 14 | 17.60    | Cyanidin-3-O-glucoside ²                    | C21H20O11        | 449.10773       | 287.0552 (100); 213.0548 (3); 137.0229 (5) | [35]      |
| 15 | 17.68    | Epicatechin ²                               | C15H14O6         | 289.07175       | 245.0818 (78); 151.0382 (31); 109.0281 (100) | [31,33]   |
| 16 | 17.81    | Cyanidin-3-O-(2'-glucosyl) rutinoside       | C33H40O20        | 757.21814       | 611.1639 (4); 287.0552 (100); 213.0547 (2) | [35]      |
| 17 | 18.20    | Coumaroylquinic acid isomer ⁴              | C16H18O8         | 337.09311       | 173.0444 (100); 163.0388 (23); 119.0487 (17) | [31,33]   |
| 18 | 18.47    | Cyanidin-3-O-rutinoside                     | C27H30O15        | 595.16626       | 449.1094 (4); 287.0552 (100); 213.0551 (2) | [35]      |
| 19 | 18.54    | Cyanidin-3-O-(2'-xylosyl)-rutinoside        | C32H38O19        | 727.20795       | 581.1515 (3); 287.0553 (100); 213.0545 (2) | [35]      |
| 20 | 18.80    | Pelargonidin-3-O-(2'-glucosyl) rutinoside   | C33H40O19        | 741.22491       | 271.0602 (100) | [35]      |
| 21 | 19.41    | Cinchonain 1 isomer ¹                       | C24H20O9         | 451.10318       | 341.0666 (100); 217.0137 (33) | [35]      |
| 22 | 19.45    | Pelargonidin-3-O-rutinoside                 | C27H30O14        | 579.17096       | 433.1135 (4); 271.0603 (100); | [35]      |
Table 1. Cont.

| No | RT [min] | Compound | Chemical Formula | Measured [M + H]^+ | Measured [M − H]^− | Calculated | ∆ ppm | Fragment Ions (relative abundance, %) | Reference |
|----|---------|----------|-----------------|--------------------|--------------------|------------|-------|--------------------------------------|-----------|
| 25 | 20.05   | Peonidin-3-O-rutinoside | C28H32O15         | 609.18152          | 609.18195          |            | −0.71 | 463.1237 (4); 301.0708 (100); 286.0474 (14) | [35]    |
| 26 | 20.35   | Cyanidin-O-pentoside | C20H18O10         | 419.09750          | 419.09783          |            | −0.79 | 287.0554 (100)                  |          |
| 27 | 20.36   | Quercetin-(hexosyl)rutinoside | C33H40O15       | 771.19940          | 771.19839          | 1.31       | 300.0276 (100); 271.0248 (40); 255.0298 (19) |          |
| 28 | 20.76   | Quercetin-(hexosyl)hexoside isomer 1 | C27H30O17       | 625.14056          | 625.14048          | 0.13       | 300.0276 (100); 271.0251 (37); 255.0305 (20) |          |
| 29 | 20.96   | Quercetin-di-O-hexoside | C27H30O17         | 625.14111          | 625.14048          | 1.01       | 463.0888 (48); 301.0356 (70); 300.0277 (100) |          |
| 30 | 21.01   | Procyanidin B isomer 3 | C30H26O12         | 577.13605          | 577.13460          | 2.51       | 407.0768 (31); 289.0724 (53); 125.0229 (100) |          |
| 31 | 21.09   | Quercetin-O-rutinoside-O-glucoside | C33H40O21       | 771.19904          | 771.19839          | 0.84       | 609.1465 (89); 301.0355 (90); 300.0277 (100) |          |
| 32 | 21.44   | Naringenin chalcone-O-hexoside | C21H22O10         | 433.11380          | 433.11348          | 0.74       | 271.0613 (100); 151.0024 (59); 119.0488 (22) |          |
| 33 | 21.81   | Cinchonain I isomer 2 | C24H20O9          | 451.10233          | 451.10291          | −3.72      | 341.0667 (100); 217.0135 (36) |          |
| 34 | 22.27   | Quercetin-(hexosyl)hexoside isomer 2 | C27H30O17       | 625.14001          | 625.14048          | −0.75      | 300.0277 (100); 271.0244 (33); 255.0288 (18) |          |
| 35 | 22.81   | Cinchonain I isomer 3 | C24H20O9          | 451.10248          | 451.10291          | −0.95      | 341.0667 (100); 217.0138 (39) |          |
| 36 | 22.88   | Di-O-cafeoylquinic acid | C25H24O12         | 515.11914          | 515.11896          | 0.35       | 353.0879 (60); 191.0553 (100); 179.0339 (62) |          |
| 37 | 22.99   | Prunin | C21H22O10         | 433.11389          | 433.11348          | 0.95       | 271.0612 (100); 151.0024 (41); 119.0487 (26) |          |
| 38 | 23.35   | Isoquercitrin 2 | C21H20O12         | 463.08810          | 463.08765          | 0.97       | 301.0354 (43); 300.0276 (100); 271.0249 (37) |          |
| 39 | 23.63   | Rutin 2 | C27H30O16         | 611.16071          | 611.16122          | −0.83      | 465.1029 (3); 303.0500 (100); 85.0289 (16) |          |
| 40 | 23.74   | Dihydroxy(iso)flavone-C-glucoside | C21H20O9         | 417.11816          | 417.11856          | −0.96      | 399.1080 (33); 381.0978 (25); 297.0760 (100) |          |
| 41 | 25.34   | Astragalin | C21H20O11         | 447.09348          | 447.09274          | 1.66       | 285.0406 (66); 284.0328 (100); 255.0297 (86) |          |
| 42 | 25.47   | Nicotiflorin | C27H30O15         | 593.15094          | 593.15065          | 0.49       | 285.0406 (100); 284.0328 (73); 255.0296 (42) |          |
| 43 | 25.51   | Cinchonain I isomer 4 | C24H20O9         | 451.10400          | 451.10291          | 1.35       | 341.0674 (100); 217.0135 (48) |          |
| 44 | 25.83   | Narcissin | C28H32O16         | 623.16132          | 623.16122          | 0.16       | 315.0512 (100); 314.0435 (44); 299.0197 (40) |          |
| 45 | 27.30   | Quercetin-3-O-(4- coumaroyl)glucoside | C30H26O14       | 609.12531          | 609.12444          | 1.43       | 463.0896 (41); 300.0279 (100); 271.0247 (25) |          |
| 46 | 27.89   | Naringenin 2 | C15H12O5          | 271.06122          | 271.06065          | 0.10       | 177.0182 (17); 151.0024 (100); 119.0488 (80) |          |

1 Identified compounds in *Prunus cerasus* L. in the literature. 2 Confirmed by standard. 3 Detected only in the ethanol/water extract (1/A + 1/B). 4 Detected in both extracts (1/A + 1/B and 2/a + 2/B).
2.2. Main Anthocyanin Compounds of Sour Cherry

The main anthocyanin components in the ‘Újfehértói fürtös’ variety (Figure 1.) were the cyanidin-3-O-glucosyl-rutinoside, (2 mg/100 g), cyanidin-3-O-rutinoside, (183 mg/100 g) and cyanidin-3-O-monoglucoside (4.29 mg/100 g).

The antioxidant capacity of these components is determined by OH groups at positions C3’ and C4’ on the chalcone.

![Figure 1. UHPLC chromatogram of sour cherry at 535 nm. Confirmed by standard.](image1)

2.3. Main Flavonoid and Phenolic Compounds of Sour Cherry

Quercetin, quercetin-3 rutinoside and apigenin (flavonoids) occur in abundant quantities in sour cherry (Figure 2.). These are precursor compounds in the biosynthesis of anthocyanins. There are other phenolic compounds like chlorogenic and caffeic acid in high concentrations in this fruit. The antioxidant activity of these compounds is also high.

![Figure 2. UHPLC chromatogram of sour cherry at 340 nm. Confirmed by standard.](image2)
2.4. Total Procyanidin Content (PAC) of Sour Cherry and Sour Cherry Residues

Several studies have been reported that procyanidins have strong protective properties regarding oxidative damage, microbial infection, prevention of colon cancer, and prevention of cardiovascular disease [37–40]. However, the degree of polymerization (DP) of procyanidins may be highly influential and determine these effects [41–43]. In the gastrointestinal tract, procyanidin monomers, dimers, and trimers are absorbed into the blood system to a much larger extent than larger oligomers and polymers [44,45]. The DP value of sour cherry procyanidins is lower than 4, indicating the relatively high levels of better absorbable short-chain procyanidin species [46], so sour cherry is an exceptional source of short-chain procyanidins, and a major food ingredient.

Measurably higher amounts of mono-, di-, and trimer procyanidins were extracted with the solvent combination 1, since fewer remained in the residue 1/R, however, the total amount of procyanidins could be not extracted (Figure 3.).

![Figure 3](image_url)

**Figure 3.** The total procyanidin content of sour cherry and sour cherry residues. * Units for different type of samples: sour cherry sample: g of procyanidin A2/100 g fresh weight sour cherry; residues: g of procyanidin A2/100 g dried residue. Abbreviations: 1/R: residue of ‘solvent combination 1; 2/R: residue of ‘solvent combination 2’. * indicates significant difference (p < 0.05) from the 1/R. # indicates significant difference (p < 0.05) between the 2/R and the sour cherry.

2.5. Identification of Cinchonain I

As you can see in the Table 1, the peaks for Cinchonain I isomers are 19.41, 21.81, 22.81, and 25.51 min. with [M – H]− ions at m/z 451.10291 were identified as distereomers of Cinchonain I on the bases of their exact molecular mass, isotopic pattern, and fragmentation. The characteristic fragment ions are 451.10400, 341.06747, and 217.0155 (Figure 4.).
2.6. Extractable Antioxidant Capacity of Sour Cherry Extracts

The antioxidant activities were compared in case of solvent combination 1 and 2, using the well-known measuring methods FRAP, DPPH, TEAC, ACL, ACW, and TPC (Figure 5A.).

Our results show that the amount of antioxidant compounds extracted by these two methods is not significantly different. The aqueous–alcoholic mixture is more advantageous for the processing industry and is sufficient to extract the most important compounds of the sour cherry (anthocyanins, procyanidins, phenolic components, and flavonoids). These compounds slightly soluble in water, but ethanol is a good solvent for them. Because of the polarity of 1/A mixture, it is also suitable for extraction of low molecular weight organic acids, which provides pH 3 in which glycolized anthocyanins maintain their chemical structure (more acid is not required.). All other the antioxidants can be extracted with ethanol (1/B).

UHPLC-MS measurements show (Table 1) that combinations of 1/A or 1/B and 2/A or 2/B are only partially applicable to the extraction of bioactive compounds, because all sour cherry extracts (1/AS, 1/BS, 2/AS, 2/BS) include ‘anthocyanin and procyanidins’, ‘flavonoids’, and ‘other polyphenols’. The difference is only the amount of extracted compounds, that depends on solubility. It is therefore not surprising that methods based on the measurement of water-soluble polar components can measure a lower antioxidant concentration.

The used assays (except for ACL, ACW) are colorimetric methods based on complexometry, and have many disadvantages. The most significant drawback of the FRAP method is that it is only suitable for measuring water soluble components. In addition, not all antioxidants are able to reduce Fe$^{3+}$, antioxidants that act by H atom transfer are not detected [47–49]. During the application of the TEAC method, the problem is that the reaction of ABTS$^{+•}$ with the antioxidant compounds is strongly time-dependent, so components with ‘slow kinetic’ do not react with the radical in time [47]. DPPH is mainly used to measure the ability of polyphenols to transfer labile H atoms to radicals [50].

Obviously, the determination of antioxidant capacity of antioxidant compounds, that are extracted with solvent combination 1 and 2 by chemiluminescence technique seem to be the most appropriate. This method is suitable for selectively determining the concentration of water-soluble and lipid-soluble antioxidants. It can be used for antioxidants that act H atom transfer and electron transfer, because both groups react with the superoxide anion, which is one of the most important radicals in the living organisms [51,52].
Figure 5. ESI-MS2 spectrum of procyanidin B. (A) Comparison of the extractable antioxidant capacity of sour cherry extracts and the total phenolic content. * Units for different measurement methods: ACL: Trolox equivalent g/100 g; TEAC: Trolox equivalent g/100 g; DPPH: Trolox equivalent g/100 g; ACW: Ascorbic acid g/100 g; FRAP: Ascorbic acid g/100 g; TPC: Gallic acid g/100 g. Abbreviations: 1/AS: evaporated ethanol:water extract; 1/BS: evaporated ethanol extract after ethanol:water extraction; 1/AS + 1/BS: 1/AS and 1/BS extracts evaporated together; 2/AS: evaporated acidic methanol:water extract; 2/BS: evaporated acetone:water extract after methanol:water extraction; 2/AS + 2/BS: 2/AS and 2/BS extracts evaporated together. The table form (Supplement 1) with the results of the statistical analysis can be found in the Supplements section. (B) Comparison of the non-extractable antioxidant capacity of sour cherry residue extracts and the total phenolic content. * Units for different measurement methods: ACL: Trolox equivalent g/100 g; TEAC: Trolox equivalent g/100 g; DPPH: Trolox equivalent g/100 g; ACW: Ascorbic acid g/100 g; FRAP: Ascorbic acid g/100 g; TPC: Gallic acid g/100 g. Abbreviations: 1/R: residue of ‘solvent combination 1’; 1/RC: supernatant from the ‘extraction of hydrolysable tannins’ of 1/R; 1/RH: supernatant from the ‘extraction of condensed tannins’ of 1/R; 1/RA: supernatant from the ‘alkaline hydrolysis’ of 1/R; 1/RPR: supernatant from the ‘protease hydrolysis’ of 1/R; 1/RPE: supernatant from the ‘pectinase hydrolysis’ of 1/R; 1/RAA: supernatant from the ‘α-amylase hydrolysis’ of 1/R; 2/R: residue of ‘solvent combination 2’; 2/RC: supernatant from the ‘extraction of hydrolysable tannins’ of 2/R; 2/RH: supernatant from the ‘extraction of condensed tannins’ of 2/R; 2/RA: supernatant from the ‘alkaline hydrolysis’ of 2/R; 2/RPR: supernatant from the ‘protease hydrolysis’ of 2/R; 2/RPE: supernatant from the ‘pectinase hydrolysis’ of 2/R; 2/RAA: supernatant from the ‘α-amylase hydrolysis’ of 2/R. The table form (Supplement 2) with the results of the statistical analysis can be found in the Supplements chapter.
2.7. Capacity of Non-Extractable Antioxidants of Sour Cherry Extracts

The enzymatic digestion not resulted significant antioxidant activity (Figure 5B.). The explanation is that sour cherry does not contain large amounts of starch, so the α-amylase hydrolisis is not relevant. It does not contain large amounts of protein, so hydrolysis with protease is not important either. The use of pectinase is also more important for apple crops.

Significant amounts of antioxidant fragments were obtained in the case of the alkaline treatment, but mainly acidic hydrolysis. The results obtained using the HCl butanol extractant are not significantly different. Low molecular weight phenolic derivatives resulting from the degradation of polyphenols may be responsible for the apparently high antioxidant activity.

Proantocianines are considered as non-extractable components but some of them can be recovered with a solvent combination 1 and 2, as can be seen from the Table 1. However, non-extractable proantocianidines can not be identified in the hydrolysates.

The amount of antioxidants that extracted from the sour cherry and obtained from hydrolysis from the remaining residues were similar in case of the solvent combination (Figure 5A,B). Thus, the acidification (HCl) and the methanol–acetone–water solvent combination that was used by Saura-Calixto and Goñi [28] did not extract significantly higher amounts of antioxidant compounds. Consequently, it is not necessary to use HCl, and the methanol can be replaced by etanol in the first step, furthermore the acetone water combination can be replaced by 100% ethanol in the second step.

3. Experimental

3.1. Plant Material

Stoned, frozen sour cherry (‘Újfehértói fúrtós’ variety) was bougth from Mirelite Mirsa Zrt (Albertirsa, Hungary) in 2017. Fruit samples were frozen (−20 °C) and stored in dark.

3.2. Chemicals and Reagents

Ethanol, acetic acid, and Folin–Ciocalteu’s reagent were purchased from VWR (Randore, PA, USA). Hydrochloric acid and 1-Butanol were obtained from Merck (Damstadt, Germany). Methanol, acetone, sulfuric acid, 2, 4, 6-Tris(2-pyridyl)-s-triazine (TPTZ), 2,2-Diphenyl-1-picrylhydrazyl (DPPH), 2,2′-azino-bis(3-ethylbenzothiazoline-6-sulfonic acid) (ABTS), 4-(dimethylamino)cinnamaldehyde (DMAC), iron(III) chloride hexahydrate, sodium acetate trihydrate, potassium persulfate, procyanidin A2, sodium carbonate anhydrous, ascorbic acid, (±)-6-Hydroxy-2,5,7,8-tetramethylchromane-2-carboxylic acid (Trolox), gallic acid, sodium hydroxide, antioxidant standards, and the enzymes (protease, α-amylase) were purchased from Sigma-Aldrich (St. Louis, MO, USA). Pectinex XXL enzyme was obtained from Novozymes (Bagsværd, Denmark). Dulbecco’s phosphate-buffered saline (DPBS) were obtained from iBioTech (Szigetszentmiklós, Hungary). ACL and ACW kits were obtained from Greenlab (Budapest, Hungary). In our experiments, all the reagents were analytical grade. HPLC-grade methanol and formic acid were purchased from Fisher Scientific (Hampton, NH, USA).

3.3. Extraction of Extractable Antioxidants

The sour cherry was defrosted and homogenized. Then two different solvents were used to extract antioxidant compounds. Each extraction was performed parallel three times.

3.3.1. Extraction with the Mixture of Ethanol and Water (Solvent Combination 1)

200 g of sample was extracted with 150 mL ethanol (96%) and 150 mL distilled water for 2 h (1/A). The samples were centrifuged (Eppendorf Cetrifuge 5810R)) for 15 min at 4000 rpm and the supernatant was recovered.

150 mL ethanol (96%) was added to the residue, and the mixture was mixed for 2 h (1/B). After centrifugation (15 min, 4000 rpm), the supernatant was recovered.
For the first time, supernatants from 1/A and 1/B were evaporated together (1/AS + 1/BS), but in the second case separately (1/AS and 1/BS). Evaporation was performed at 40 °C, 10 mbar (Figure 6). The residue (1/R) was lyophilised (ScanVac CoolSafe 55-4 Pro lyophilizer), homogenized (Gorenje SMK 150 B coffee grinder), and stored in freezer (−20 °C) before use.

![Figure 6](https://example.com/figure6.png)

**Figure 6.** Scheme of the extraction of antioxidants from sour cherry. Abbreviations: 1/AS: evaporated ethanol:water extract; 1/BS: evaporated ethanol extract after ethanol:water extraction; 1/AS + 1/BS: 1/AS and 1/BS extracts evaporated together; 2/AS: evaporated acidic methanol:water extract; 2/BS: evaporated acetone:water extract after methanol:water extraction; 2/AS + 2/BS: 2/AS and 2/BS extracts evaporated together; 1/R: residue of ‘solvent combination 1’; 1/RC: supernatant from the ‘extraction of hydrolysable tannins’ of 1/R; 1/RH: supernatant from the ‘extraction of condensed tannins’ of 1/R; 1/RA: supernatant from the ‘alkaline hydrolysis’ of 1/R; 1/RPR: supernatant from the ‘protease hydrolysis’ of 1/R; 1/RPE: supernatant from the ‘pectinase hydrolysis’ of 1/R; 1/RAA: supernatant from the ‘α-amilase hydrolysis’ of 1/R; 2/R: residue of ‘solvent combination 2’; 2/RC: supernatant from the ‘extraction of hydrolysable tannins’ of 2/R; 2/RH: supernatant from the ‘extraction of condensed tannins’ of 2/R; 2/RA: supernatant from the ‘alkaline hydrolysis’ of 2/R; 2/RPR: supernatant from the ‘protease hydrolysis’ of 2/R; 2/RPE: supernatant from the ‘pectinase hydrolysis’ of 2/R; 2/RAA: supernatant from the ‘α-amilase hydrolysis’ of 2/R.

3.3.2. Extraction of Extractable Antioxidants According to Saura-Calixtoa and Goñi [28] (Solvent Combination 2)

200 g of sample was extracted with 300 mL acidic methanol/water/HCl (50:50; pH 2) were added (2/A) for 2 h, then centrifuged (15 min, 4000 rpm) and the supernatant was recovered.

300 mL acetone/water (70:30, v/v) was added to the residue, and the mixture was mixed for 2 h (2/B). After centrifugation (15 min, 4000 rpm), the supernatant was recovered [28].
The extraction was performed twice. For the first time, supernatants from 2/A and 2/B were evaporated together (2/AS + 2/BS), but in the second case separately (2/AS and 2/BS). Evaporation was performed at 40 °C, 10 mbar (Figure 6).

The residue (2/R) was lyophilised, homogenized, and stored in freezer (−20 °C) before use.

3.3.3. Preparation of Extracts from UHPLC

The purification of anthocyanins, a simple fractionation of sour cherry extracts (1/A + 1/B and 2/A + 2/B) was performed using preconditioned Supelclean ENVI-18 SPE tubes [53]. The tubes were conditioned with 5 mL MeOH then with 5 mL H₂O and finally 1 mL of fruit sample was applied. The anthocyanins were eluted with methanol:water; 80:20. Solvent was evaporated at 40 °C with Heidolph Hei-VAP Value rotary evaporator (Schwabach, Germany).

3.4. Acid Hydrolysis

3.4.1. Extraction of Hydrolysable Tannins

10–10 mg dried sour cherry residue powder from the two extractions were subjected to hydrolysis with 2 mL methanol and 200 µL sulphuric acid for 20 h at 85 °C. Samples are then centrifuged (2500 g, 10 min) and supernatants recovered. The residues was washed with 2–2 mL distilled water two times [54]. The supernatant from 1/R residue is 1/RH, and the extract from 2/R residue is 2/RH.

3.4.2. Extraction of Condensed Tannins

10–10 mg dried sour cherry residue powder from the two extractions were treated with 3 mL HCl/butanol (5:95) and 100 µL FeCl₃ (2 wt %) at 100 °C for 3 h. After centrifugation (2500 g, 10 min), the supernatant was recovered. The residues were washed with 2–2 mL HCl/butanol (5:95) two times. The supernatant from 1/R residue is 1/RC, and the extract from 2/R residue is 2/RC [55,56].

3.5. Alkaline Hydrolysis

0.1 g dried sour cherry residue powder from the two extractions were treated with 5 mL NaOH (4 mM) at 25 °C for 1 h. After centrifugation (4000 rpm, 10 min), the supernatant was recovered. The supernatant from 1/R residue is 1/RA, and the hydrolysate from 2/R residue is 2/RA [23].

3.6. Enzymatic Hydrolysis

3.6.1. Enzymatic Hydrolysis with Protease

5 mL DPBS and 10 µL tyrosine (50 mg/mL; 3550 tyrosine units/mL) was added to 0.1 g dried sour cherry residue powder from the two extractions. The samples were incubated at 60 °C for 1 h. To stop the enzymatic hydrolysis, the hydrolysates were placed in a water bath at 100 °C for 10 min. After centrifugation (5 min, 4000 rpm), the supernatant was recovered. The hydrolysate from 1/R residue is 1/RPR, and the supernatant from 2/R residue is 2/RPR.

3.6.2. Enzymatic Hydrolysis with Pectinase

5 mL DPBS and 10 µL pectinase (Pectinex XXL) was added to 0.1 g dried sour cherry residue powder from the two extractions. The samples were incubated at room temperature for 1 h. To stop the enzymatic hydrolysis, the hydrolysates were placed in a water bath at 100 °C for 5 min. After centrifugation (5 min, 4000 rpm), the supernatant was recovered. The supernatant from 1/R residue is 1/RPE, and the hydrolysate from 2/R residue is 2/RPE [22].

3.6.3. Enzymatic Hydrolysis with α-amilase

5 mL DPBS and 10 µL α-amilase (3000 Units/mL) was added to 0.1 g dried sour cherry residue powder from the two extractions. The samples were incubated at 37 °C for 10 min. To stop the
enzymatic hydrolisis, the hydrolysates were placed in a water bath at 100 °C for 20 min. After centrifugation (5 min, 4000 rpm), the supernatant was recovered. The supernatant from 1/R residue is 1/RAA, and the hydrolysate from 2/R residue is 2/RAA [23].

3.7. Determination of Total Phenolic Content (TPC)

Total phenolics in all extracts were determined with the Folin–Ciocalteu assay [57] with minor modifications. 10 µL of appropriately diluted extracts, standard gallic acid solutions (50, 100, 200, 400, 800, and 1600 µg/mL) or water (blank) was mixed with 190 µL of distilled water in a well of a 96-well plate; 25 µL of Folin–Ciocalteu reagent solution was then added. After 6 min, 75 µL of 7% Na₂CO₃ was added. The mixture was shaken gently and incubated in the pre-heated chamber (50 °C) for 10 min, and its absorbance was measured at 765 nm, using the microplate reader (SPECTROstar®Nano, BMG Labtech, Ortenberg, Germany). TPC was expressed as milligrams of gallic acid equivalents (mg GAE/100 g fresh weight sour cherry/dried residue). In some cases, the extract was too dilute, so 100 µL of sample and 100 µL of H₂O were mixed in a well.

3.8. Determination of Total Procyanidin Content (PAC)

Total procyanidin content was measured using the method of Prior et al. [58]. Dried powder of residuals and sour cherry were weighed (500 mg) into a 50 mL conical tube. 20 mL extraction solution (acetone/deionized water/acetic acid 75:24.5:0.5) was added to the samples. The samples were vortexed for 30 s followed by sonication for 1 h at room temperature. After centrifugation (4000 rpm, 10 min), the supernatant was collected for analysis.

70 µL of 96% ethanol for blank; or 70 µL of control (100 µg mL⁻¹ Procyanidin A2 in ethanol)/standard/samples were added to 210 µL DMAC solution (0.1 wt % in ethanol) in a well of 96-well plate. The mixture was shaken gently and incubated in the pre-heated chamber (25 °C). The microplate was read for 25 min.

The plate reader protocol was set to read the absorbance (640 nm) of each well in the plate every min for 30 min. The maximum absorbance readings were used for calculation.

3.9. Determination of Antioxidant Capacity

In this experiment, FRAP, DPPH, TEAC, and PCL (Photochemiluminescence assay) methods were used to measure, the antioxidant capacity.

3.9.1. FRAP

The ferric reducing antioxidant power assay was performed as previously described by Benzie and Strain [59]. It is based on the reduction of the Fe³⁺-TPTZ complex to the ferrous form at low pH. This reduction is monitored by measuring the absorption change at 593 nm.

The reaction was carried out in a microtiter plate. 30 µL of distilled water and 10 µL properly diluted samples/standard were pipetted in a well of 96-well plate, then 200 µL FRAP reagent (10 volumes of 250 mM acetate buffer (pH 3.6), one volume of 20 mM ferric chloride solution and one volume of 10 mM tripyridyl-s-triazin (TPTZ) in 40 mM HCl) were added. The mixture was incubated at 37 °C and the absorbance was taken after 8 min at 593 nm.

The FRAP values was calculated and expressed as ascorbic acid equivalents per 100 g sample (fresh weight sour cherry/dried residue).

3.9.2. DPPH

The DPPH free radical scavenging activity was measured using the method of Brand-Williams with modified as follows: 10 µL of appropriately diluted sample or Trolox solution (31.25, 62.5, 125, 250, 500, 750, and 1000 µM) and 50 µL distilled water was added to 190 µL of DPPH solution (0.1 mM in methanol) in a well of a 96-well plate. The mixture was shaken gently, incubated at 25 °C and the
absorbance was taken after 30 min. The absorbance was measured at 517 nm, using the microplate reader. DPPH was expressed as milligrams of trolox equivalents (mg TE/100 g fresh weight sour cherry/dried residue) [26,60].

3.9.3. TEAC

Determination of Trolox Equivalent Antioxidant Capacity (TEAC). This assay was performed as reported previously with slight modification. ABTS radical cations were prepared by mixing equal volumes of ABTS (7 mM in H$_2$O) and potassium persulfate (4.9 mM in H$_2$O), and the solution was left to stand in the dark for 12–16 h at room temperature; then the above solution was filtered and diluted with 80% ethanol to an absorbance of about 2 at 734 nm. 70 µL of 80% ethanol and 10 µL properly diluted samples/standard were pipetted in a well of 96-well plate, then 190 µL ABTS solution in a well of a 96-well plate, and the absorbance was recorded at 734 nm after 30 min of incubation at room temperature. Trolox was used as standard, and a standard calibration curve was obtained for Trolox at concentrations of 15.65, 31.25, 62.5, 125, 250, 500, and 1000 µM. The TEAC of samples was calculated from the standard curve of Trolox and expressed as gram of Trolox equivalents (TE) per 100 g of fresh weight sour cherry/dried residue (g TE/100 g) [61].

3.9.4. Photochemiluminescence Assay (PLC)

This assay was described by Popov and Lewin, and distributed as a complete system under the name Photochem® by Analytik Jena AG (Jena, Germany).

In the PCL assay, the photochemical generation of (O$_2$•−) free radicals is combined with the sensitive detection by using chemiluminescence. The assay is initiated by optical excitation of photosensitizer (S), resulting in the generation of the superoxide radical anion.

\[ S + h\nu + O_2 \rightarrow [S*O_2] \rightarrow S^{*+} + O_2^{•−} \]  

There are two different protocols: ACW (water-soluble antioxidant capacity) and ACL (lipid-soluble antioxidant capacity) so both of the hydrophilic and the lipophilic antioxidants can be measured separately. These are standardized conditions, so the results are comparable to other assays. The antioxidant potential was assayed by means of the lag phase (ACW) or by means of the area under the curve (ACL) at different concentrations [51,52].

A. ACL

The lipophilic antioxidants were measured with the ACL kit. The reaction solutions were prepared by mixing 2.3 mL Reagent 1 (methanol), 200 µL Reagent 2 (buffer solution), 25 µL Reagent 3 (photosensitizer and detection reagent), and 0–30 µL of Reagent 4 (calibration standard for quantification of lipophilic antioxidants in Trolox equivalents) or 10 µL of sample (beverage diluted with Reagent 1) were mixed and measured. The detector measures the current proportion to the generated luminescence as a function of measurement time. The detector signal, monitored for 180 s, Includes a lag phase in which no luminescence can be detected. When the antioxidants are exhausted, the amount of radicals in the sample increases until the detected signal reaches the maximum. The length of the
lag phase increases in function of the amount of the antioxidants in the sample, and it is calculated by determining the first derivative and the maximum point of the detected curve. The intersection point of the slope of the straight line with the x-axis defines the lag time. Results are expressed as mg equivalents of ascorbic acid per 100 g fresh weight sour cherry/dried residue [51].

3.10. UHPLC Analysis

Measurements were carried out using CromasterUltraRs UHPLC, equipped with diode array detector, automatic sampler and Agilent OpenLAB software. The sample components were separated on a Phenomenex Kinetex column (2.6μ, XB.C18, 100A, 100 × 4.6 mm).

UHPLC running conditions consisted of the following linear gradient steps

0 min solvent A 15%,
0–25 min solvent A to 30%,
25–30 min solvent A to 40%,
30–40 min solvent A to 50%.
Solvent A: MeOH; Solvent B: 3% HCOOH (Formic acid) in water.

Flow rate was 0.7 mL min⁻¹ and oven temperature was kept at 25 °C. The anthocyanin content was analyzed quantitatively by comparison with the corresponding authentic standards. UV–vis detection was used at 535 nm wavelength for anthocyanins and 340 nm for flavonoid and phenolic compounds. The appropriate amounts of sour cherry extracts were measured and dissolved in solvent A. Injection volume was 10 μL.

3.11. UHPLC-MS Analysis

The UHPLC system (Dionex Ultimate 3000RS) was coupled to a Thermo Q Exactive Orbitrap mass spectrometer (Thermo Fisher Scientific Inc., Waltham, USA) equipped with an electrospray ionization source (ESI). The HPLC separation was achieved on a Thermo Accucore C18 column (100 mm × 2.1 mm × 2.6 μm). Sampler and oven temperature were maintained at 25 °C, flow rate was 200 μL min⁻¹. Eluent A was water containing 0.1% formic acid and eluent B was methanol containing 0.1% formic acid. The following gradient elution program was used: 0 min, 95% A; 0–3 min, 95% A; 3–43 min, →0% A; 43–61 min, 0% A; 61–62 min, →95% A; 62–70 min, 95% A. 2 μL of the samples were injected in every run. The Q Exactive hybrid quadrupole-orbitrap mass spectrometer was operated with the following parameters: capillary temperature 320 °C, spray voltage 4.0 kV in positive and 3.8 kV in negative ionization mode. The resolution was set to 35,000. The mass range scanned was 150–1500 m/z. The maximum injection time was 100 ms. The resolution was set to 17,500 in the cases of MS2 scans. The collision energy was 35 NCE. Sheath gas and aux gas flow rates were 32 and 7 arb, respectively. Xcalibur 4.0 (Thermo Fisher Scientific Inc., Waltham, USA) software was used to collect and analyze data.

3.12. Statistical Analysis

Data were expressed as means ± standard errors. Data were statistically analyzed using the SPSS statistical software, version 23. (SPSS Inc, Chicago, IL, USA). One-way analysis of variance (ANOVA) with Tukey’s honestly significant difference test (homogeneity of variances) and Games–Howell significant difference test (not equal variances) were used to compare means among groups. The level of significance was set at p < 0.05.

4. Conclusions

It is known that different types of extractants can extract a variety of bioactive compounds. Accordingly, just one solvent is not sufficient to extract the total antioxidant compounds of foods, solvent combination is necessary, and the choice of suitable solvents is essential. As a result of our investigation, we found that the amounts and antioxidant properties of the dissolved compounds are not significantly different in case of solvent combination 1 and 2. However, Solvent combination 2
(methanol–acetone–water) can not be used in the food technology, only alcohol–water extraction is suitable.

The choice of the appropriate solvent is just the first step, the next is the determination of the exact antioxidant capacity. To do this you will need to find the most appropriate measurement method for the food type, because they contain different antioxidant compounds. FRAP method is good for the determination of water soluble, low molecular weight components. However, anthocyanins and associated flavonoids are only slightly soluble in water, so this method can not be measured these well. Although the solvent of DPPH and TEAC methods is ethanol, there are certain limitations that block measurement of the anthocyanins rich in sour cherry. For instance, because of the steric accessibility of DPPH\(\bullet\) radical, the DPPH method can not adequately measure anthocyanin compounds [47], and in case of TEAC, the degree and position of hydroxylation and methoxylation in the B ring of anthocyanins, affects the stability and reactivity and thereby the antioxidant capacity [62]. However, the ACL and ACW methods are suitable to determine the antioxidant activity of various chemical components in SC.

Supplementary Materials: The following are available.

Author Contributions: J.R. conceived and designed the experiments. A.N., E.S., L.S., A.B., J.R.H., M.M.S., and Z.C. performed the experiments. P.B., Z.C., and J.R. analyzed the data. A.N. and J.R. wrote the paper. All authors read and approved the final manuscript.

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Dedication: This study is dedicated to Professor Sándor Biró on the occasion of his 70th birthday, and we honor his undisputed merits in the field of the microbial genetics.

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