Healthy Benefits of Natural Pigments

Subjects: Nutrition & Dietetics Contributor: Zhengguo Li

Natural pigments, including carotenoids, flavonoids and anthocyanidins, determine the attractive color of fruits. These natural pigments are essential secondary metabolites, which play multiple roles in the whole life cycle of plants and are characterized by powerful antioxidant activity. After decades of research and development, multiple benefits of these natural pigments to human health have been explored and recognized and have shown bright application prospects in food, medicine, cosmetics and other industries.

carotenoids flavonoids anthocyanidins fruits antioxidant health benefits

1. Introduction

Eating fresh fruits is a major source for people to intake natural nutrients; beyond that, fruit can also be fermented or processed into fruit wine, fruit juice, preserved fruit and other products for consumption. Fruits contain a variety of natural pigments, which are secondary metabolites with important biological activities ^{[1][2]}. In addition, fruit color is also an important commodity quality that determines the consumers' choice. These pigments play essential roles in plant growth and development, photosynthesis, attracting pollinators and seed carriers, and resisting biotic and/or abiotic stresses ^[3]. The main pigments in fruits include carotenoids contributing red, yellow, and orange (for example, apricot and tomato); flavonoids contributing yellow (for example, citrus); anthocyanidins contributing red, purple, and blue (for example, grape and blueberry). These pigments have powerful antioxidant activities and multiple health benefits, such as delaying aging, repairing the nervous system, anti-atherogenicity, anticancer, and anti-inflammation ^{[4][5][6]}. Studies on the natural fruit pigments and their antioxidant activities and functional mechanisms are of guiding significance for targeted breeding and help improve the nutritional and commodity quality of fruits.

2. Carotenoids

Carotenoids, a class of irreplaceable micronutrients in the human diet, are widely found in various bacteria, fungi, algae, and plants. So far, more than 800 natural carotenoids have been found, with colors of red, orange, yellow, etc. ^[2]. In plants, carotenoids are a category of lipophilic natural pigment stored in chloroplast and chromoplast membranes. They make plant organs, including fruits, appear red, yellow and/or orange ^[8]. Carotenoids also exist in green tissues as photosynthetic pigments. They can capture light energy and deliver it to the chlorophyll a in the excited state, which performs the function of transforming light energy ^[9]. Moreover, carotenoids are effective active oxygen scavengers with provitamin activity and can reduce oxidative stress in the human body. They have an outstanding effect on chronic diseases, preventing cardiovascular and cerebrovascular diseases, eye diseases,

osteoporosis and cancer ^[10]. There are abundant and diverse carotenoids in fruits, as more than 100 kinds have been identified ^[8].

2.1. Chemical Structure and Classification of Carotenoids

Carotenoids are a class of tetraterpenoid fat-soluble polyene pigments, and they are C40 terpenoid compounds, and the derivatives are composed of eight isoprenoid units. The polyene skeleton is the most prominent feature of carotenoids, and most of the molecules have bilaterally symmetric and varied numbers of conjugated double bonds, generally 3–15, which confer carotenoids a strong ability of light harvesting and oxidation-reduction ^[11]. The structural characteristics enable carotenoids to be precursors and converted into a variety of active substances in the living body through various pathways ^{[8][11]}. In plants, most carotenoids exist in the form of more stable trans isomers, which have strong lipophilicity and polarity, and can be either free in cells or esterified with fatty acids or complexed with sugars and proteins, which depend on the molecular structure ^[12]. Depending on the presence or absence of the oxygen atom, carotenoids can be roughly divided into xanthophylls ($C_{40}H_{56}O_2$ or $C_{40}H_{56}O$) and carotenes in plants include α -carotene, β -carotene, γ -carotene, lycopene, phytoene, and phytofluene, and the first three are biosynthesis precursors of vitamin A ^[13]. Depending on whether there is a terminal ring, carotenoids can also be classified into cyclic and acyclic carotenoids ^[8].

2.2. Carotenoids in Fruits

Almost all fruits contain carotenoids. Some typical fruits that are rich in carotenoids are mentioned here, including goji berries (*Lycium barbarum*), tomato (*Solanum lycopersicum*), and apricot (*Prunus armeniaca*). Goji berries are recognized as a healthy food for both medicine and diet, the orange-red color of goji fruits is mainly determined by carotenoids. The dry wild goji fruits contain about 6 mg/g of carotenoids, of which zeaxanthin dipalmitate accounts for about 70–80%, which endowed the powerful ability to scavenge oxygen free radicals and active oxygen [14]. Tomatoes are important commercial crops, and the content of carotenoids is an important indicator of their quality. Tomato fruits contain at least twelve kinds of carotenoids, of which lycopene accounts for 70–90% of the total carotenoids, and β -carotene accounts for 5–26%. Lycopene confers tomato fruits with a strong antioxidant capability and high nutritional value ^[15]. Apricot fruits are also rich in carotenoids, and seven types of carotenoids have been identified in apricots. The content of *trans*- β -carotene is approximately 61.2 mg/kg (dry weight, DW), accounting for 39–65% of total carotenoids; 9-*cis*- β -carotene accounting for 4%; zeaxanthin accounting for about 3%. To sum up, β -carotene makes up the predominant proportion of total carotenoids in apricot fruits, which makes it one of the best sources of β -carotene l^[16].

2.3. Biosynthesis of Carotenoids

Carotenoids are tetraterpenoids composed of eight isoprenes. There are two independent terpenoids biosynthesis pathways in plants, namely, the mevalonic acid (MVA) pathway in cytoplasmic and the 2-*C*-Methyl-d-erythritol-4-phosphate (MEP) pathway in plastid ^[8]. Both pathways can produce isopentenyl pyrophosphate (IPP) and

dimethylpropenyl diphosphate (DMAPP), which are precursors of carotenoids biosynthesis ^{[17][18]}. Pyruvate and glyceraldehyde 3-phosphate (G3P) generate MEP under the catalysis of 1-deoxy-d-xylulose-5-phosphate (DXP) synthase (DXS) and DXP reductoisomerase (DXR) ^[19]. Additionally, DXS is the first rate-limiting enzyme, and MEP finally formed IPP and DMAPP after a series of enzymatic reactions ^[20].

Then, DMAPP and IPP (1:3) are condensed under the catalysis of geranylgeranyl pyrophosphate synthase (GGPPS) to produce a variety of direct precursors of carotenoids biosynthesis. Under the catalysis of phytoene synthase (PSY), two molecules of geranylgeranyl pyrophosphate (GGPP) undergo a condensation reaction to produce colorless linear phytoene ^[8]. PSY is the most critical rate-limiting enzyme in carotenoids biosynthesis; antisense inhibition of the tomato *SIPSY1* gene significantly reduced the carotenoid content in the fruit, resulting in yellow fruit at maturity ^[21]. Under the catalysis of phytoene dehydrogenase (PDS) and ζ -carotene dehydrogenase (ZDS), *cis*-lycopene is formed after two dehydrogenation reactions. Finally, under the catalysis of carotene isomerase (CRTISO), the conformation changes to produce pink-colored all-*trans* lycopene ^[3]. Another important point in the catalytic action of β -carotene hydroxylase (BCH) and ϵ -carotene, and α -carotene transforms into lutein under the catalytic action of β -carotene hydroxylase (BCH) and ϵ -cyclohydroxylase (ECH) ^[22]. Zeaxanthin goes through two consecutive reactions under the continuous catalysis of zeaxanthin cyclooxygenase (ZEP) and neoxanthin synthase (NSY) to synthesize violaxanthin and then further converts into neoxanthin. This is the final step of carotenoids biosynthesis ^[22].

Tomato is a model material for studying fruit ripening and metabolisms. In recent years, there have been many studies on transcriptional regulation of carotenoids' biosynthesis and metabolism in tomatoes. RIN is one of the most deeply studied MADS-box transcription factors, and the tomato *rin* mutant cannot turn red normally and contains a small amount of lycopene or even no lycopene ^[23]. The expression of the *PSY1* gene is hardly detected in the *rin* mutant, and recent studies have shown that RIN directly binds to the *PSY1* promoter to regulate carotenoids biosynthesis in fruit ^[24]. In addition, several other members of the MADS-box transcription factor family also play important roles in the regulation of carotenoids biosynthesis, including SITAGL1, SIMBP15, SIMADS1, and SIMBP8 ^{[25][26][27][28]}. Other transcription factors also participate in this process, but most of which indirectly regulate the carotenoids biosynthesis genes ^[29].

3. Flavonoids

Flavonoids belong to the polyphenol superfamily and are synthesized by plants and widely exist in stems, leaves, flowers, and fruits. Flavonoids are usually appeared yellow, light yellow, or white. Seed disseminators are attracted by the attractive color of fruits, which is prominently contributed by flavonoids. Plants produce flavonoids in response to various abiotic and biotic stresses, and flavonoids also participate in regulating plant growth and development ^[30]. There are a variety of flavonoids with rich contents in fruits, and more than 5000 different kinds have been identified from plants ^[31]. In recent years, more and more studies have reported multiple benefits of flavonoids on human health. Flavonoids have gradually become a research hotspot in biology, food science, medicine, and other fields.

3.1. Chemical Structure and Classification of Flavonoids

Flavonoids are a category of phenols with a basic C15 phenyl-benzopyrone skeleton formed by a C6-C3-C6 structure, and most contain the 2-phenylchroman core [32]. The typical flavonoid compound is a flavone, which was first identified from stems and leaves of the *Primula* plant ^[33]. The 2-phenylchroman core is constituted by two aromatic rings (A and B) linked by a central three-carbon chain that may or may not form a third ring (C). According to the connection position of B-ring (C-2, C-3 or C-4), most classes of this secondary metabolite known as flavonoids (flavones, flavonols, flavonones, and flavanols), isoflavonoids (isoflavones, isofavonones, isoflavonols, isoflavans, rotenoids, coumestans, pterocarpans, and isoflavenes) and neoflavonoids (arylcoumarins, neoflavenes, etc.), respectively [34]. Moreover, flavonoids can be divided into flavonols, flavones, flavanols (catechins), flavanones, anthocyanins, and isoflavonoids based on variations and saturation of the C-ring ^[35]. There are some special cases, including aurones, chalcones, and dihydrochalcones etc. are the primary product of flavonoids biosynthesis, which are also precursors of other classes [34]. For example, chalcone is not considered a true flavonoid due to the lack of the aromatic C-ring but is still considered a member of the flavonoid family ^[36]. On the other hand, various modifications of flavonoids are very common. Hydroxyl, methoxyl, methyl, isopentenyl, methylenedioxyl, benzyl, nitro, etc., are usually substituted on A-ring and/or B-ring. Glycosidated and methylated derivatives are the main forms of plant natural flavonoids. d-glucose, I-rhamnose, glucorhamnose, galactose, and arabinose can be formed glycosidic bonds with flavonoids at the C-3 or C-7 positions $\frac{37}{2}$.

3.2. Flavonoids in Fruits

People in different regions have different dietary patterns, which was influenced by geographical factors 37. However, no matter what kind of dietary habits, fruits play an indispensable role and are an excellent bank of dietary flavonoids. Flavonoids in berry fruits are mainly found in the peels and seeds, grape (Vitis vinifera), citrus fruit (family Rutaceae) and several tropical fruits mentioned here are typically fruits rich in flavonoids. Flavonoids have long been a mainstay of researches in grape and wine because this class of substances not only affects the color of the wine but also the aging capacity, bitterness, and astringency [38]. Anthocyanins, flavan-3-ols (tannins, generate proanthocyanidins), and flavonols are the main flavonoids found in grapes and their derivatives, and the first two kinds will be discussed in the next section. Flavonols only accumulate in the skin of grapefruit, guercetin, myricetin, and kaempferol occupy 90% of flavonols, while laricitin, isorhamnetin, and syringetin make up the remaining 10% ^[39]. Citrus fruit is an import bulk fruit all over the world, which is rich in a variety of physiologically active substances. Citrus polyphenols (including flavonoids) are mainly found in the peel, which accounts for 40-50% of the citrus fruit biomass [40]. Citrus flesh also contains many kinds of flavonoids, but the content is significantly lower than that in peels [41]. Citrus peel contains abundant glycosylated flavones and polymethoxylated flavones (PMFs) ^[42]. The glycosylated flavones in citrus include hesperidin, neoeriocitrin, poncirin, dydimin, narirutin, naringin, diosmin, and isorhoifolin ^{[43][44]}. The PMFs are a unique class of highly methoxylated flavonoids, which exist almost exclusively in citrus peels (Lu et al., 2020). Nobiletin and tangeretin are the most abundant PMFs in citrus, accounting for 25.1% and 16.9% of the total content, respectively [42][45]. As a by-product of the citrus industry, citrus peels have been widely used in the food, pharmaceutical, and cosmetic industries [41]. In addition, tropical fruits also contain abundant flavonoids. From an evaluation of fruit by-product water extract (FWE) by HPLC-DAD, passion fruit presented rutin (7.0 mg/L FWE) and quercitin (4.0 mg/L FWE). Acerola fruit only presented rutin (8.0 mg/L FWE). Additionally, mango fruit presented rutin (29.0 mg/L FWE), quercitin (4.0 mg/L FWE), and epicatechin (2.0 mg/L FWE) [46].

3.3. Biosynthesis of Flavonoids

Flavonoid biosynthesis has been widely studied and well understood. Flavonoids are produced in the cytoplasm, catalyzed by a series of enzymes that belong to shikimic acid and the phenylpropanoid pathway and the acetate pathway, and then are transported to vacuoles for storage [47][48]. Each step of flavonoids biosynthesis is catalyzed by a specific enzyme family that is conserved in higher plants [49]. Phenylalanine, at the beginning of flavonoid biosynthesis, is deaminated by phenylalanine ammonia lyase (PAL) and hydroxylated by cinnamate-4-hydroxylase (C4H). Then, it is converted to 4-coumarate ^[50]; 4-coumarate contains a carboxyl group, which could be activated by 4-coumarate-CoA ligase (4CL) to produce 4-coumarate-CoA. Then, 4-coumarate-CoA acts as a substrate for subsequent enzymes, which also makes it become a branching point for either phenolic acids or flavonoids. The synthesis of chalcone is considered to be the beginning of flavonoids' biosynthesis. Under the action of chalcone synthase (CHS), three molecules of malonyl-CoA, the product of the acetate pathway, are condensed with one molecule of 4-coumarate-CoA to form A-ring and B-ring of typical flavonoids C6-C3-C6 skeleton. Additionally, the C-ring is formed by chalcone catalyzed by chalcone isomerase (CHI), and flavanone (naringenin) is formed in this step. A branch is arisen here for the production of flavone, and a double bond is formed between C-2 and C-3 of the naringin C-ring catalyzed by flavone synthase (FNS). Returning to the trunk, naringenin is hydroxylated by flavonone-3-hydroxylase (F3H), flavonoid-3'-hydroxylase (F3'H), or flavonoid-3'-5'-hydroxylase (F3'5'H) to form dihydrokaempferol, dihydroguercetin, or dihydromyricetin, respectively. Subsequently, they are reduced by dihydroflavonol reductase (DFR) to form leucocyanidin, leucopelargonidin and leucodelphinidin, respectively. These leucoanthocyanidins are then catalyzed by flavonol synthase (FLS) to form kaempferol, quercetin, or myricetin, respectively [47][48][49][50].

There have been many studies on the regulation of flavonoid biosynthesis. R2R3-MYB transcription factors occupy the core position in the regulatory network, which could directly regulate the key biosynthesis genes ^[51]. In the transgenic *Arabidopsis thaliana* that is heterologously expressed *VvMYBF1*, the accumulation of flavonoids was significantly increased ^[52]. Another study also showed that overexpression of *VvMYBF1* in the transgenic *Arabidopsis thaliana* induced the expression of *AtPAL*, *AtCHS*, *AtCHI*, *AtF3'H*, *AtFS'H*, *AtFLS*, *AtDFR*, *AtLDOX*, and *AtUFGT* and enhanced the activity of corresponding enzymes ^[53]. In addition, basic helix-loop-helix (bHLH) and WD40 protein also participate in the regulation of flavonoids accumulation. The MYB-bHLH-WDR (MBW) protein complex efficiently regulates flavonoids accumulation ^[54]. Beyond that, other transcription factors are also involved in the regulatory network. For example, a recent study indicated that three citrus AP2/ERF subfamily members act as positive factors to promote the expression of *CitCHIL1*, which enhances the accumulation of flavanones and flavones ^[55].

4. Anthocyanidins

Given the prominent specificity and high nutritional value of anthocyanidins, they are discussed separately in this section. There are a large number of studies on anthocyanidins, which have different colors compared with other flavonoids. Anthocyanidins appear in different colors that are affected by the basic structure, co-pigmentation, metal ion complexation, and vacuole pH ^[4]. They make fruits and flowers appear in orange, red, purple, or blue and cooperate with other natural pigments to build gorgeous colors in plants. Proanthocyanidins (Pas) and anthocyanidins come from the same branch of flavonoids biosynthesis pathway, which mainly includes catechins, epicatechins, and their polymers, which are unique reddish-brown substances found in fruits ^[56]. They also determine the bitterness and astringency of the fruits and their secondary products ^[57]. So far, more than 600 different anthocyanidins have been identified and widely distributed in at least 27 families, 73 genera, and countless species ^[58].

4.1. Chemical Structure and Classification of Anthocyanidins

Anthocyanidins belong to a category of typical flavonoids, also contain the basic C6-C3-C6 skeleton. The unique color of anthocyanidins comes from their conjugated bond ^[4]. Cyanidin, pelargonidin, delphinidin, peonidin, malvidin, and petunidin are the most common anthocyanidins. Among them, cyanidin is the most abundant in fruits that accounting for about 50% ^[50]. Same as the flavonoids, the chemical diversity and bioavailability of anthocyanidins are contributed by the number and location of glucosides ^[59]. Anthocyanidins in fruit also exist mainly in the form of glycosylation ^[59]. In addition, acylated anthocyanin, caffeoylated anthocyanin, and malonylated anthocyanin ^[59].

4.2. Anthocyanidins in Fruits

Anthocyanidins are widely distributed in a variety of fruits and are abundant in fruits, making them ready to obtain. Grapes are the most common fruit that rich in anthocyanins, and there are a large number of anthocyanins in grape skins and wine. Proanthocyanidins are mainly found in peels and seeds but very little in the pulp ^[60]. The quality of wine is mainly contributed by proanthocyanidins, which determine the astringency, browning, and turbidity ^[60]. Blueberry (*Vaccinium corymbosum*) is also the most representative and well-known fruit that rich in anthocyanins, about 22 kinds of anthocyanins have been detected in blueberry ^[61]. Wang et al. ^[62] measured and analyzed the metabolic profile of 14 varieties of blueberry; five anthocyanins including cyanidin (154.5–1001 µg/g FW, fresh weight), delphinidin (3.3–69.7 µg/g FW), malvidin (3.0–55.8 µg/g FW), petunidin (5.6–51.9 µg/g FW), and peonidin (4.0–164.3 µg/g FW) were identified by HPLC-PAD. Although the content of different kinds of anthocyanins in blueberry varies greatly, in general, blueberry contains very high levels of anthocyanins. Purple kiwifruit (*Actinidia species*) contains cyanidin and delphinidin ^[63]. Peng et al. used ultra-high performance liquid chromatography (UHPLC) and liquid chromatography/mass spectrometry analysis (LC-MS) to analyze anthocyanins in different varieties of purple kiwifruit at different development stages. According to their research, cyanidin-based anthocyanins were distributed in the pericarp and flesh, and the anthocyanis in the pericarp were higher than that in the flesh at different developmental stages ^[64].

4.3. Biosynthesis of Anthocyanidins

Leucoanthocyanidins (leucocyanidin, leucopelargonidin, and leucodelphinidin), which are derived from the flavonoid biosynthesis pathway, are common precursors of anthocyanidins and flavan-3-ols ^{[48][49]}. Anthocyanidin synthase (ANS), also known as leucoanthocyanidin dioxygenase (LDOX), catalyzes three leucoanthocyanidins into cyanidin, pelargonidin, and delphinidin, respectively. As substrates, leucoanthocyanidins and anthocyanidins are reduced to (+)-catechin and (–)-epicatechin under the catalytic action of leucoanthocyanidin reductase (LNR) and anthocyanidin reductase (ANR) ^{[48][50]}. Finally, anthocyanidins undergo structural modification, including glycosylation, acylation, and methylation under the catalytic action of UDP-glucose flavonoid-3-*O*-glucosyltransferase (UFGT), acyl transferase (AT), and methyl transferase (MT), respectively ^[49].

As a kind of important flavonoid, anthocyanidins biosynthesis is also tightly regulated by the evolutionarily conserved MBW complexes ^[54]. In tomato, SIAN2-like (SIMYB114) is a major transcriptional regulator that promotes anthocyanidins biosynthesis ^[65]. Our study also presented that overexpression of *SIMYB75* (*SIAN2*) in tomato results in high-level anthocyanidins in various tissues ^[66]. In apple, a long terminal repeat (LTR) retrotransposon was found to insert the upstream of the *MdMYB1* gene and enhance its expression and then activate anthocyanidins biosynthesis ^[67].

5. Antioxidant Capacity and Health Benefits

5.1. Antioxidant Mechanisms

Reactive oxygen species (ROS) are produced by oxidation reactions in organisms ^[68]. The biochemical imbalance caused by ROS may damage many biological macromolecules, including DNA, RNA, lipid, and protein, that lead to degenerative diseases such as multiple sclerosis, cancer and Alzheimer's disease ^[69]. The characteristic structure of carotenoids is conjugated polyunsaturated double bond, and this particular chemical structure confers the ability to inhibit free radicals. The conjugated polyunsaturated double bond also confers carotenoids a lipophilic character and, thus, protects lipoproteins and cell membranes from free radical attack ^[70]. Flavonoids, including anthocyanidins, are capable of directly scavenging free radicals by donating the hydrogen atom or indirectly inhibiting free radicals by chelating free metal ions ^[71][72]. In addition, flavonoids are also closely related to the cellular oxidant-antioxidant enzyme system and inhibit some of the enzymes that produce ROS, including NADH oxidase, glutathione S-transferase, mitochondrial succinoxidase, and microsomal monooxygenase. On the other hand, flavonoids enhance the activity of antioxidant enzymes ^[71]. For example, hesperidin could enhance the activity of antioxidant enzymes (SOD), glutathione peroxidase (GPX), glutathione reductase (GRX), and catalase (CAT) ^[73], and quercetin could activate glyoxalase activity [^{74]}.

5.2. Antioxidant Capacity and Detection Methods

Different in vitro assay methods may generate different results when evaluating the antioxidant capacity of natural pigments in fruits. This is because a single analysis method cannot take into account the following factors: the diversity and complexity of the antioxidant reaction mechanism, the distribution effect of antioxidants in a heterogeneous system, and the influence of other components in the test system. Thus, multiple assays are often

used simultaneously to measure antioxidant capacity objectively ^[75]. The following assays are commonly used to evaluate the antioxidant capacity of fruit pigments: 1,1-dipheny-1-2-picrylhydrazyl (DPPH), and 2,2-azinobis (3ethylbenzothiazoline-6-sulphonic acid) (ABTS) free radical scavenging assay, total radical-trapping antioxidant parameter (TRAP), oxygen radical absorbance capacity (ORAC), and ferric reducing antioxidant power (FRAP). These reaction principles are essentially based on the ability of artificially produced free radical scavenging to evaluate the antioxidant capacity of the substance under test ^[76]. Besides, the total phenolic content (TPC) and total flavonoid content (TFC) also partly represent the antioxidant capacity ^[77].

The composition and content of natural pigments vary greatly among different fruits that caused significant differences in antioxidant capacity between different fruits, even in the same variety ^{[46][62]}. When people eat fruits or other related products, a variety of natural pigments with antioxidant properties intake, it seems more practical to study the overall antioxidant capacity. We summarize the detailed information of several common fruits on the main natural pigments and their overall antioxidant capacity in <u>Table 1</u>.

Fruit	Antioxidant Capacity	Methods	Main Pigments	References
	27.42 μmol TE/g DW	PCL		[<u>78][79]</u>
	32.37 μg DW	DPPH	Carotenoids: lycopene, lutein, α -carotene, β -carotene	
Tomato (Solanum	13.32 ± 0.79%			
lycopersicum)	10.51 ± 0.31 mg/100 g FW	TPC	quercetin	
	8.02 ± 0.37 mmol/L 100/g FW	FRAP		
Grape (Vitis	2605 ± 487 μmol TE/100 g FW	ORAC	Flavonoids: quercetin, kaempferol Anthocyanins: delphinidin, malvidin, cyanidin, petunidin, peonidin	[<u>80][81][82][83]</u> [<u>84][85]</u>
viiniciaj	42.18 mmol TE/g (seed)			
	36.40 mmol TE/g (skin)			
	70.14 ± 1.54 mg of catechin/100 g	TPC		
	281.3 μmol TE/g (seed) 12.8 μmol TE/g (skin)	TEAC		

Table 1. The antioxidant capacity and main antioxidant pigments of common fruits.

	Fruit	Antioxidant Capacity	Methods	Main Pigments	References
		2.4 μmol TE/g (flesh)			
		58.04 mol TE/100 g (seed)	FRAP		
		32 mmol Fe ²⁺ /L (juice)			
		16.8 to 92 mmol TE/g (seed)			
		15.7 to 113.3 mmol TE/g (skin)	DPPH		
		15 mmol TE/L (juice)			
	Blueberry (Vaccinium corymbosum)	4521 μmol TE/100 g FW	ORAC	Flavonoids: quercetin glycosides, myricetin glycosides Anthocyanins: cyanidin glycosides, peonidin glycosides, delphinidin glycosides, malvidin	[<u>62][86][87][88]</u>
		4669 μmol TE/100 g			
		1287 µmol Vit. C equiv./100 g FW	PSC	giycosides, peturildiri giycosides	
		128 ± 30 μmol of QE/100 g (no PBS wash) 19.0 ± 4.7 μmol of QE/100 g (PBS wash)	CAA		
		5.88 ± 1.17 μmol/g FW	ABTS		
		7775.45 ± 1009.60 EC ₅₀ —g FW/g	DPPH		
		99 ± 5.28 μg/mL (juice)			
		0.3336 ± 0.1219 μg/mL (juice)	MAO-A		
		0.1064 ± 0.03630 μg/mL (juice)	Tyrosinase		

Fruit	Antioxidant Capacity	Methods	Main Pigments	References
	1.26 ± 0.109 μg/mg (juice)	Glucosidase		
	0.453 ± 0.07156 μg/mg (juice)	DPP-4		
	13.23 ± 1.37 μmol/g FW	ABTS		
	2142.42 ± 125.64 EC ₅₀ —g FW/g	DPPH		
Blackberry	5905 μmol TE/100 g	ORAC	Flavonoids: kaempferol glycosides, quercetin	
Blackberry (Rubus fruticosus)	201.05 ± 4.69 µmol Fe ²⁺ /g DM (mature)	EDAD	glycosides, myricetin glycosides Anthocyanins: pelargonidin glycosides, cyanidin glycosides, peonidin glycosides	[<u>86][88][89][90]</u>
	241.06 ± 4.44 μmol Fe ²⁺ /g DM (fully mature)	FIXAF		
	20.35 ± 3.25 μM/g FW	TEAC		
Strawberry (Fragaria	8348 ± 888 μmol TE/100 g FW	0540	Flavonoids: kaempferol glycosides, quercetin	[<u>81][86][88][91]</u> [<u>92</u>]
ananassaj	4302 μmol TE/100 g	URAC	Anthocyanins: pelargonidin glycosides, cyanidin glycosides, neonidin glycosides	
	136 ± 18 μmol of QE/100 g (no PBS wash) 42.2 ± 3.3 μmol of QE/100 g (PBS wash)	CAA	gij ocolaoo, poormani gij ooolaoo	
	7.87 ± 0.87 μmol/g FW	ABTS		
	3778.94 ± 333.88 EC ₅₀ —g FW/g			
	417.9 mg AA/100 g FW	UFFN		
	2.15 mmol	FRAP		

Fruit	Antioxidant Capacity	Methods	Main Pigments	References
	Fe ²⁺ /100 g FW			
Cherry	5945 ± 978 μmol TE/100 g FW	ORAC		
	27.4 ± 4.1 μmol of QE/100 g (no PBS wash) 6.8 ± 0.8 μmol of QE/100 g (PBS wash)	CAA	Flavonoids: quercetin glycosides, myricetin glycosides, kaempferol	[<u>81][86][93][94</u>]
serotina)	8.83 ± 1.32 μmol/g FW		Anthocyanins: cyanidin glycosides, peonidin glycosides, pelargonidin glycosides	
	50.03 ± 1.32 mg ascorbic acid eq. 100/g FW	ABTS		
	6065.68 ± 563.46 EC ₅₀ —g FW/g	DPPH		
Peach (Prunus persica)	2235 ± 278 μmol TE/100 g FW	ORAC	Carotenoids: β-cryptoxanthin, β-carotene	[<u>81][95][96]</u>
	1586 ± 51 μmol TE/100 g FW (nectarine)	ORAC	quercetin glycosides, isorhamnetin glycosides, kaempferol glycosides Anthocyanins:	
	7.1 to 88.7 mg/serving (white-flesh nectarine)		Gyarmann grybbolaeo	
	9.1 to 40.0 mg/serving (yellow-flesh nectarine)			
	19.6 to 107.3 mg/serving (white-flesh peach)			
	13.0 to 50.5 mg/serving (yellow-flesh peach)			

Fruit	Antioxidant Capacity	Methods	Main Pigments	References
	14.4 to 105.5 mg/serving (white-flesh nectarine)	FRAP		
	15.9 to 49.5 mg/serving (yellow-flesh nectarine)			
	27.9 to 119.6 mg/serving (white-flesh peach)			
	19.0 to 72.2 mg/serving (yellow-flesh peach)			
Plum (Prunus	5661 ± 440 μmol TE/100 g FW	ORAC	Carotenoids: β-cryptoxanthin, β-carotene Flavonoids: quercetin, myricetin, eriodictyol Anthocyanins: cyanidin glycosides	[<u>81][96][97][98]</u>
salicina)	1181 to 2366 μmol TE/100 g (summer varieties)			
	1510 to 3444 μmol TE/100 g (autumn varieties)			
	33.5 ± 8.6 μmol of QE/100 g (no PBS wash) 12.9 ± 0.1 μmol of QE/100 g (PBS wash)			
	27.4 to 61.1 mg/serving	DPPH		
	203 to 730 μmol TE/100 g (summer varieties)			
	132 to 554 μmol TE/100 g			

Fruit	Antioxidant Capacity	Methods	Main Pigments	References
	(autumn varieties)			
	40.5 to 127.2 mg/serving	FRAP		
	863 to 4215 mmol AAE/100 g (summer varieties)	Haomolycic		
	2012 to 12449 mmol AAE/100 g (autumn varieties)	Haemolysis		
Apricot	21.68–69.78% inhibition	DPPH	Carotenoids: β-cryptoxanthin, lutein, β-carotene, γ- carotene. zeaxanthin Flavonoids: rutin, quercetin 3-rutinoside, kaempferol 3-	[16][99][100]
(Prunus armeniaca)	1.24 to 11.47 μmol TE/g FW	TEAC	rhamnosyl-hexoside and quercetin 3- acetyl-hexoside Anthocyanins: cyanidin 3-rutinoside, and cyanidin 3- glucoside	
Citrus fruit (family <i>Rutaceae</i>)	2887 ± 717 μmol TE/100 g FW (orange)	ORAC	Carotenoids: violaxanthin, antheraxanthin, lutein, zeaxanthin, β-cryptoxanthin, β-carotene, α-carotene Flavonoids: naringin, neohesperidin, neoeriocitrin, kaempferol, rutin, narirutin, hesperidin, nobiletin, tangeritin, quercetin, hesperitin, naringenin, tangeretin, sinensetin, diosmin, poncirin, didymin, isorhoifolin, myricetin, resveratrol	[<u>68][81][101]</u> [<u>102]</u>
	1848 ± 186 μmol TE/100 g FW (lemon),			
	1640 ± 299 μmol TE/100 g FW (grapefruit)			
	1033.8 to 1331.7 μmol TE/g (orange peel)			
	189.5 to 256.2 μmol TE/g (orange peel)	TEAC		
	0.51 to 0.68 mg/mL (orange peel)	DPPH		

Fruit	Antioxidant Capacity	Methods	Main Pigments	References	
	80.93 mg TE/100 g (lemon)				
	53.39 mg TE/100 g (lime)				
	69.03 mg TE/100 g (mandarin)				
	451.56 mg TE/100 g (lemon)) benefits. oids and
	235.38 mg TE/100 g (lime)	ABTS			ntiallergic, Inosomal,
	301.08 mg TE/100 g (mandarin)				ases such
Apple (Malus pumila)	4592 ± 201 μmol TE/100 g FW [<u>107]</u> 8.6 to 31.4 mmol	ORAC	Flavonoids: hyperoside, isoquercitrin, rutin, reynoutrin avicularin, quercitrin Anthocyanins:	(<u>105)</u> [<u>81][103][104]</u> ,	ncer to a reventing -carotene
[]	2.4 to 8.1 mmol TE/100 g (flesh)		cyanian giycosides	[<u>109][110]</u> [111]	addition, les in the
	21.9 ± 4.0 μmol of QE/100 g (no PBS wash) 17.2 ± 2.0 μmol of QE/100 g (PBS wash)	[<u>8]</u> CAA			aceutical,
	2.4 to 12.8 mmol TE/100 g (peel)			[42]	vrotective,
	0.8 to 2.3 mmol TE/100 g (flesh)	TEAC	[112]		damage
	94.6 to 465.4 mg/mL (red flesh)	DPPH	<u></u>		I flavones
	7.4 ₩5፻፻፲፭ TE μmol/g FW (red flesh)	ABTS	[<u>42</u>]		mbranes, drug), the
	2.5 to 1 <mark>93</mark> TE µmol/g FW (red	FRAP			ticals and

Hyperlipidemia and hyperglycemia are common and frequently occurring diseases, which seriously threaten human health and are often difficult to be completely cured. They will cause a variety of complications, including

Fruit	Antioxidant Capacity	Methods	Main Pigments	References ful health
	flesh)			and their
complications.	ימוועבובנווו מווע ווטג	חופנווו נמוו ופטלנפ	e serum cholesterol and triacylglyce	erol ^[116] , possibly through
inhibit sterol r	egulatory element-	binding protein	^{45]} . The mechanism underlying th	e antibacterial activity of
antioxidants is	still unclear. Howev	ver, some studies	have shown that there are three po	ossible mechanisms for its
antibacterial ac	ctivity: outer membr	ane permeability,	cytoplasmic leakage, and inhibition	of nucleic acid formation
[<u>117][118]</u> . Flavor	noids can directly a	affect the growth	of bacteria and inhibit their pathoge	enic activity ^[119] , such as
Cutibacterium a	acnes (lead to acne	e), Streptococcus	mutans, and Lactobacillus acidophi	ilus (lead to dental caries)
[120]				

On the other hand, independent studies have demonstrated the multiple human health benefits of anthocyanins in the diet ^{[121][122]}. These health benefits include inhibition of various cancers, protection of cardiovascular and nervous systems, and balance of metabolisms, such as blood glucose and lipids, which have been demonstrated in experimental models, such as in vitro, mammalian, and human clinical and epidemiological studies ^[123].

5.4. Bioaccessibility and Bioavailability

Fruit natural pigments exhibit strong antioxidant activity and multiple health benefits. However, whether these beneficial bioactive compounds can be effectively used by the human body depend on their bioaccessibility and bioavailability. Bioaccessibility indicates the proportion of ingested natural pigments that are absorbed by the intestine, and bioavailability indicates the number of natural pigments that enter the blood circulation of the human body after digestion and absorption. Only this part can finally play function or be stored in the human body ^[68]. The primary digestive mechanisms of natural pigments determine their bioaccessibility and bioavailability. In addition, the chemical properties also affect their bioaccessibility and bioavailability. However, the chemical properties of fruit natural pigments are more unstable than other nutrients, such as sugar and organic acids. Thus, as the stability is enhanced, so does the bioavailability ^[124].

Carotenoids are abundant in fruits, but their bioavailability is not as high as expected (10–65%) ^[125]. Carotenoids perform powerful physiological functions only after entering the human bloodstream. Before that, carotenoids are involved in a series of digestive processes and are eventually taken by intestinal mucosal cells and then enter the lymphatic system and circulation. Many factors work together in this process, including release from the food matrix, mass transfer phenomenon, and micelle formation ^[126].

The bioaccessibility of oral phenolic compounds (including flavonoids and anthocyanidins) is 5–10% ^[127]. Flavonoids in fruits usually exist as flavonoid glycosides and PMFs; the modification of these substituents has a great influence on the bioactivity of flavonoids and their bioavailability. The bioaccessibility and bioavailability of flavonoid glycosides depend on the aglycone, both in its type and position. The number and position of the methoxyl group of PMFs have a decisive effect on its biological activity ^[68]. With the increase of the methoxyl group, PMFs exhibit higher hydrophobicity and stronger biological activity. However, it is also a double-edged sword, a high-level hydrophobicity results in poor oral bioavailability, which limits its potential application ^[128]. In

general, compared with flavonoid glycosides, PMFs have lower polarity and approximate planar structure, higher permeability to biofilm, and easier to enter into the blood circulation in vivo ^[42].

The bioaccessibility and bioavailability of anthocyanidins were generally considered to be low, even lower than other flavonoids commonly found in fruits ^[123]. The conclusion was drawn by previous studies only focused on the result that anthocyanidins content in the bloodstream was very low after intake ^{[129][130]}. More and more recent studies revealed the existence of anthocyanidins molecular intermediates, which retain the unique flavonoid C6-C3-C6 backbone structure and have the characteristics that include qualitative diversity, relatively high concentrations, and tenacity ^{[123][131]}. Therefore, the present evaluation method of anthocyanidins bioavailability may be not accurate, but more evidence is needed.

Flavonoids, especially anthocyanidins and proanthocyanidins, have much a lower antioxidant capacity than carotenoids ^[3]. However, due to the water-soluble characteristic, flavonoids are more easily absorbed by the human body than liposoluble carotenoids, which means higher bioaccessibility and bioavailability ^{[80][130]}. In addition, the contents of flavonoids in fruits were more abundant than that of carotenoids ^[3]. Therefore, the health function of flavonoids is more powerful. However, this does not mean that carotenoids are unimportant because some of their properties cannot be replaced, such as the prevention of certain cancers. Therefore, for the highest efficacy, reasonable and balanced intake of these pigment substances is necessary.

References

- 1. Lai, A.; Santangelo, E.; Soressi, G.; Fantoni, R. Analysis of the main secondary metabolites produced in tomato (Lycopersicon esculentum, Mill.) epicarp tissue during fruit ripening using fluorescence techniques. Postharvest Biol. Technol. 2007, 43, 335–342.
- Karppinen, K.; Zoratti, L.; Nguyenquynh, N.; Häggman, H.; Jaakola, L. On the Developmental and Environmental Regulation of Secondary Metabolism in Vaccinium spp. Berries. Front. Plant Sci. 2016, 7, 655.
- 3. Chen, C. Pigments in Fruits and Vegetables; Springer: New York, NY, USA, 2015; ISBN 978-1-4939-2355-7.
- Khoo, H.E.; Azlan, A.; Tang, S.T.; Lim, S.M. Anthocyanidins and anthocyanins: Colored pigments as food, pharmaceutical ingredients, and the potential health benefits. Food Nutr. Res. 2017, 61, 1361779.
- 5. Baby, B.; Antony, P.; Vijayan, R. Antioxidant and anticancer properties of berries. Crit. Rev. Food Sci. Nutr. 2017, 58, 2491–2507.
- 6. Young, A.J.; Lowe, G.L. Carotenoids—Antioxidant Properties. Antioxidants 2018, 7, 28.

- 7. Cazzonelli, C.I.; Pogson, B.J. Source to sink: Regulation of carotenoid biosynthesis in plants. Trends Plant Sci. 2010, 15, 266–274.
- Rodriguez-Concepcion, M.; Avalos, J.; Bonet, M.L.; Boronat, A.; Gomez-Gomez, L.; Hornero-Mendez, D.; Limon, M.C.; Meléndez-Martínez, A.J.; Olmedilla-Alonso, B.; Palou, A.; et al. A global perspective on carotenoids: Metabolism, biotechnology, and benefits for nutrition and health. Prog. Lipid Res. 2018, 70, 62–93.
- 9. Frank, H.A.; Cogdell, R.J. Carotenoids in Photosynthesis. Photochem. Photobiol. 1996, 63, 257–264.
- 10. Roohbakhsh, A.; Karimi, G.; Iranshahi, M. Carotenoids in the treatment of diabetes mellitus and its complications: A mechanistic review. Biomed. Pharmacother. 2017, 91, 31–42.
- 11. Merhan, O. The biochemistry and antioxidant properties of carotenoids. In Carotenoids; Cvetkovic, D.J., Nikolic, G.S., Eds.; InTechOpen: London, UK, 2017; ISBN 978-953-51-3211-0.
- 12. Pérez-Gálvez, A.; Mínguez-Mosquera, M.I. Esterification of xanthophylls and its effect on chemical behavior and bioavailability of carotenoids in the human. Nutr. Res. 2005, 25, 631–640.
- 13. DellaPenna, D.; Pogson, B.J. VITAMIN SYNTHESIS IN PLANTS: Tocopherols and Carotenoids. Annu. Rev. Plant Biol. 2006, 57, 711–738.
- Lu, Y.; Guo, S.; Zhang, F.; Yan, H.; Qian, D.-W.; Wang, H.-Q.; Jin, L.; Duan, J.-A. Comparison of Functional Components and Antioxidant Activity of Lycium barbarum L. Fruits from Different Regions in China. Molecules 2019, 24, 2228.
- 15. Sérino, S.; Gomez, L.; Costagliola, G.; Gautier, H. HPLC Assay of Tomato Carotenoids: Validation of a Rapid Microextraction Technique. J. Agric. Food Chem. 2009, 57, 8753–8760.
- Zhang, L.; Zhang, Q.; Li, W.; Zhang, S.; Xi, W. Identification of key genes and regulators associated with carotenoid metabolism in apricot (Prunus armeniaca) fruit using weighted gene coexpression network analysis. BMC Genom. 2019, 20, 1–15.
- 17. Tritsch, D.; Hemmerlin, A.; Bach, T.J.; Rohmer, M. Plant isoprenoid biosynthesis via the MEP pathway: In vivo IPP/DMAPP ratio produced by (E)-4-hydroxy-3-methylbut-2-enyl diphosphate reductase in tobacco BY-2 cell cultures. FEBS Lett. 2009, 584, 129–134.
- 18. Ruiz-Sola, M. Águila; Rodríguez-Concepción, M. Carotenoid Biosynthesis in Arabidopsis: A Colorful Pathway. Arab. Book 2012, 10, e0158.
- 19. Kuzuyama, T.; Seto, H. Two distinct pathways for essential metabolic precursors for isoprenoid biosynthesis. Proc. Jpn. Acad. Ser. B 2012, 88, 41–52.
- 20. Hunter, W.N. The Non-mevalonate Pathway of Isoprenoid Precursor Biosynthesis. J. Biol. Chem. 2007, 282, 21573–21577.

- 21. Luo, Z.; Zhang, J.; Li, J.; Yang, C.; Wang, T.; Ouyang, B.; Li, H.; Giovannoni, J.; Ye, Z. A STAY-GREEN protein S I SGR 1 regulates lycopene and β-carotene accumulation by interacting directly with S I PSY 1 during ripening processes in tomato. New Phytol. 2013, 198, 442–452.
- 22. Nisar, N.; Li, L.; Lu, S.; Khin, N.C.; Pogson, B.J. Carotenoid Metabolism in Plants. Mol. Plant 2015, 8, 68–82.
- 23. Li, S.; Chen, K.; Grierson, D. A critical evaluation of the role of ethylene and MADS transcription factors in the network controlling fleshy fruit ripening. New Phytol. 2019, 221, 1724–1741.
- 24. Li, L.; Wang, X.; Zhang, X.; Guo, M.; Liu, T. Unraveling the target genes of RIN transcription factor during tomato fruit ripening and softening. J. Sci. Food Agric. 2016, 97, 991–1000.
- Pan, I.L.; McQuinn, R.; Giovannoni, J.J.; Irish, V.F. Functional diversification of AGAMOUS lineage genes in regulating tomato flower and fruit development. J. Exp. Bot. 2010, 61, 1795– 1806.
- Yin, W.; Yu, X.; Chen, G.; Tang, B.; Wang, Y.; Liao, C.; Zhang, Y.; Hu, Z. Suppression of SIMBP15 Inhibits Plant Vegetative Growth and Delays Fruit Ripening in Tomato. Front. Plant Sci. 2018, 9, 938.
- Dong, T.; Hu, Z.; Deng, L.; Wang, Y.; Zhu, M.; Zhang, J.; Chen, G. A Tomato MADS-Box Transcription Factor, SIMADS1, Acts as a Negative Regulator of Fruit Ripening. Plant Physiol. 2013, 163, 1026–1036.
- Yin, W.; Hu, Z.; Cui, B.; Guo, X.; Hu, J.; Zhu, Z.; Chen, G. Suppression of the MADS-box gene SIMBP8 accelerates fruit ripening of tomato (Solanum lycopersicum). Plant Physiol. Biochem. 2017, 118, 235–244.
- Zhu, M.; Chen, G.; Zhou, S.; Tu, Y.; Wang, Y.; Dong, T.; Hu, Z. A New Tomato NAC (NAM/ATAF1/2/CUC2) Transcription Factor, SINAC4, Functions as a Positive Regulator of Fruit Ripening and Carotenoid Accumulation. Plant Cell Physiol. 2013, 55, 119–135.
- 30. Peer, W.A.; Murphy, A.S. Flavonoids and auxin transport: Modulators or regulators? Trends Plant Sci. 2007, 12, 556–563.
- 31. Cetinkaya, H.; Kulak, M.; Karaman, M.; Karaman, H.S.; Kocer, F. Flavonoid Accumulation Behavior in Response to the Abiotic Stress: Can a Uniform Mechanism Be Illustrated for All Plants? In Flavonoids—From Biosynthesis to Human Health; IntechOpen: London, UK, 2017.
- Corradini, E.; Foglia, P.; Giansanti, P.; Gubbiotti, R.; Samperi, R.; Laganà, A. Flavonoids: Chemical properties and analytical methodologies of identification and quantitation in foods and plants. Nat. Prod. Res. 2011, 25, 469–495.
- 33. Wollenweber, E.; Mann, K. Neue Flati onoide aus Primelmehl. Biochem. Physiol. Pflanz. 1986, 181, 665–669.

- 34. Awouafack, M.D.; Tane, P.; Morita, H. Isolation and Structure Characterization of Flavonoids. In Flavonoids—From Biosynthesis to Human Health; InTechOpen: London, UK, 2017.
- 35. E Heim, K.; Tagliaferro, A.R.; Bobilya, D.J. Flavonoid antioxidants: Chemistry, metabolism and structure-activity relationships. J. Nutr. Biochem. 2002, 13, 572–584.
- 36. Yerragunta, V.; Kumaraswamy, T.; Suman, D.; Anusha, V.; Patil, P.; Samhitha, T. A review on chalcones and its importance. Pharma Tutor 2013, 1, 54–59.
- Escobar-Cévoli, R.; Castro-Espín, C.; Béraud, G.B.V.; Buckland, G.; Zamora-Ros, R.; Béraud, V. An Overview of Global Flavonoid Intake and its Food Sources. In Flavonoids—From Biosynthesis to Human Health; IntechOpen: London, UK, 2017.
- Drappier, J.; Thibon, C.; Rabot, A.; Geny-Denis, L. Relationship between wine composition and temperature: Impact on Bordeaux wine typicity in the context of global warming—Review. Crit. Rev. Food Sci. Nutr. 2019, 59, 14–30.
- 39. Mattivi, F.; Guzzon, R.; Vrhovsek, U.; Stefanini, A.M.; Velasco, R. Metabolite Profiling of Grape: Flavonols and Anthocyanins. J. Agric. Food Chem. 2006, 54, 7692–7702.
- 40. Sagar, N.A.; Pareek, S.; Sharma, S.; Yahia, E.M.; Lobo, M.G. Fruit and Vegetable Waste: Bioactive Compounds, Their Extraction, and Possible Utilization. Compr. Rev. Food Sci. Food Saf. 2018, 17, 512–531.
- 41. Mahato, N.; Sharma, K.; Sinha, M.; Cho, M.H. Citrus waste derived nutra-/pharmaceuticals for health benefits: Current trends and future perspectives. J. Funct. Foods 2018, 40, 307–316.
- 42. Gao, Z.; Gao, W.; Zeng, S.-L.; Li, P.; Liu, E.-H. Chemical structures, bioactivities and molecular mechanisms of citrus polymethoxyflavones. J. Funct. Foods 2018, 40, 498–509.
- 43. Manthey, J.A.; Grohmann, K. Phenols in Citrus Peel Byproducts. Concentrations of Hydroxycinnamates and Polymethoxylated Flavones in Citrus Peel Molasses. J. Agric. Food Chem. 2001, 49, 3268–3273.
- 44. Gómez-Mejía, E.; Rosales-Conrado, N.; León-González, M.E.; Madrid, Y. Citrus peels waste as a source of value-added compounds: Extraction and quantification of bioactive polyphenols. Food Chem. 2019, 295, 289–299.
- 45. Duan, L.; Dou, L.-L.; Yu, K.-Y.; Guo, L.; Bai-Zhong, C.; Li, P.; Liu, E.-H. Polymethoxyflavones in peel of Citrus reticulata 'Chachi' and their biological activities. Food Chem. 2017, 234, 254–261.
- 46. de Albuquerque, M.A.C.; Levit, R.; Beres, C.; Bedani, R.; LeBlanc, A.D.M.D.; Saad, S.M.I.; LeBlanc, J.G. Tropical fruit by-products water extracts as sources of soluble fibres and phenolic compounds with potential antioxidant, anti-inflammatory, and functional properties. J. Funct. Foods 2019, 52, 724–733.

- Petrussa, E.; Braidot, E.; Zancani, M.; Peresson, C.; Bertolini, A.; Patui, S.; Vianello, A. Plant Flavonoids—Biosynthesis, Transport and Involvement in Stress Responses. Int. J. Mol. Sci. 2013, 14, 14950–14973.
- Nabavi, S.M.; Šamec, D.; Tomczyk, M.; Milella, L.; Russo, D.; Habtemariam, S.; Suntar, I.; Rastrelli, L.; Daglia, M.; Xiao, J.; et al. Flavonoid biosynthetic pathways in plants: Versatile targets for metabolic engineering. Biotechnol. Adv. 2020, 38, 107316.
- 49. Gouot, J.C.; Smith, J.P.; Holzapfel, B.P.; Walker, A.R.; Barril, C. Grape berry flavonoids: A review of their biochemical responses to high and extreme high temperatures. J. Exp. Bot. 2018, 70, 397–423.
- 50. Gutierrez, E.; Velasco, A.G.; Lucas, J.A.; Gutierrez-Mañero, F.J.; Ramos-Solano, B. The Flavonol-Anthocyanin Pathway in Blackberry and Arabidopsis: State of the Art. In Flavonoids—From Biosynthesis to Human Health; IntechOpen: London, UK, 2017.
- 51. Liu, J.; Osbourn, A.; Ma, P. MYB Transcription Factors as Regulators of Phenylpropanoid Metabolism in Plants. Mol. Plant 2015, 8, 689–708.
- Czemmel, S.; Stracke, R.; Weisshaar, B.; Cordon, N.; Harris, N.N.; Walker, A.R.; Robinson, S.P.; Bogs, J. The Grapevine R2R3-MYB Transcription Factor VvMYBF1 Regulates Flavonol Synthesis in Developing Grape Berries. Plant Physiol. 2009, 151, 1513–1530.
- 53. Wang, J.; Wang, F.; Jin, C.; Tong, Y.; Wang, T. A R2R3-MYB transcription factor VvMYBF1 from grapevine (Vitis vinifera L.) regulates flavonoids accumulation and abiotic stress tolerance in transgenic Arabidopsis. J. Hortic. Sci. Biotechnol. 2019, 95, 147–161.
- 54. Xu, W.; Dubos, C.; Lepiniec, L. Transcriptional control of flavonoid biosynthesis by MYB–bHLH– WDR complexes. Trends Plant Sci. 2015, 20, 176–185.
- 55. Zhao, C.; Liu, X.; Gong, Q.; Cao, J.; Shen, W.; Yin, X.; Grierson, D.; Zhang, B.; Xu, C.; Li, X.; et al. Three AP2/ERF family members modulate flavonoid synthesis by regulating type IV chalcone isomerase in citrus. Plant Biotechnol. J. 2021, 19, 671–688.
- 56. Perezjimenez, J.; Neveu, V.; Vos, F.; Scalbert, A. Identification of the 100 richest dietary sources of polyphenols: An application of the Phenol-Explorer database. Eur. J. Clin. Nutr. 2010, 64, S112–S120.
- 57. Zhou, H.; Lin-Wang, K.; Wang, F.; Espley, R.V.; Ren, F.; Zhao, J.; Ogutu, C.; He, H.; Jiang, Q.; Allan, A.C.; et al. Activator-type R2R3-MYB genes induce a repressor-type R2R3-MYB gene to balance anthocyanin and proanthocyanidin accumulation. New Phytol. 2019, 221, 1919–1934.
- 58. Martín, J.; Kuskoski, E.M.; Navas, M.J.; Asuero, A.G. Antioxidant Capacity of Anthocyanin Pigments. In Flavonoids—From Biosynthesis to Human Health; IntechOpen: London, UK, 2017.

- 59. Bueno, J.M.; Sáez-Plaza, P.; Ramos-Escudero, F.; Jiménez, A.M.; Fett, R.; Asuero, A.G. Analysis and Antioxidant Capacity of Anthocyanin Pigments. Part II: Chemical Structure, Color, and Intake of Anthocyanins. Crit. Rev. Anal. Chem. 2012, 42, 126–151.
- 60. Blancquaert, E.; Oberholster, A.; Ricardo-Da-Silva, J.; Deloire, A. Effects of Abiotic Factors on Phenolic Compounds in the Grape Berry—A Review. S. Afr. J. Enol. Vitic. 2018, 40, 40.
- 61. Norberto, S.; Silva, S.; Meireles, M.; Faria, A.; Pintado, M.; Calhau, C. Blueberry anthocyanins in health promotion: A metabolic overview. J. Funct. Foods 2013, 5, 1518–1528.
- Wang, H.; Guo, X.; Hu, X.; Li, T.; Fu, X.; Liu, R.H. Comparison of phytochemical profiles, antioxidant and cellular antioxidant activities of different varieties of blueberry (Vaccinium spp.). Food Chem. 2017, 217, 773–781.
- 63. Montefiori, M.; Comeskey, D.J.; Wohlers, M.; McGhie, T.K. Characterization and Quantification of Anthocyanins in Red Kiwifruit (Actinidia spp.). J. Agric. Food Chem. 2009, 57, 6856–6861.
- 64. Peng, Y.; Lin-Wang, K.; Cooney, J.M.; Wang, T.; Espley, R.V.; Allan, A.C. Differential regulation of the anthocyanin profile in purple kiwifruit (Actinidia species). Hortic. Res. 2019, 6, 1–16.
- Sun, C.; Deng, L.; Du, M.; Zhao, J.; Chen, Q.; Huang, T.; Jiang, H.; Li, C.-B.; Li, C. A Transcriptional Network Promotes Anthocyanin Biosynthesis in Tomato Flesh. Mol. Plant 2020, 13, 42–58.
- 66. Jian, W.; Cao, H.; Yuan, S.; Liu, Y.; Lu, J.; Lu, W.; Li, N.; Wang, J.; Zou, J.; Tang, N.; et al. SIMYB75, an MYB-type transcription factor, promotes anthocyanin accumulation and enhances volatile aroma production in tomato fruits. Hortic. Res. 2019, 6, 1–15.
- 67. Zhang, L.; Hu, J.; Han, X.; Li, J.; Gao, Y.; Richards, C.M.; Zhang, C.; Tian, Y.; Liu, G.; Gul, H.; et al. A high-quality apple genome assembly reveals the association of a retrotransposon and red fruit colour. Nat. Commun. 2019, 10, 1–13.
- 68. Singh, B.; Singh, J.P.; Kaur, A.; Singh, N. Phenolic composition, antioxidant potential and health benefits of citrus peel. Food Res. Int. 2020, 132, 109114.
- 69. Bakoyiannis, I.; Daskalopoulou, A.; Pergialiotis, V.; Perrea, D. Phytochemicals and cognitive health: Are flavonoids doing the trick? Biomed. Pharmacother. 2019, 109, 1488–1497.
- Santos-Sánchez, N.F.; Salas-Coronado, R.; Villanueva-Cañongo, C.; Hernández-Carlos, B. Antioxidant Compounds and Their Antioxidant Mechanism. In Antioxidants; Shalaby, E., Ed.; IntechOpen: London, UK, 2019; pp. 1–28.
- 71. Mishra, A.; Kumar, S.; Pandey, A.K. Scientific Validation of the Medicinal Efficacy of Tinospora cordifolia. Sci. World J. 2013, 2013, 1–8.
- 72. Mishra, A.; Sharma, A.K.; Kumar, S.; Saxena, A.K.; Pandey, A.K. Bauhinia variegataLeaf Extracts Exhibit Considerable Antibacterial, Antioxidant, and Anticancer Activities. BioMed Res. Int. 2013,

2013, 1–10.

- 73. Kheradmand, E.; Moghaddam, A.H.; Zare, M. Neuroprotective effect of hesperetin and nanohesperetin on recognition memory impairment and the elevated oxygen stress in rat model of Alzheimer's disease. Biomed. Pharmacother. 2018, 97, 1096–1101.
- 74. Frandsen, J.; Narayanasamy, P. Flavonoid Enhances the Glyoxalase Pathway in Cerebellar Neurons to Retain Cellular Functions. Sci. Rep. 2017, 7, 1–10.
- Cai, Y.; Qin, W.; Ketnawa, S.; Ogawa, Y. Impact of particle size of pulverized citrus peel tissue on changes in antioxidant properties of digested fluids during simulated in vitro digestion. Food Sci. Hum. Wellness 2020, 9, 58–63.
- 76. Alam, N.; Bristi, N.J.; Rafiquzzaman. Review on in vivo and in vitro methods evaluation of antioxidant activity. Saudi Pharm. J. 2013, 21, 143–152.
- 77. Mohamed, S.A.; Awad, M.A.; El-Dengawy, E.-R.F.A.; Abdel-Mageed, H.M.; El-Badry, M.O.; Salah, H.A.; Abdel-Aty, A.M.; Fahmy, A.S. Total phenolic and flavonoid contents and antioxidant activities of sixteen commercial date cultivars grown in Saudi Arabia. RSC Adv. 2016, 6, 44814–44819.
- Li, H.; Deng, Z.; Liu, R.; Loewen, S.; Tsao, R. Ultra-performance liquid chromatographic separation of geometric isomers of carotenoids and antioxidant activities of 20 tomato cultivars and breeding lines. Food Chem. 2012, 132, 508–517.
- Fattore, M.; Montesano, D.; Pagano, E.; Teta, R.; Borrelli, F.; Mangoni, A.; Seccia, S.; Albrizio, S. Carotenoid and flavonoid profile and antioxidant activity in "Pomodorino Vesuviano" tomatoes. J. Food Compos. Anal. 2016, 53, 61–68.
- 80. Tagliazucchi, D.; Verzelloni, E.; Bertolini, D.; Conte, A. In vitro bio-accessibility and antioxidant activity of grape polyphenols. Food Chem. 2010, 120, 599–606.
- Wolfe, K.L.; Kang, X.; He, X.; Dong, M.; Zhang, Q.; Liu, R.H. Cellular Antioxidant Activity of Common Fruits. J. Agric. Food Chem. 2008, 56, 8418–8426.
- Avidzba, A.M.; Kubyshkin, A.V.; I Guguchkina, T.; A Markosov, V.; Katsev, A.M.; Naumova, N.V.; I Shramko, Y.; Zaytsev, G.P.; Chernousova, I.V.; A Ogay, Y.; et al. The antioxidant activity of the products of processing of red grape of Cabernet Sauvignon, Merlot, Saperavi. Vopr. Pitan. 2016, 85, 99–109. (In Russian)
- 83. Xia, E.-Q.; Deng, G.-F.; Guo, Y.-J.; Li, H.-B. Biological Activities of Polyphenols from Grapes. Int. J. Mol. Sci. 2010, 11, 622–646.
- 84. Pastrana-Bonilla, E.; Akoh, C.C.; Sellappan, S.; Krewer, G. Phenolic Content and Antioxidant Capacity of Muscadine Grapes. J. Agric. Food Chem. 2003, 51, 5497–5503.
- 85. Poudel, P.R.; Tamura, H.; Kataoka, I.; Mochioka, R. Phenolic compounds and antioxidant activities of skins and seeds of five wild grapes and two hybrids native to Japan. J. Food Compos.

Anal. 2008, 21, 622–625.

- 86. de Souza, V.R.; Pereira, P.A.P.; da Silva, T.L.T.; Lima, L.C.D.O.; Pio, R.; Queiroz, F. Determination of the bioactive compounds, antioxidant activity and chemical composition of Brazilian blackberry, red raspberry, strawberry, blueberry and sweet cherry fruits. Food Chem. 2014, 156, 362–368.
- 87. Cásedas, G.; Les, F.; Gómez-Serranillos, M.P.; Smith, C.; López, V. Anthocyanin profile, antioxidant activity and enzyme inhibiting properties of blueberry and cranberry juices: A comparative study. Food Funct. 2017, 8, 4187–4193.
- Haytowitz, D.B.; Bhagwat, S.A. USDA Database for the Oxygen Radical Absorbance Capacity (ORAC) of Selected Foods, Release 2. USDA National Nutrient Database for Standard Reference. 2010; pp. 10–48. Available online: (accessed on 3 May 2021).
- Schulz, M.; Seraglio, S.K.T.; Della Betta, F.; Nehring, P.; Valese, A.C.; Daguer, H.; Gonzaga, L.V.; Costa, A.C.O.; Fett, R. Blackberry (Rubus ulmifolius Schott): Chemical composition, phenolic compounds and antioxidant capacity in two edible stages. Food Res. Int. 2019, 122, 627–634.
- 90. Sellappan, S.; Akoh, C.C.; Krewer, G. Phenolic Compounds and Antioxidant Capacity of Georgia-Grown Blueberries and Blackberries. J. Agric. Food Chem. 2002, 50, 2432–2438.
- Aaby, K.; Ekeberg, A.D.; Skrede, G. Characterization of Phenolic Compounds in Strawberry (Fragaria × ananassa) Fruits by Different HPLC Detectors and Contribution of Individual Compounds to Total Antioxidant Capacity. J. Agric. Food Chem. 2007, 55, 4395–4406.
- 92. Van De Velde, F.; Tarola, A.M.; Güemes, D.; Pirovani, M.E. Bioactive Compounds and Antioxidant Capacity of Camarosa and Selva Strawberries (Fragaria x ananassa Duch.). Foods 2013, 2, 120– 131.
- 93. Serrano, M.; Guillén, F.; Martínez-Romero, D.; Castillo, S.; Valero, D. Chemical Constituents and Antioxidant Activity of Sweet Cherry at Different Ripening Stages. J. Agric. Food Chem. 2005, 53, 2741–2745.
- 94. Kirakosyan, A.; Seymour, E.; Llanes, D.E.U.; Kaufman, P.B.; Bolling, S.F. Chemical profile and antioxidant capacities of tart cherry products. Food Chem. 2009, 115, 20–25.
- 95. Liao, X.; Greenspan, P.; Pegg, R.B. Characterizing the phenolic constituents and antioxidant capacity of Georgia peaches. Food Chem. 2019, 271, 345–353.
- 96. Gil Muñoz, M.I.; Tomás-Barberán, F.A.; Hess-Pierce, B.; Kader, A.A. Antioxidant Capacities, Phenolic Compounds, Carotenoids, and Vitamin C Contents of Nectarine, Peach, and Plum Cultivars from California. J. Agric. Food Chem. 2002, 50, 4976–4982.
- 97. Arion, C.M.; Tabart, J.; Kevers, C.; Niculaua, M.; Filimon, R.; Beceanu, D.; Dommes, J. Antioxidant potential of different plum cultivars during storage. Food Chem. 2014, 146, 485–491.

- 98. Kim, S.-N.; Kim, M.-R.; Cho, S.-M.; Kim, S.-Y.; Kim, J.-B.; Cho, Y.-S. Antioxidant activities and determination of phenolic compounds isolated from oriental plums (Soldam, Oishiwase and Formosa). Nutr. Res. Pract. 2012, 6, 277–285.
- 99. Leccese, A.; Bartolini, S.; Viti, R. From Genotype to Apricot Fruit Quality: The Antioxidant Properties Contribution. Plant Foods Hum. Nutr. 2012, 67, 317–325.
- 100. Xi, W.; Feng, J.; Liu, Y.; Zhang, S.; Zhao, G. The R2R3-MYB transcription factor PaMYB10 is involved in anthocyanin biosynthesis in apricots and determines red blushed skin. BMC Plant Biol. 2019, 19, 1–14.
- 101. Chen, X.-M.; Tait, A.R.; Kitts, D.D. Flavonoid composition of orange peel and its association with antioxidant and anti-inflammatory activities. Food Chem. 2017, 218, 15–21.
- 102. Ernawita; Wahyuono, R.A.; Hesse, J.; Hipler, U.-C.; Elsner, P.; Böhm, V. Carotenoids of indigenous citrus species from Aceh and its in vitro antioxidant, antidiabetic and antibacterial activities. Eur. Food Res. Technol. 2016, 242, 1869–1881.
- 103. Kschonsek, J.; Wolfram, T.; Stöckl, A.; Böhm, V. Polyphenolic Compounds Analysis of Old and New Apple Cultivars and Contribution of Polyphenolic Profile to the In Vitro Antioxidant Capacity. Antioxidants 2018, 7, 20.
- 104. Wang, X.; Li, C.; Liang, D.; Zou, Y.; Li, P.; Ma, F. Phenolic compounds and antioxidant activity in red-fleshed apples. J. Funct. Foods 2015, 18, 1086–1094.
- 105. Dorgan, J.F.; Ziegler, R.G.; Schoenberg, J.B.; Hartge, P.; McAdams, M.J.; Falk, R.T.; Wilcox, H.B.; Shaw, G.L. Race and sex differences in associations of vegetables, fruits, and carotenoids with lung cancer risk in New Jersey (United States). Cancer Causes Control 1993, 4, 273–281.
- 106. Astorg, P. Food carotenoids and cancer prevention: An overview of current research. Trends Food Sci. Technol. 1997, 8, 406–413.
- 107. Perez-Cornago, A.; Travis, R.C.; Appleby, P.N.; Tsilidis, K.K.; Tjønneland, A.; Olsen, A.; Overvad, K.; Katzke, V.; Kuehn, T.; Trichopoulou, A.; et al. Fruit and vegetable intake and prostate cancer risk in the European Prospective Investigation into Cancer and Nutrition (EPIC). Int. J. Cancer 2017, 141, 287–297.
- 108. Goralczyk, R. ß-Carotene and Lung Cancer in Smokers: Review of Hypotheses and Status of Research. Nutr. Cancer 2009, 61, 767–774.
- 109. Williams, A.W.; Boileau, T.W.-M.; Zhou, J.R.; Clinton, S.K.; Erdman, J.J.W. β-Carotene Modulates Human Prostate Cancer Cell Growth and May Undergo Intracellular Metabolism to Retinol. J. Nutr. 2000, 130, 728–732.
- 110. Gill, J.K.; Franke, A.A.; Morris, J.S.; Cooney, R.V.; Wilkens, L.R.; Le Marchand, L.; Goodman, M.T.; Henderson, B.E.; Kolonel, L.N. Association of selenium, tocopherols, carotenoids, retinol,

and 15-isoprostane F2t in serum or urine with prostate cancer risk: The multiethnic cohort. Cancer Causes Control 2009, 20, 1161–1171.

- 111. Lindbergh, C.A.; Renzi-Hammond, L.M.; Hammond, B.R.; Terry, U.P.; Mewborn, C.M.; Puente, A.N.; Miller, L.S. Lutein and Zeaxanthin Influence Brain Function in Older Adults: A Randomized Controlled Trial. J. Int. Neuropsychol. Soc. 2017, 24, 77–90.
- 112. Asadi-Samani, M.; Farkhad, N.K.; Mahmoudian-Sani, M.R.; Shirzad, H. Antioxidants as a Double-Edged Sword in the Treatment of Cancer. Antioxidants 2019.
- 113. S, A.N.; Sr, R.K.; Nair, A.S.; Baby, S. Citrus peels prevent cancer. Phytomedicine 2018, 50, 231– 237.
- 114. Chait, A. Hyperlipidemia: Forestalling complications in older diabetics. Geriatrics 1985, 40, 71–73.
- 115. Schrijvers, B.F.; De Vriese, A.S.; Flyvbjerg, A. From Hyperglycemia to Diabetic Kidney Disease: The Role of Metabolic, Hemodynamic, Intracellular Factors and Growth Factors/Cytokines. Endocr. Rev. 2004, 25, 971–1010.
- 116. Kurowska, E.M.; Manthey, J.A. Hypolipidemic Effects and Absorption of Citrus Polymethoxylated Flavones in Hamsters with Diet-Induced Hypercholesterolemia. J. Agric. Food Chem. 2004, 52, 2879–2886.
- 117. Ajiboye, T.; Mohammed, A.; Bello, S.; Yusuf, I.; Ibitoye, O.; Muritala, H.; Onajobi, I. Antibacterial activity of Syzygium aromaticum seed: Studies on oxidative stress biomarkers and membrane permeability. Microb. Pathog. 2016, 95, 208–215.
- 118. Zhang, Y.; Wu, Y.-T.; Zheng, W.; Han, X.-X.; Jiang, Y.-H.; Hu, P.-L.; Tang, Z.-X.; Shi, L.-E. The antibacterial activity and antibacterial mechanism of a polysaccharide from Cordyceps cicadae. J. Funct. Foods 2017, 38, 273–279.
- Papuc, C.; Goran, G.V.; Predescu, C.N.; Nicorescu, V.; Stefan, G. Plant Polyphenols as Antioxidant and Antibacterial Agents for Shelf-Life Extension of Meat and Meat Products: Classification, Structures, Sources, and Action Mechanisms. Compr. Rev. Food Sci. Food Saf. 2017, 16, 1243–1268.
- 120. Shetty, S.B.; Mahin-Syed-Ismail, P.; Varghese, S.; Thomas-George, B.; Thajuraj, P.K.-; Baby, D.; Haleem, S.; Sreedhar, S.; Divakar, D.D. Antimicrobial effects of Citrus sinensis peel extracts against dental caries bacteria: An in vitro study. J. Clin. Exp. Dent. 2016, 8, e71–e77.
- 121. Yue, E.; Tuguzbaeva, G.; Chen, X.; Qin, Y.; Li, A.; Sun, X.; Dong, C.; Liu, Y.; Yu, Y.; Zahra, S.M.; et al. Anthocyanin is involved in the activation of pyroptosis in oral squamous cell carcinoma. Phytomedicine 2019, 56, 286–294.
- 122. Blagojević, B.; Agić, D.; Serra, A.T.; Matić, S.; Matovina, M.; Bijelić, S.; Popović, B.M. An in vitro and in silico evaluation of bioactive potential of cornelian cherry (Cornus mas L.) extracts rich in

polyphenols and iridoids. Food Chem. 2021, 335, 127619.

- 123. Lila, M.A.; Burton-Freeman, B.; Grace, M.; Kalt, W. Unraveling Anthocyanin Bioavailability for Human Health. Annu. Rev. Food Sci. Technol. 2016, 7, 375–393.
- 124. Cortez, R.; Luna-Vital, D.A.; Margulis, D.; De Mejia, E.G. Natural Pigments: Stabilization Methods of Anthocyanins for Food Applications. Compr. Rev. Food Sci. Food Saf. 2016, 16, 180–198.
- 125. Saini, R.K.; Nile, S.H.; Park, S.W. Carotenoids from fruits and vegetables: Chemistry, analysis, occurrence, bioavailability and biological activities. Food Res. Int. 2015, 76, 735–750.
- 126. Lemmens, L.; Colle, I.; Van Buggenhout, S.; Palmero, P.; Van Loey, A.; Hendrickx, M. Carotenoid bioaccessibility in fruit- and vegetable-based food products as affected by product (micro)structural characteristics and the presence of lipids: A review. Trends Food Sci. Technol. 2014, 38, 125–135.
- 127. Cardona, F.; Andrés-Lacueva, C.; Tulipani, S.; Tinahones, F.J.; Queipo-Ortuño, M.I. Benefits of polyphenols on gut microbiota and implications in human health. J. Nutr. Biochem. 2013, 24, 1415–1422.
- 128. Lu, X.; Zhang, H.; Zheng, T.; Liu, Q.; Zhu, J.; Huang, Q. Evaluation of Oral Bioaccessibility of Aged Citrus Peel Extracts Encapsulated in Different Lipid-Based Systems: A Comparison Study Using Different In Vitro Digestion Models. J. Agric. Food Chem. 2020, 68, 97–105.
- 129. Fernandes, I.; Faria, A.; Calhau, C.; de Freitas, V.; Mateus, N. Bioavailability of anthocyanins and derivatives. J. Funct. Foods 2014, 7, 54–66.
- 130. Sengul, H.; Surek, E.; Nilufer-Erdil, D. Investigating the effects of food matrix and food components on bioaccessibility of pomegranate (Punica granatum) phenolics and anthocyanins using an in-vitro gastrointestinal digestion model. Food Res. Int. 2014, 62, 1069–1079.
- 131. Zhang, H.; Hassan, Y.I.; Renaud, J.; Liu, R.; Yang, C.; Sun, Y.; Tsao, R. Bioaccessibility, bioavailability, and anti-inflammatory effects of anthocyanins from purple root vegetables using mono- and co-culture cell models. Mol. Nutr. Food Res. 2017, 61, 61.

Retrieved from https://encyclopedia.pub/entry/history/show/23164