

## Flavonoid-rich foods (FRF): A promising nutraceutical approach against lifespan-shortening diseases

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### ABSTRACT

It is well documented that life expectancy in developed countries at birth is going to surpass the 20<sup>th</sup> century. However, regrettably, a potential decline in life expectancy has been proposed for these nations in the 21<sup>st</sup> century due to a rapid upsurge in the prevalence of fatal degenerative diseases like cardiovascular diseases (CVD), cancer and diabetes. Collectively, these three diseases accounted for 65% of all deaths in urbanized societies and were considered as a dynamic issue for shortening the genetically determined lifespan through increased mortalities, morbidities, disabilities, immense sufferings, and premature aging. These fatal degenerative diseases and premature aging are closely associated with oxidative stress produced by the free radicals in the body. In epidemiologic studies, flavonoid-rich foods (FRF) like fruits, vegetables, and beverages have been associated as protective agents against these diseases. These also have been observed for their geroprotective effects and help in preventing premature aging and deterioration of brain function, which is related to Alzheimer's disease and dementia. In this review, we presented a comprehensive overview of the FRF for their potential role against lifespan-shortening complications, i.e., CVD, cancer, and diabetes. We also have drawn the future perspective and dietary guidelines to reduce the fatal disease burden in urban populations.

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### Introduction

Food concept in the world is changing from a past emphasis on hunger satisfaction, the absence of the classical nutrient deficiency diseases and survival to an increasing focus on foods promising use as functional ingredients to provide better well-being and health (1). Substantial data from epidemiological surveys and nutritional intervention studies indicate that different components of food have biological characteristics that exhibit activities in modulating animal and human metabolism in a manner favorable for the longevity and several fatal disease prevention (2-4), and foods containing these components are called "functional foods", which are sometimes dubbed as 'superfoods' in medical communities (5, 6). The term "superfood" has been valuable for marketing purpose, having no precise scientific definition (6). Nevertheless, it is rising in the scientific literature with the sense of plant foods that contain phytochemicals that confer remarkable health benefits thus potentially increasing life expectancy. Until recently, the relationship between diet and lifespan has been hardly understood. Nevertheless, animal studies

have revealed that dietary manipulation could extend the mean and maximum lifespan and significantly slow down the progression or even completely prevent the age-dependent pathologies (7).

It is well documented that life expectancy at birth in developed countries has been increasing by three months per year since 1850 (8). Life expectancy at birth was 47.3 years at the beginning of the 20<sup>th</sup> century, but now after a dramatic increase in the past 100 years, it is nearly 77 years, because of reduced mortalities from infections, increased income of people, and several other dietary, lifestyle-, and health-related factors (6, 7, 9). Regrettably, in the 21<sup>st</sup> century, a potential decline in life expectancy has been proposed for developed nations (10) due to rapid trends in the prevalence of fatal diseases like cardiovascular diseases, cancer, and diabetes (10, 11), which accounted for 65% of all deaths in the USA. These diseases shorten life expectancy through increased mortalities, disabilities, and enormous sufferings and undermine health conditions (11). There is growing evidence that oxidative stress produced by some reactive oxygen compounds as

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well as free radicals such as nitric oxide (\*NO) radical, hydroxyl (\*OH) radical, lipid peroxy (LOO\*) radical, and superoxide (O<sub>2</sub><sup>-</sup>) radical are the causative agents for a number of human diseases (12, 13). Therefore, particular attention is given to search for powerful antioxidant agents to attenuate the incidence and progression of lifespan-shortening diseases and to reinforce the genetically determined lifespan (13).

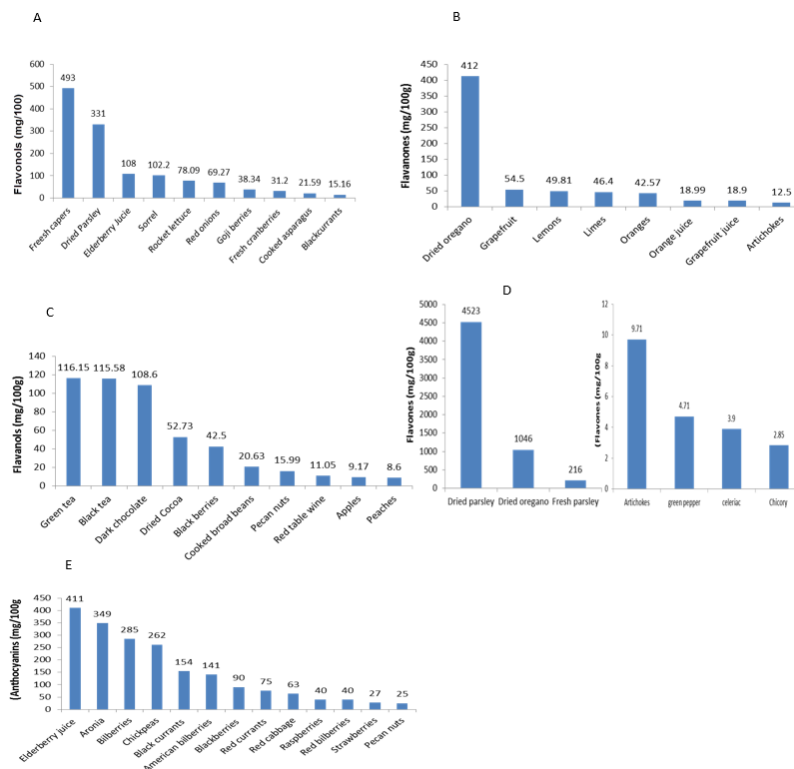
Among natural antioxidants, phytochemicals which are ubiquitous in plants with their major nutraceutical part called flavonoids (14), are considered as potent natural antioxidants. Over 8000, various natural flavonoids have been already described as mentioned by Croft (15), and this list is still growing. Some health-oriented biological effects are attributed to their antioxidant potency (16). Flavonoids have been shown to possess antioxidant properties within *in vitro* experimental systems (4) and anti-inflammatory (17), immune-modulator (16), antiviral (18), antiallergic (19), and anticarcinogenic properties (20). Their antioxidant and inflammatory properties help in toxin-mediated stress and chronic disease prevention (21). Due to their surprising biological health effects, flavonoids are considered 'disease-preventing, health-promoting dietary supplements' (22). So, more than 30000 publications/ year in the past few years were focused on health-promoting impacts of flavonoids (23). Foods containing these flavonoids have also been attracting considerable attention in the medical and public communities because of evidence from various literature which suggest that consumption of Flavonoid-rich foods (FRF) could potentially improve human health and well-being (24). Numerous dietary intervention studies have proven that consumption of plant products (e.g., fruits and vegetables) declines the development of pathological condition risks, which include genetic

and nervous system disorders, inflammatory and cardiovascular diseases, and cancer (25, 26). A series of studies in China indicated that high intake of fruits and vegetables could prevent breast cancer (27), coronary heart disease (28), cataracts, diabetes, Alzheimer's disease, and even asthma (29, 30). According to the world health report, 4.4% of the overall disease burden including disability and mortality in Europe could be attributed to low fruit and vegetable intake (31).

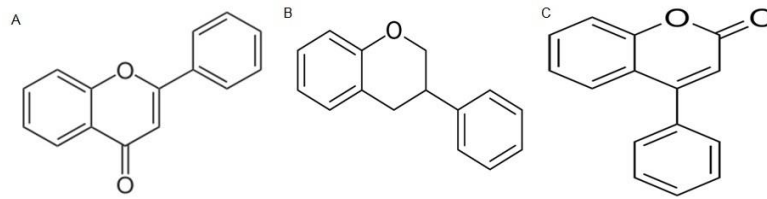
All these studies revealed that plant-derived FRF could reinforce life expectancy by cutting or preventing the risk of chronic lifespan-shortening disorders including diabetes, cancer, and cardiovascular diseases (10, 11) by dropping their associated risk factors, which positively affect the lifespan (32). The present review has focused on the evidence and mechanisms involved in the protective role of FRF against global lifespan-shortening complications including cardiovascular diseases, cancer, and diabetes. We also have drawn the evidence-based dietary guidelines for the readers to minimize the effects of aging, reduce the occurrence of degenerative diseases, and to get the genetically determined lifespan. Moreover, we have proposed a future perspective to identify priorities in food for a particular condition.

### Sources and phytochemistry

Flavonoid-rich foods, based on their surprising health effects are well described as superfoods. These include all plant origin foods mainly tea, fruit, vegetables, grains, legumes, nuts, and wine (33). Tea and wine are the primary dietary sources of flavonoids in eastern and western societies, respectively. Besides, leafy vegetables, onions, apples, berries, cherries, soybeans, and citrus fruits are considered an important source of dietary flavonoids (34-36). The content of flavonoids in different food sources is shown in Figures 1, A-E.



**Figure 1.** A. Mean concentration of flavonols (mg/100 g) in different food sources. B. Mean concentration of flavanonones (mg/100 g) in various food sources. C. Mean concentration of flavanolols (mg/100 g) in different food sources. For green and black tea, leaves to water ratio are 1:20 (w/v). D. Mean concentration of anthocyanins (mg/100 g) in different food sources E. Mean concentration of flavones (mg/100 g) in various food sources (37)



**Figure 2.** Basic chemical structural and types of flavonoids. A: Molecular structure of flavones backbone (2-phenyl-1,4-benzopyrene). B & C: isoflavan and neoflavonoids structures

The chemical structure of flavonoids consists of a 15-carbon skeleton, which contains two phenyl (A & B) rings and (C) heterocyclic ring. The carbon skeleton can be abbreviated C6-C3-C6. According to the IUPAC nomenclature, they can be categorized into three categories: flavonoids or bioflavonoids, isoflavonoids (3-phenyl-1,4-benzopyrone), and neoflavonoids (4-phenyl-1,2-benzopyrone) (33, 34). Flavonoids could be further classified into flavonols (e.g., quercetin, rutin), flavanones (e.g., naringenin, hesperidin), flavanols (e.g., epicatechin, gallic catechin), flavones (e.g., luteolin, apigenin), and anthocyanins (e.g., pelargonidin, malvidin). The typical examples of isoflavonoids (also called isoflavone) are genistein and daidzein; whereas, neoflavonoids occur in the form of isoflavones and are rarely found in plant foods (24). The chemical structure and types of flavonoids are shown in Figure 2.

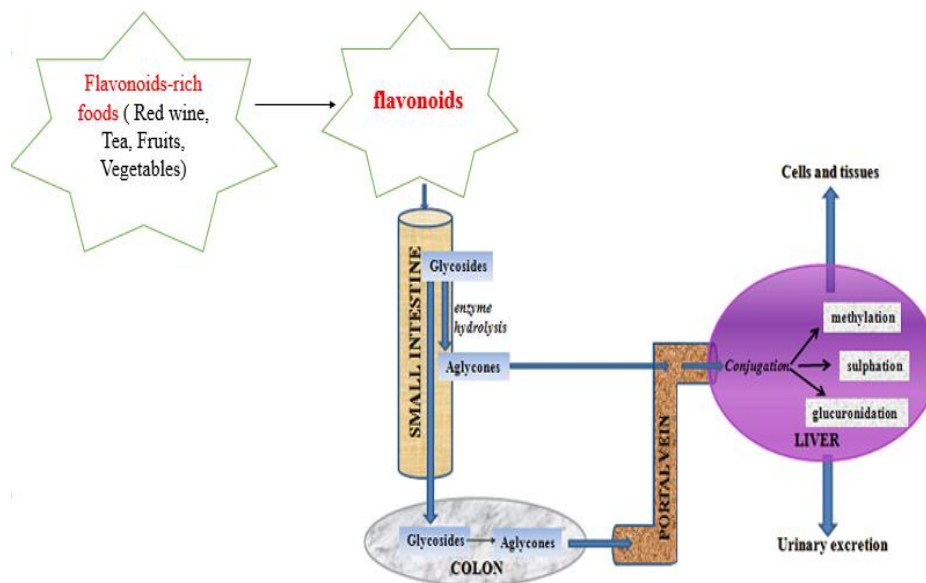
**Mechanism of action of FRF**

Significant work has been conducted in recent years to explore the working mechanisms by which FRF exert their effects against pathological conditions. Most of the investigations have communicated that beneficial activity of these foods from anti-mutagenicity to anti-aging are attributed to antioxidant potency of flavonoids (38, 39). These flavonoids exert their anti-oxidative action in several ways including direct trapping and scavenging of free radicals, decreasing leukocyte immobilization, and regulation of nitric

oxide and xanthine oxidase activity (40, 41). Several flavonoids, including quercetin, have been reported for the reduction in ischemia-reperfusion injury by interfering with inducible nitric-oxide synthase activity (42). Nitric oxide itself can be viewed as a radical, and it was reported that nitric oxide molecules are directly scavenged by flavonoids. Therefore, it has been speculated that nitric oxide scavenging plays a role in the therapeutic effects of flavonoids (41). The significant effects of flavonoids are due to radical scavenging but another possible mechanism by which flavonoids act is through interaction with various enzyme systems such as superoxide dismutase, catalase, and glutathione peroxidase (41). Furthermore, *in vitro* studies have declared the anti-proliferative activity of flavonoids through inhibition of polyamine biosynthesis and signal transduction enzymes like protein tyrosine kinase (PTK), protein kinase C (PKC) and phosphoinositide 3-kinases (PI3K), induction of apoptosis and cell cycle arrest at G1/G2, differentiation of transformed cells, and rehabilitation of cellular homeostasis (43, 44).

**Pharmacokinetics of flavonoids**

The absorption, distribution, and biotransformation of flavonoids from FRF are crucial for their biological effects against lifespan-shortening degenerative diseases as shown in Figure 3. The majority of flavonoids found in FRF are in the form of  $\beta$ -glycosides, and hydrolysis of the glycoside moiety is an essential



**Figure 3.** Pharmacokinetics of dietary flavonoids

step for absorption. Hydrolysis occurs in the cecum and colon by enterobacteria, aglycones are absorbed by gut epithelial cells and entered the circulation to metabolize in the liver (45). It is also reported that lactase-phlorizin hydrolase (LPH, EC 3.2.1.62), which is found on the brush border of the mammalian small intestine for the breakdown of lactose, is responsible for hydrolysis of flavonoids (46). By using healthy ileostomy volunteers in an experiment, it is demonstrated that sodium-dependent glucose transporter-1 (SGLT-1) is responsible for the transport of quercetin glucosides from intestinal epithelial cells (46).

Conversely, in another experiment, it was claimed that quercetin glucosides are entirely hydrolyzed to their aglycone form before transport from the intestinal tract in ileostomy patients (47). Overall, the intestinal absorption, distribution, and metabolism of flavonoids are not well elucidated at present; thus, the events in the intestinal tract should be clarified to observe the desirable health effects of FRF. Moreover, research on the mechanisms for aglycone transfer across the gut wall need to be illuminated.

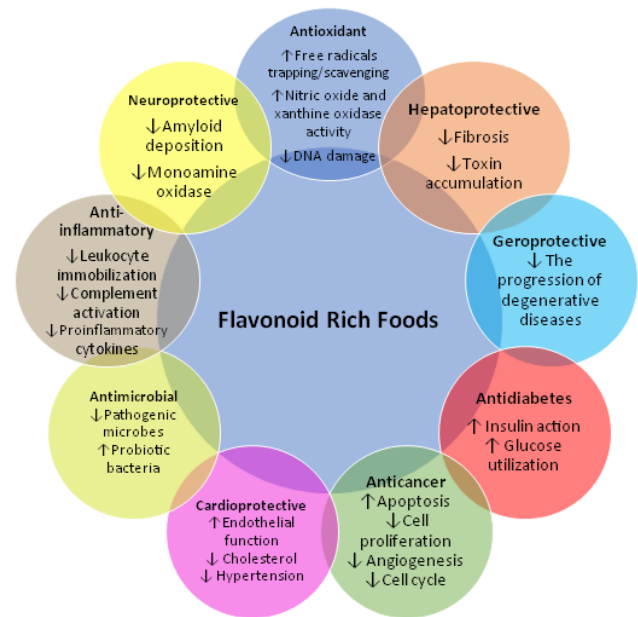
It has been observed that bioavailability of flavonoids is very low, i.e., 2–20% on an intake of FRF like vegetables, fruit juices, and beverages (48). Therefore, this area of research needs more attention to find possible ways for enhancement of flavonoids absorption.

Food matrix also plays a vital role in the bioavailability and pharmacokinetics of flavonoids from FRF. A liquid matrix yields a faster absorption rate and higher peak plasma concentrations than a solid matrix, whereas aglycones in fermented foods are absorbed more rapidly than glucoside conjugates (49). Dietary fat (3–5 g/meal) is also reported as an enhancer for the absorption of phytochemicals from vegetables in children (50).

It is also well established that processing (mechanical or heat treatment) could increase the biological activity of FRF by promoting the bioaccessibility of polyphenols in food matrix (51). Tomato polyphenols (e.g., naringenin) are trapped in the cutin matrix of the membrane of the ripe fruit where it strongly interacts with insoluble polyesters. Mechanical and heat treatments may break the interactions, thus improving flavonoids bioaccessibility *in vivo* (52). Nevertheless, a few studies have reported the adverse effect of processing on garlic's ability to alter the bioactivation of a known experimental mammary carcinogen (53). This might indicate the different response of each FRF to processing. Thus, studies are needed to unravel the individual response of processing for all FRF with significant insight into processing time. Some creative tools are required to explore the exact relationship of FRF with other components of diet and food matrix to make their possible synergisms and to avoid antagonistic combinations to enhance their bio-functionality against lifespan-shortening complications.

### Pharmacological values and therapeutic properties

At the end of the 20<sup>th</sup> century, it was concluded for the first time that traditional Mediterranean diet meets several important criteria (Figure 4), which is responsible for extended lifespan (54) and low incidence of fatal degenerative diseases like cardiovascular diseases, cancer, and diabetes (55, 56). This conclusion



**Figure 4.** The schematic overview of the therapeutic properties of flavonoid-rich foods

was drawn by intriguing evidence accumulated over the last three decades that supported the Mediterranean diet having a high amount of FRF, i.e., olive oil, legumes, cereals, vegetables, and fruits for these effects. In Greece, in addition to high vegetable diet, the wild edible greens are eaten frequently in the form of pies and salads and contain a very high concentration of flavonoids (more than those in torn or wine) as reported by a study (54). The antioxidants found in these foods (FRF) such as polyphenolic flavonoids correct the free radicals generated in all cells from normal oxidative reactions, which if left uncorrected may damage cellular proteins, lipids, and nucleic acids, resulting in the onset of lifespan-shortening diseases (Table 1).

Fatal degenerative diseases including CVD, cancer, and diabetes are the primary health burdens, which reduce the average life expectancy and impair the health status. CVD is the leading cause of death followed by cancer, and both together account for almost one-half of all deaths (9) whereas, collectively all three (cardiovascular diseases, cancer, and diabetes) account for nearly two of every three persons in the USA and comprise 32% health cost of total illness costs (11). A summary of studies indicating the effects of FRF in reducing mortalities have been summarized in Table 2.

### Protective effect against diabetes

Diabetes is an epidemic health concern in both developed and developing countries, characterized by impaired insulin production and function. In the United States, 7.8% of the population have diabetes, 35.4% have impaired fasting glucose (IFG) levels, 15.4% have impaired glucose tolerance (IGT), and 40.1% have pre-diabetes (IFG, IGT, or both) (57), while in China 9.7% and 15.5% of people are positive for diabetes and pre-diabetes, respectively (58). Diabetes could reduce life expectancy up to 15 years and account as a significant cause of mortalities in both developed and developing countries (59). It is also associated with other chronic



diseases like stroke, heart diseases, nervous system disorders, kidney diseases, and vision problems. Its prevention is more accessible than a cure. In randomized trials, it was consistently observed that increased physical activity and dietary manipulation are the ideal approaches to prevent diabetes (56).

Numerous physiological studies had declared that free radicals might contribute to the autoimmune destruction of pancreatic  $\beta$  cells, leading to diabetes (60), and may impair insulin action (61). Fruits, vegetables, and whole grains possess strong scavenging ability against these radicals resulting in reduced risk of type I and type II diabetes mellitus (62, 63). It was also

hypothesized that plant-derived foods like fruits and vegetables are low in carbohydrate contents, therefore could prevent the rise of blood sugars (56). Other dietary factors that have been related to reducing the risk of type II diabetes include coffee (64), berries (63), and tea (65). High intakes of quercetin and myricetin, mainly from dietary consumption of apples and berries also associated with reduced risk of type 2 diabetes (20). Among European adult persons who drank coffee frequently ( $\geq 7$  cups/d) had a 29% to 52% reduced risk for diabetes compared with those who drank less coffee ( $\leq 2$  cups/d or no cups/d) (64, 66). Moreover, in another study in Japan, it was reported that consumption of

**Table 1.** Effects of flavonoids rich-foods against lifespan-shortening diseases

Type of flavonoids rich-food	Biological and pharmacological effects	References
Fruits/vegetables	Anti-hypertension, reduced risk of diabetes, anti-hypercholesterolemia, anti-obesity, $\downarrow$ cardiovascular diseases, $\downarrow$ breast cancer, $\downarrow$ coronary heart diseases	(27, 28, 62, 93, 94, 95, 97)
Whole grains	Reduced risk of diabetes, anticancer	(63, 82)
Coffee	Reduced risk of type 2 diabetes	(64)
Berries	$\downarrow$ Prostate cancer, reduced risk of type 2 diabetes	(20, 63)
Green tea	Reduced risk of type 2 diabetes, $\downarrow$ blood glucose, anticancer	(65, 68, 79)
Apple	Reduced risk of type 2 diabetes, $\downarrow$ breast cancer, $\downarrow$ cardiovascular diseases	(20, 36)
Black tea	$\downarrow$ Blood glucose, $\downarrow$ total and LDL cholesterol, $\downarrow$ myocardial infarction, reduced risk of coronary heart disease	(69, 89, 121)
Onion	Antihyperglycemic effects, $\downarrow$ breast cancer	(20, 70)
Garlic	Anti-platelet aggregation, modification of LDL, antihyperglycemic effects, anticancer	(70, 82)
Cruciferous vegetables	Anticancer	(73)
Cabbage	Anticancer, $\downarrow$ vascular diseases	(20, 76)
Broccoli	Anticancer, $\downarrow$ prostate cancer	(76)
Cauliflower	Anticancer, $\downarrow$ prostate cancer	(76)
Brussels sprouts	Anticancer	(76)
Soy	Reduced risk of breast and prostate cancer	(77)
Citrus fruits	Antiproliferative, $\downarrow$ vascular diseases	(20)
Tomato	$\downarrow$ Prostate cancer	(78)
Turmeric	Anti-hepatocarcinogenesis, anticancer	(80, 81)
Ginger	Inhibit platelet aggregation, anticancer, anti-thrombotic	(82, 122)
Carrots	Anticancer	(82)
Pomegranate	Anticancer	(44)

$\downarrow$ : decreased; LDL: low-density lipoprotein

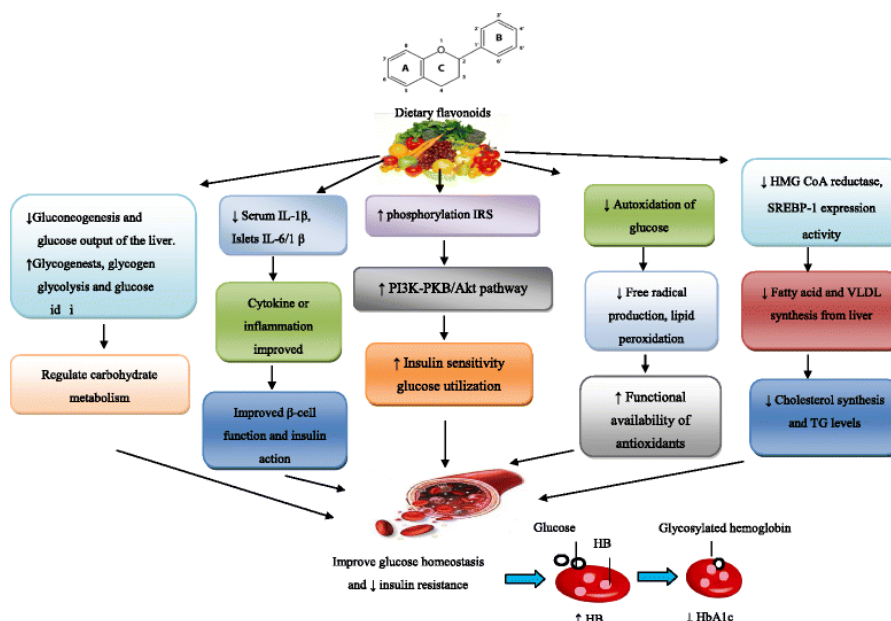
**Table 2.** Summary of studies indicating reduced mortalities by intake of flavonoids rich-foods (FRF)

Type of FRF	Dose	Type of study	Number of participants	Follow-up period	Outcomes	References
Fruit and vegetable	≥5 servings/day	Population-based cohort	71,706	13 year	53 percent higher all-cause mortality rate in those who never consumed fruits and vegetables than those who consumed five servings/day	(123)
Cruciferous vegetable	144, 232, 307, 398 & 583 g/day	Prospective cohort study	134,796	4.6 year	A dose-response pattern was evident for increasing quintiles of cruciferous vegetables intake and reduction in total mortality as well as CVD mortality	(124)
Fruit and vegetables	178.8, 316.8, 468.4 & 725.4 g/day	Population cohort study	451,151	10 year	A stronger association was observed for raw vegetable and fruit consumption and reduction in all-cause mortality as well as CVD mortality	(125)
Coffee	1 to >6 cups/day	Cohort study	41,836	15 year	1-3 cups/d significantly reduced the mortality from CVD and other inflammatory diseases in postmenopausal women	(126)
Green tea	1 to >5 cups/day	A population-based, prospective cohort study	40,530	11 year	A 16% lowered mortality from all-cause and CVD in people who consumed 5 or more cups/day than those consuming less than 1 cup/day	(127)
Fruit, vegetable, and beans	Fruit: 0.9, 2.3, 3.9 & 5.9 servings/wk; Vegetable: 1.2, 2.3, 3.4 & 5.2 servings/wk; Bean: 0.8, 1.8, 3.0 & 4.5 servings/wk	Cohort study	59,485	13 year	Fruit intake followed by vegetable and bean intake exhibited the significant inverse association for total and CVD mortality	(128)
Whole grains	≥3 servings/day	Cross-sectional study	535	3 year	A significant inverse association was recorded for whole-grain intake and mortality from CVD	(129)
Green-yellow vegetables and fruits	≤ 1, 2-4 servings/wk and one serving daily	Prospective study	38,540	18 year	Subjects had 12% lower mortality from all cancers and 20% lower mortality from lung and stomach cancer; as well as 8% lower mortality from all cancers and 25% lower mortality from liver cancer, those consuming 1 or about 1 serving/day of fruits and green-yellow vegetables, respectively, comparing with those who ate these foods once per week or less	(130)
Fruit and vegetables	0.87, 1.61, 2.31, 3.21 & 4.89 servings/day	Prospective study	6,151	13 year	Compared with the bottom fifth, highest fifth of fruit and vegetable consumers had a significantly lower all-cause, cancer, and CVD mortality	(131)
Fruit and vegetables	< 5 to ≥5 servings/d	Cohort study	501	18 year	Individuals consumed the combination of ≥5 servings/day of fruits and vegetables and ≤12 energy from saturated fat had lower all-cause (31%) and CVD (76%) mortality, compared with those consuming <5 servings/day of fruits and vegetables and >12% saturated fats	(132)
Fruit and vegetables	<1 to 8 times/day	Cohort study	9,608	19 year	Fruit and vegetable consumption ≥3 times/day as compared with <1 time/day significantly reduce stroke mortality (42%), ischemic heart disease mortality (24%), CVD mortality (27%), and all-cause mortality (15%)	(133)
Fruits	0-1, 2-3, 4-5, 6-7 times/wk	Prospective cohort study	792	26 year	Significantly lower total and CVD mortality were recorded in men with high fruit consumption. This association existed up to 16 year follow-up (at the age of 70)	(134)
Fruit, berry, and vegetables	<133, 133-214, 215-293, 294-408 & >408 g/day	Prospective cohort study	2,641	12.8 year	A significantly lower CVD, non-CVD, and all-cause mortality was observed in men with the highest consumption of fruit, vegetable, and berries	(135)

green tea, coffee, and large total caffeine was associated with a reduced risk for type 2 diabetes (67).

Furthermore, antidiabetic (blood glucose reducing) effect of green tea (68) and black tea (69) has also been observed, indicating them as potent preventive and

curative agents. Onion and garlic also have been found to have antihyperglycemic effects (70), probably based on their phytochemical contents. The antidiabetic results of FRF have been summarized in Figure 5.



**Figure 5.** The proposed antidiabetic effects of flavonoids foods. ↑ increase; ↓ decrease; AKT: v-akt murine thymoma viral oncogene homolog; HB: hemoglobin; IRS: insulin receptor substrate; HbA1c: glycated hemoglobin; IL-1β: interleukin-1 beta; HMG-CoA: 3-hydroxy-3-methyl glutaryl-coenzyme A; PI3K: phosphatidylinositol-3-kinase; TG: triglycerides; SREBP-1c: sterol regulatory element-binding protein; VLDL: very low density Lipoprotein (71)

**Protective effect against cancer**

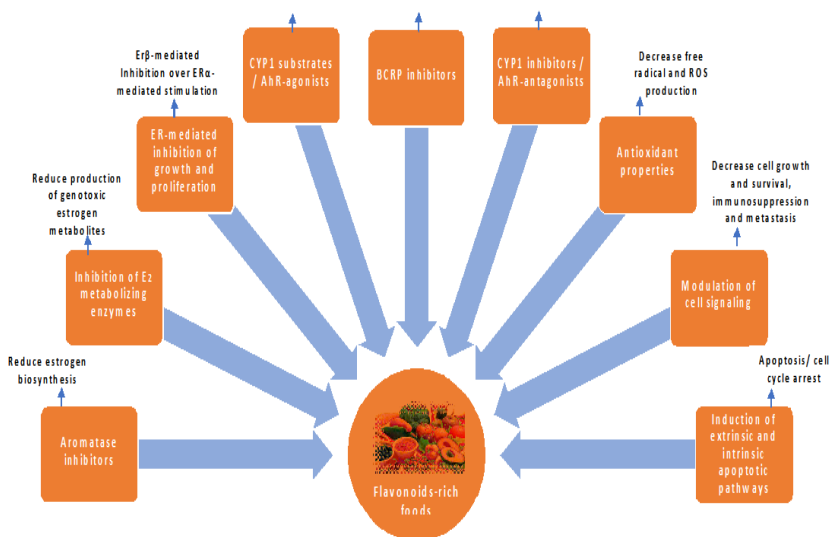
The oxidative stress from exposure to industrial chemicals, air pollutants, ionizing radiation, or ultraviolet light, might overwhelm the antioxidant system of the body and cause oxidative damage of nuclear acids and proteins, which leads to cancer initiation in addition to other degenerative diseases as explained by another study (13). Until now, no data are available assuring anticarcinogenic impacts of foods in humans. However, there are many rodent *in vivo* and *in vitro* studies confirming chemopreventive impact of certain plant foods against early stages of cancer (72). Epidemiological studies are also supporting FRF for their cancer preventing effects particularly, cruciferous vegetables due to the abundant presence of anticarcinogenic compounds such as polyphenols and isothiocyanates (73, 74). Verhoeven *et al.* (75) claimed inverse associations between intake of crucifer and the incidence of skin, pancreas, lung, prostate, bladder, stomach, colon, and thyroid cancers. In case-control studies, inverse associations between risk of cancer and intakes of broccoli, cauliflower, brussels sprouts, or cabbage were noted in 70%, 56%, 67%, and 29%, respectively (76). A Finnish study specified that risk of prostate cancer lowered at higher intakes of myricetin (from berries), and risk of breast cancer lowered at higher intakes of quercetin (from apples and onions) (20). Soybean consumption is demonstrated as a contributing factor in lowering prostate and breast cancer in Japanese men and women, respectively. This effect is considered due to isoflavone genistein that functions as an estrogen antagonist and reduces the risk of estrogen-sensitive tumors (77). Tomato and tomato products are also being investigated for their significant role in cancer chemoprevention, especially in prostate cancer (78). In another animal study, the chemopreventive effect of tea was observed for various

types of cancers and recognized for the antioxidant property of polyphenolic components known as catechins (79). Among spices, chemopreventive effects of turmeric against cancers of the skin, mouth, fore stomach, liver, and colon are well documented (80, 81). These spices contain several natural water-soluble phenolic acids and flavonoids, such as caffeic acid and quercetin that attribute to these effects by inhibition of procarcinogen activators or induction of carcinogen deactivation enzymes (72). The National Cancer Institute, after five years of research, revealed the anticancer potential of plant foods. The foods and herbs with the highest anticancer activity include garlic, soybeans, cabbage, ginger, carrots, celery, cilantro, parsley, and parsnips, while those with a modest level of cancer-protective activity include onions, citrus, turmeric, cruciferous vegetables (broccoli, brussels sprouts, cabbage, and cauliflower), tomatoes, peppers, and whole wheat (82) and their anticancer effects have been summarized in Figure 6.

**Estrogenic effect**

Some dietary flavonoids (including isoflavones and prenylflavonoids) are known as phytoestrogens, which can interact with estrogen receptors (ER) or modulate estrogen action *in vivo*. These phytoestrogens (e.g., genistein) are non-steroidal in chemical structure, but due to the presence of phenolic rings, particularly the 4'-hydroxyl, they can bind estrogen receptors. At specific concentrations, which may depend on many factors including receptor numbers, occupancy, and competing for estrogen concentration, they may antagonize and inhibit estrogen action (83).

There are inconsistent reports for the effects of phytoestrogens found in FRF on breast cancer. Setchell (83) demonstrated that soy isoflavones could stimulate the growth of ER-positive breast cancer cells through



**Figure 6.** The schematic overview of the anticancer effects of flavonoid-rich foods. CYP1: cytochrome P450; BCRP: breast cancer resistance protein; AhR: aryl hydrocarbon receptor; Erβ: estrogen receptor β; ROS: reactive oxygen species

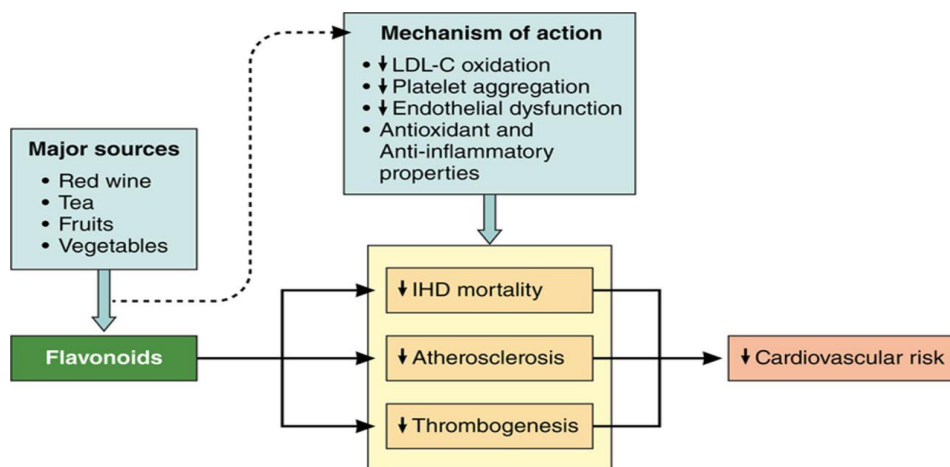
the ER signaling pathway; while, some other studies stated the putative effect of isoflavones to inhibit carcinogenesis (84, 85). These chemopreventive effects may be through impairment of protein tyrosine kinases (86) or topoisomerase II inhibition (87). From the reports in support of the positive and negative effects of phytoestrogens on breast cancer, it is indicated that more clinical studies are warranted to clarify this critical issue. Other factors like hormonal status, age, the timing of exposure, and individual metabolism should also be considered. At this stage, recommendation of nutritionists to avoid soy foods is not logical (83), and the use of phytoestrogens as supplements will be entirely acquitted.

**Protective effect against cardiovascular diseases**

Potential health benefits of chocolate (88) and tea (89) regarding heart health have been well-documented and attributed to flavonoid procyanidins, which reduce low-density lipoprotein (LDL) cholesterol by 11.1%. This LDL

is oxidized directly by peroxynitrite, which is produced by the reaction of nitric oxide with free radicals. The nitric oxide release through the constitutive nitric oxide synthase activity is essential in the maintenance of blood vessels dilation (40). Flavonoids are capable of affecting different cells included in atherosclerosis development, one of the leading reasons for cardiovascular diseases. The chemokine monocyte chemoattractant protein 1 (MCP-1) is well known to mediate macrophage recruitment to infection or inflammation sites, and direct involvement of MCP-1 on atherogenesis has been established. Furthermore, flavonoids could inhibit aggregation of TRAP-induced platelet (90) and protect endothelial cells from CD40-induced pro-inflammatory signaling as found by a study (91).

Several epidemiological studies proved that diets rich in specific antioxidants from fruits, some vegetable oils, and vegetables diminish the relative risk of premature death from CVD (25, 92) as shown in Figure 7. Findings of observational population studies support the hypothesis



**Figure 7.** The cardioprotective effect and implicated mechanisms of flavonoids. (IHD) indicates ischemic heart disease and (LDL), low-density lipoprotein cholesterol (92)



that fruit and vegetable consumption have a regulating effect on hypertension (93), hypercholesterolemia (94), and obesity (95), which are important cardiovascular risk factors. In a study related to Dietary Approaches to Stop Hypertension (DASH), it was concluded that intake of 1-2 cups of vegetable juice daily is associated with a reduction of blood pressure in subjects who were prehypertensive at the start of the trial (96). Furthermore, the decreasing trend for ischemic heart disease mortality was observed with the consumption of apples and onions (20). These findings are consistent with a large prospective study of postmenopausal women with 16 years follow-up that indicated dietary intakes of apples, pears, and red wine were associated with a lower risk of all-cause mortality, death due to coronary heart disease (CHD), and death due to CVD (36). In a cohort study of about 100,000 people, it was concluded that five servings of fruits and vegetables daily cause 28% reduction in risk of cardiovascular diseases (97), which is also in agreement with American Heart Association (AHA) recommendation for consuming at least five servings of fruits and vegetables per day (98).

### FRF as geroprotective

Aging is an essential part of life, which is directly proportional to lifespan. It may not be eliminated from life, but it could be potentially accelerated or decelerated (99). Target health issues related to the aging process include the process of oxidation, the promotion of bone health, memory retention and cognition. Decrements in motor function and memory are two main behavioral parameters that altered senescence in both humans and animals; however, they appear due to increased amounts of inflammatory markers and/or enhanced susceptibility to oxidative stress caused by reactive species from oxygen and nitrogen and subsequent induction of peroxidative reactions that result in damage to biomolecules (100).

The potential substances that can slow down the aging process, prevent premature aging, and increase life expectancy, are known as geroprotectors (99). Natural antioxidants such as flavonoids have been observed as efficient geroprotectors and lifespan extending compounds through down-regulating the progression of degenerative diseases (101), and are also called lifespan-essential ingredients (102). Therefore, the use of FRF with strong antioxidant potential may reduce age-related disorders. This may be achieved by scavenging damaging ROS, preventing the formation of lipid peroxides, protecting proteins and DNA from oxidative damage, decreasing inflammation, and protecting against ROS-mediated apoptosis (103). In addition to free radical scavenging, anti-inflammatory role of flavonoids is also crucial for overreaction of microglial cells for signals thus reducing the production of cytokines causing behavioral pathology, including cognition and restore the population of microglial brain cells to put the elder brain in the youthful state (104). It was also observed that these FRF could directly alter the neuronal communication, calcium buffering, neuroprotective stress shock proteins, and stress signaling pathways for the amelioration of age-related deficits (105).

The role of diet in minimizing the adverse effects

of aging has been extensively investigated. It has been concluded that nutritional interventions, via the polyphenolics present in plant foods like fruits and vegetables, may correct the age deficits and age-related deterioration of brain function (17, 106). Furthermore, Joseph and coworkers expressed that anthocyanin-rich fruits such as blueberry, spinach, and strawberry may help reverse the course of neuronal and behavioral aging (107). Moreover, by using transgenic mice as a model for Alzheimer's disease, the same group reported the beneficial effect of blueberry extracts on the outcome of this neurodegenerative illness (108). Studies also suggested the antiaging and brain protective role of garlic linked to dementia and Alzheimer's disease (103). In another study, green tea and its flavonoids constituents were proven for efficacy as prophylactic and neuroprotective agents against age-related neurodegenerative and neuroinflammatory diseases such as Parkinson's disease and multiple sclerosis (109). The study also confirmed that (-)-Epigallocatechin gallate inhibits lipopolysaccharide-induced microglial activation and protects against inflammation-mediated dopaminergic neuronal injury, Alzheimer's disease via modulation of cell survival/death genes, and mitochondrial function, which contributes to neuronal viability (109).

### How to get maximum health benefits from FRF

Epidemiological and clinical evidence indicate that like essential nutrients of foods (e.g., vitamins), which are vital for several physiological and pathological conditions, flavonoids are necessary for full genetically-determined lifespan through down-regulation of chronic degenerative diseases (102). Therefore, it is essential to intake the optimum amount of flavonoids by including the reference quantity of FRF on the regular menu.

Flavonoid-rich green tea, isoflavone-rich soy, flax seed, flavonol quercetin, and isoflavones are popular supplements among consumers. People are showing their interest in these supplements because they assume that they are not consuming sufficient quantities of dietary flavonoids and flavonoids supplements are devoid of toxicity because these compounds are "natural." However, there is evidence, which indicates the harmful effects of flavonoids supplements. In both animal and human studies, anti-thyroid and goitrogenic activities were observed by a high dose of green tea extracts and isoflavones (110, 111). Likewise, another adverse effect of high flavonoids doses includes inhibition of vitamin C transport, decreased trace element bioavailability, and impaired foliate uptake (112).

Feeling the gravity of this ambiguous situation it is advisable to use all flavonoid supplements with the guidelines of a health practitioner/nutritionist because these are providing hundreds of times higher doses than a regular diet and only use the optimum amounts of fresh fruits and vegetables. A large body of nutritionists, medical experts, and the American Heart Association (AHA) (97, 98) also suggests this. The instructions of regulatory bodies like WHO (world health organization), AHA, ACS (American cancer society), and ADA (American diabetes association) should strictly be followed for consumption of at least 5 servings of fresh fruits and vegetables, whole grains, legumes, flavonoid-

rich beverages, and fruit juices to get the potential health benefits from these FRF. It is also acceptable to take a moderate amount of wines, meat, milk, and other dairy products to keep in mind the possible role/effects of these substances in the food matrix. A moderate amount of heart-healthy fat (proportional monounsaturated and polyunsaturated fats) should also be included in the dietary menu for its helping role in the absorption of FRF. To ensure the maximum health benefits of FRF, healthy lifestyle, including regular exercise, should also be adapted to maintain the active physiological status of the body.

### **Biomarkers**

Intake biomarkers are useful for reflecting the amount of food or metabolites present in the body cells or fluids, but there is a need to know the best time for their measurement after consumption. Evidence is emerging that some biomarkers for cardiac health and other fatal degenerative diseases have been used successfully to differentiate disease and non-disease states and to predict the association between dietary intake and future susceptibility to infections (113, 114). For evaluation of the merits of nutritional habits, many genes, associated products, and receptors have been investigated for fatal degenerative and age-related diseases (like cancer, diabetes, Parkinson's disease, etc.) as markers of genetic susceptibility including p53, PPAR- $\alpha$ , APOA1, OB, and BCL (115, 116). Overall, authentic intake and susceptibility biomarkers are probably needed to develop a profile for an individual to approach health and longevity and significantly reduce the incidence of diet-related diseases through improved management of disease and ultimately its prevention. However, the relationship between dietary intake and lifespan related biomarkers is often highly complex.

### **Dietary/lifestyle patterns**

Dietary factors are seen as contributing to the leading causes of death of urban people, including CHD, diabetes, and certain types of cancers. Inappropriate dietary habits are responsible for poor health and reduced lifespan, which are also a major reasons for public interest in the use of alternative medicines and functional foods (117). Advances in nutritional genomics like development of the concept of gene-nutrient interaction gets significant emphasize from dietetics profession particularly the clinical dietitians to suggest appropriate foods to their patients. Unquestionably, dietary habits are not the sole determinant of disease states, because adjusting the dietary menu with FRF represents a significant way of reducing risk. Further studies are recommended to characterize the strength of diet-health association, its generalizability, the dose-response relationship, and the timing of diet for potential benefits.

Some reviews have been published on the potential role of FRF in the prevention of cancer (73, 75) and other lifestyle-related diseases like hypertension, diabetes, obesity, and aging (118). However, there is much diversity in the cultural dietary patterns and lifestyles such as the degree of physical activity and consumption of low fat and low-calorie diet, which adds complexity in understanding the exact biological role of FRF in disease prevention. Therefore, more epidemiological studies are

needed in different geographical locations to correlate the dietary patterns with cultural and individual habits and to minimize the wide divergence of the population.

### **DRI in terms of effective dosages**

Regrettably, dietary reference intake (DRI) for flavonoids could not be established until now due to insufficient data regarding flavonoid contents of different foods and variations in different varieties and cultivars and grown in different environmental conditions, production methods (organic/conventional), post-harvest processing, and storage effect on flavonoids concentration of foods (48). Another difficulty is the unavailability of authentic and certified means for determining the flavonoid contents of several foods (FRF). On the other hand, there is growing interest in developing dietary supplements containing flavonoids or flavonoid-rich foods. In such dietary supplements usually, active components (a mixture of polyphenols) from FRF were purified or concentrated to boost the antioxidant status of the consumer (119). Nevertheless, presently all dietary recommendations by scientific communities like AHA for consumption of five servings of fruits and vegetables per day are based on nutritional epidemiological surveys or *in vitro* studies but not supported by *in vivo* clinical literature. Therefore, there is a need to use the tools of food science and technology to establish a data bank for flavonoid contents of all foods with an estimated difference of biological and physical factors to determine the therapeutic/preventive dosages/amount of FRF for fatal degenerative diseases. Clinical trials are also necessary to know the relationship between specific FRF and lifespan-shortening disorders. Consequences of long-term intake of FRF should also be explored.

### **Conclusion and future perspectives**

Dietary factors play a vital role in the development and preclusion of premature aging, and some fatal degenerative diseases ultimately shorten the genetically determined lifespan. These life-threatening neurodegenerative diseases and age-related metabolic disorders are closely associated with oxidative stress produced by free radicals in the body. It has been estimated that free radicals are involved in the etiology of several (>100) human diseases and the aging process. Based on their potent antioxidant properties, flavonoids protect against these diseases and could potentially modulate life expectancy. Clinical evidence supports the health promoting, disease preventing, and life-extending effects of FRF like fruits including citrus fruits, berries, apples, vegetables particularly deep-colored green vegetables and onion, beverages such as tea and red wine, and cocoa that make these foods approach the superfoods of the millennium. It is strongly recommended that consumers eat 5 to 10 servings (one serving: about 40 g) of a wide variety of fruits and vegetables daily to reduce the risk of fatal degenerative lifespan-shortening diseases and to meet the nutrient requirements for optimum lifespan. However, one should keep in mind that FRF are not "magic bullets" for disease-free long life (120). It is only one part of a comprehensive lifestyle, which should be adjoined by physical activity (at least 30 min/day), smoking

avoidance, stress reduction, moderate consumption of alcohol, meat, eggs, and dairy products, low consumption of fat and sugars, maintaining healthy body weight (BMI: <25 kg/m<sup>2</sup>), maintenance of healthy environment, and other positive health practices. When all of these issues are addressed together, then FRF becomes part of an efficient strategy to maximize lifespan and cut the risk of lifespan-shortening degenerative diseases.

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### Conflicts of Interest

The authors declare that there are no conflicts of interest. All authors read and approved the final manuscript.

### Competing Interest

The authors declare that there are no competing financial interests.

### References

- De Leo F, Del Bosco F. Citrus flavonoids as bioactive compounds: Role, bioavailability, socio-economic impact and biotechnological approach for their modification, 9th ICABR International Conference on Agricultural Biotechnology: Ten Years Later, Ravello, Italy; 2005.
- Shen JZ, Ng LJT, Ho SW. Therapeutic potential of phytochemicals in combination with drugs for cardiovascular disorders, *Curr Pharm Des* 2017;23: 961-966.
- Kamboh AA, Arain MA, Mughal MJ, Zaman A, Arain ZM, Soomro AH. Flavonoids: health promoting phytochemicals for animal production - a review, *J Anim Health Prod* 2015; 3: 6-13.
- Kris-Etherton PM, Harris WS, Appel LJ. Omega-3 fatty acids and cardiovascular disease. *Arterioscler Thromb Vasc Biol* 2003; 23:151-152.
- Setchell KD, Radd S. Soy and other legumes: 'Bean' around a long time but are they the 'superfoods' of the millennium and what are the safety issues for their constituent phytoestrogens? *Asia Pac J Clin Nutr* 2000;9:1-10.
- Hancock RD, McDougall GJ, Stewart D. Berry fruit as 'superfood': hope or hype, *Biologist* 2007;54:73-79.
- Bishop NA, Guarente L. Genetic links between diet and lifespan: shared mechanisms from yeast to humans. *Nat Rev Genet* 2007;8:835-844.
- Tuljapurkar S, Li N, Boe C. A universal pattern of mortality decline in the G7 countries. *Nature* 2000;405:789-792.
- Murphy SL, Xu J, Kochanek KD. National vital statistics reports. National vital statistics reports. 2013;8:61.
- Olshansky SJ, Passaro DJ, Hershov RC, Layden J, Carnes BA, Brody J, et al. A potential decline in life expectancy in the United States in the 21st century, *N Engl J Med* 2005; 352:1138-1145.
- Eyre H, Kahn R, Robertson RM, Clark NG, Doyle C, Gansler T, et al. Preventing cancer, cardiovascular disease, and diabetes: a common agenda for the American Cancer Society, the American Diabetes Association, and the American Heart Association, *CA. Cancer J Clin* 2004;54:190-207.
- Kehrer JP, Klotz LO. Free radicals and related reactive species as mediators of tissue injury and disease: implications for Health. *Critic Rev Toxicol* 2015;45:765-798.
- Mojsilove G, Kuchta M. Dietary flavonoids and risk of coronary heart disease. *Physiol Res* 2001;50:529-535.
- Yang X, Jiang Y, Yang J, He J, Sun J, Chen F, et al. Prenylated flavonoids, promising nutraceuticals with impressive biological activities. *Trends Food Sci Technol* 2015; 44:93-104.
- Croft KD. The chemistry and biological effects of flavonoids and phenolic acids. *Ann N Y Acad Sci* 1998; 854:435-442.
- Catoni C, Schaefer HM, Peters A. Fruit for health: the effect of flavonoids on humoral immune response and food selection in a frugivorous bird. *Funct Ecol* 2008;22:649-654.
- Shukitt-Hale B, Galli RL, Meterko V, Carey A, Bielinski DF, McGhie T, et al. Dietary supplementation with fruit polyphenolics ameliorates age-related deficits in behavior and neuronal markers of inflammation and oxidative stress. *Age (Dordr)* 2005;27:49-57.
- Saravanan D, Thirumalai D, Asharani IV. Anti-HIV flavonoids from natural products: A systematic review. *Int J Res Pharm Sci* 2016;6:248-255.
- Liang Q, Chen H, Zhou X, Deng Q, Hu E, Zhao C, Gong X. Optimized microwave-assisted extraction combined ultrasonic pretreatment of flavonoids from *Periploca forrestii* Schltr. and evaluation of its anti-allergic activity. *Electrophoresis* 2017; 38:1113-1121.
- Knekt P, Kumpulainen J, Jarvinen R, Rissanen H, Heliovaara M, Reunanen A, et al. Flavonoid intake and risk of chronic diseases, *Am J Clin Nutr* 2002; 76:560-568.
- Aldrak N, Abudawood M, Hamed SS, Ansar S. Effect of rutin on proinflammatory cytokines and oxidative stress in toxin-mediated hepatotoxicity. *Toxin Rev* 2017;37:1-8.
- Middleton J, MD E. Biological properties of plant flavonoids: an overview, *Int J Pharm* 1996;34:344-348.
- Scalbert A, Johnson IT, Saltmarsh M. Polyphenols: antioxidants and beyond. *Am J Clin Nutr* 2005; 81:215-217.
- Kamboh AA. Flavonoid-rich Foods - Super Foods of the Millennium, LAP LAMBERT academic publishing Deutschland, Germany; 2012.
- Behzad S, Sureda A, Barreca D, Nabavi SF, Rastrelli L, Nabavi SM. Health effects of phloretin: from chemistry to medicine. *Phytochem Rev* 2017;16:527-533.
- Stanner S, Hughes J, Kelly C, Buttriss J. A review of the epidemiological evidence for the 'antioxidant hypothesis'. *Public Health Nutr* 2004;7:407-422.
- Malin AS, Qi D, Shu XO, Gao YT, Friedmann JM, Jin F, et al. Intake of fruits, vegetables and selected micronutrients in relation to the risk of breast cancer, *Int J Cancer* 2003; 105:413-418.
- Zhang X, Shu XO, Gao YT, Yang G, Li Q, Li H, et al. Soy food consumption is associated with lower risk of coronary heart disease in Chinese women. *J Nutr* 2003;133:2874-2878.
- Huang SL, Lin KC, Pan WH. Dietary factors associated with physician-diagnosed asthma and allergic rhinitis in teenagers: analyses of the first Nutrition and Health Survey in Taiwan. *Clin Exp Aller* 2001;31:259-264.
- Liu RH. Health benefits of fruit and vegetables are from additive and synergistic combinations of phytochemicals. *Am J Clin Nutr* 2003;78:517-520.
- WHO. The world health report: reducing risks, promoting healthy life, World Health Organization; 2002.
- Pallauf K, Duckstein N, Rimbach G. A literature review of flavonoids and lifespan in model organisms. *Proc Nutr Soc* 2017;76:145-162.
- Harborne JB, Williams CA. Advances in flavonoid research since 1992. *Phytochemistry* 2000; 55:481-504.
- Liu RH. Health-promoting components of fruits and vegetables in the diet. *Adv Nutr* 2013; 4:384S-392S.
- Butt MS, Imran A, Sharif MK, Ahmad RS, Xiao H, Imran M, et al. Black tea polyphenols: a mechanistic treatise. *Crit Rev Food*



- Sci Nutr 2014; 54:1002-1011.
36. Mink PJ, Scrafford CG, Barraji LM, Harnack L, Hong CP, Nettleton JA, *et al.* Flavonoid intake and cardiovascular disease mortality: a prospective study in postmenopausal women. *Am J Clin Nutr* 2007; 85:895-909.
37. Bhagwat S, Haytowitz DB, Holden JM. USDA Database for the flavonoid content of selected foods. Nutrient Data Laboratory, Beltsville Human Nutrition Research Center Agricultural Research Service U.S. Department of Agriculture 2011; 1:159.
38. Giampieri F, Forbes-Hernandez TY, Gasparrini M, Alvarez-Suarez JM, Afrin S, Bompadre S, *et al.* Strawberry as a health promoter: an evidence based review. *Food Funct* 2015; 6:1386-1398.
39. de Pascual-Teresa S, Sanchez-Ballesta MT. Anthocyanins: from plant to health. *Phytochem Rev* 2008; 7:281-299.
40. Nijveldt RJ, van Nood E, van Hoorn DE, Boelens PG, van Norren K, van Leeuwen PA. Flavonoids: a review of probable mechanisms of action and potential applications. *Am J Clin Nutr* 2001; 74:418-425.
41. Abarikwu SO, Olufemi PD, Lawrence CJ, Wekere FC, Ocholor AC, Barikuma AM. Rutin an antioxidant flavonoid, induces glutathione and glutathione peroxidase activities to protect against ethanol effects in cadmium-induced oxidative stress in the testis of adult rats. *Andrologia* 2017; 49:7.
42. Atef Y, El-Fayoumi HM, Abdel-Mottaleb Y, Mahmoud MF. Quercetin and tin protoporphyrin attenuate hepatic ischemia reperfusion injury: role of HO-1. *Naunyn-Schmiedeberg's Arch Pharmacol* 2017; 390:871-881.
43. Ren W, Qiao Z, Wang H, Zhu L, Zhang L. Flavonoids: promising anticancer agents. *Med Res Rev* 2003; 23:519-534.
44. Kawaii S, Lansky EP. Differentiation-promoting activity of pomegranate (*Punica granatum*) fruit extracts in HL-60 human promyelocytic leukemia cells. *J Med Food* 2004; 7:13-18.
45. Gonzales GB, Smaghe G, Grootaert C, Zotti M, Raes K, Camp JV. Flavonoid interactions during digestion, absorption, distribution and metabolism: a sequential structure-activity/property relationship-based approach in the study of bioavailability and bioactivity. *Drug Metab Rev* 2015; 47:175-190.
46. Day AJ, Canada FJ, Diaz JC, Kroon PA, McLauchlan R, Faulds CB, *et al.* Dietary flavonoid and isoflavone glycosides are hydrolysed by the lactase site of lactase phlorizin hydrolase. *FEBS Lett* 2000; 468:166-170.
47. Walle T, Otake Y, Walle UK, Wilson FA. Quercetin glucosides are completely hydrolyzed in ileostomy patients before absorption. *J Nut* 2000; 130:2658-2661.
48. Martin KR, Appel CL. Polyphenols as dietary supplements: a double-edged sword. *Nutr Diet Suppl* 2010; 2:1-12.
49. Cassidy A, Brown JE, Hawdon A, Faughnan MS, King LJ, Millward J, *et al.* Factors affecting the bioavailability of soy isoflavones in humans after ingestion of physiologically relevant levels from different soy foods. *J Nutr* 2006; 136:45-51.
50. Jalal F, Nesheim MC, Agus Z, Sanjur D, Habicht JP. Serum retinol concentrations in children are affected by food sources of beta-carotene, fat intake, and anthelmintic drug treatment. *Am J Clin Nutr* 1998; 68:623-629.
51. Porrini M, Riso P, Testolin G. Absorption of lycopene from single or daily portions of raw and processed tomato. *Br J Nutr* 1998; 80:353-361.
52. Bugianesi R, Salucci M, Leonardi C, Ferracane R, Catasta G, Azzini E, *et al.* Effect of domestic cooking on human bioavailability of naringenin, chlorogenic acid, lycopene and beta-carotene in cherry tomatoes. *Eur J Nutr* 2004; 43:360-366.
53. Kun S, Milner J. Heating blocks garlic's ability to suppress 7, 12-dimethylbenz (a) anthracene bioactivation. *J Nutr* 1999; 129:657-661.
54. Trichopoulou A, Vasilopoulou E. Mediterranean diet and longevity. *Br J Nutr* 2000; 84: S205-S209.
55. Barringer TA. Mediterranean diets and cardiovascular disease. *Curr Atheroscler Rep* 2001; 3:437-445.
56. Salas-Salvado J, Bullo M, Babio N, Martinez-Gonzalez MA, Ibarrola-Jurado N, Basora J, *et al.* Reduction in the incidence of type 2 diabetes with the Mediterranean diet: results of the PREDIMED-Reus nutrition intervention randomized trial. *Diabetes Care* 2011; 34:14-19.
57. Centers for Disease Control and Prevention CDC. National diabetes fact sheet: general information and national estimates on diabetes in the united states, 2007. Atlanta, GA: U.S. Department of health and human services, centre for disease control and prevention; 2008.
58. Yang W, Lu J, Weng J, Jia W, Ji L, Xiao J, *et al.* Prevalence of diabetes among men and women in China. *New Eng J Med* 2010; 362:1090-1101.
59. Gillies CL, Abrams KR, Lambert PC, Cooper NJ, Sutton AJ, Hsu RT, *et al.* Pharmacological and lifestyle interventions to prevent or delay type 2 diabetes in people with impaired glucose tolerance: systematic review and meta-analysis. *BMJ* 2007; 334:299.
60. Oberley LW. Free radicals and diabetes. *Free Radic Biol Med* 1988; 5:113-124.
61. Asmat U, Abad K, Ismail K. Diabetes mellitus and oxidative stress-a concise review. *Saudi Pharma J* 2016; 24:547-553.
62. Wang PY, Fang JC, Gao ZH, Zhang C, Xie SY. Higher intake of fruits, vegetables or their fiber reduces the risk of type 2 diabetes: A meta-analysis. *J Diabet Invest* 2016; 7:56-69.
63. Meinilä J, Valkama A, Koivusalo SB, Stach-Lempinen B, Lindström J, Kautiainen H, *et al.* Healthy Food Intake Index (HFII)-Validity and reproducibility in a gestational-diabetes-risk population. *BMC Public Health* 2016; 16:680-690.
64. Tuomilehto J, Hu G, Bidel S, Lindström J, Jousilahti P. Coffee consumption and risk of type 2 diabetes mellitus among middle-aged Finnish men and women. *JAMA* 2004; 291:1213-1219.
65. Huxley R, Lee CM, Barzi F, Timmermeister L, Czernichow S, Perkovic V, *et al.* Coffee, decaffeinated coffee, and tea consumption in relation to incident type 2 diabetes mellitus: a systematic review with meta-analysis. *Arch Intern Med* 2009; 169:2053-2063.
66. Rosengren A, Dotevall A, Wilhelmsen L, Thelle D, Johansson S. Coffee and incidence of diabetes in Swedish women: a prospective 18-year follow-up study. *J Intern Med* 2004; 255:89-95.
67. Iso H, Date C, Wakai K, Fukui M, Tamakoshi A. The relationship between green tea and total caffeine intake and risk for self-reported type 2 diabetes among Japanese adults. *Ann Intern Med* 2006; 144:554-562.
68. Saeed M, Naveed M, Arif M, Kakar MU, Manzoor R, Abd EHM, *et al.* Green tea (*Camellia sinensis*) and l-theanine: Medicinal values and beneficial applications in humans-A comprehensive review. *Biomed Pharmacol* 2017; 95:1260-1275.
69. Satoh T, Igarashi M, Yamada S, Takahashi N, Watanabe K. Inhibitory effect of black tea and its combination with acarbose on small intestinal  $\alpha$ -glucosidase activity. *J Ethnopharmacol* 2015; 161:147-155.
70. El-Demerdash FM, Yousef MI, El-Naga NI. Biochemical study on the hypoglycemic effects of onion and garlic in alloxan-induced diabetic rats. *Food Chem Toxicol* 2005; 43:57-63.
71. Vinayagam R, Xu B. Antidiabetic properties of dietary flavonoids: a cellular mechanism review. *Nutr Metab* 2015;

- 12:60-79.
72. Bahmani M, Nejad ASM, Shah NA, Shah SA, Rafieian-Kopaei M, Mahmoodnia L. Survey on ethnobotanical uses of anti-cancer herbs in Southern region of Ilam, West Iran. *J Biol Res-BollettinodellaSocietàItaliana di Biologia Sperimentale* 2017; 90:19-25.
73. Higdon JV, Delage B, Williams DE, Dashwood RH. Cruciferous vegetables and human cancer risk: epidemiologic evidence and mechanistic basis. *Pharmacol Res* 2007; 55:224-236.
74. Lam TK, Gallicchio L, Lindsley K, Shiels M, Hammond E, Tao XG, et al. Cruciferous vegetable consumption and lung cancer risk: a systematic review. *Cancer Epidemiol Biomarkers Prev* 2009; 18:184-195.
75. Verhoeven DT, Goldbohm RA, van Poppel G, Verhagen H, van den Brandt PA. Epidemiological studies on brassica vegetables and cancer risk. *Cancer Epidemiol Biomarkers Prev* 1996; 5:733-748.
76. Keck AS, Finley JW. Cruciferous vegetables: cancer protective mechanisms of glucosinolate hydrolysis products and selenium. *Integr Cancer Ther* 2004; 3:5-12.
77. Tian T, Li J, Li B, Wang Y, Li M, Ma D, et al. Genistein exhibits anti-cancer effects via down-regulating FoxM1 in H446 small-cell lung cancer cells. *Tumor Biol* 2014; 35:4137-4145.
78. Giovannucci E, Rimm EB, Liu Y, Stampfer MJ, Willett WC. A prospective study of tomato products, lycopene, and prostate cancer risk. *J Natl Cancer Inst* 2002; 94:391-398.
79. Hayakawa S, Saito K, Miyoshi N, Ohishi T, Oishi Y, Miyoshi M, et al. Anti-cancer effects of green tea by either anti- or pro-oxidative mechanisms. *Asian Pacific J Cancer Preve* 2016; 17:1649-1654.
80. Mandal S. Curcumin, a promising anti-cancer therapeutic: it's bioactivity and development of drug delivery vehicles. *Inter J Drug ResTechnol* 2017; 6:14.
81. Volate SR, Davenport DM, Muga SJ, Wargovich MJ. Modulation of aberrant crypt foci and apoptosis by dietary herbal supplements (quercetin, curcumin, silymarin, ginseng and rutin). *Carcinogenesis* 2005; 26:1450-1456.
82. Caragay AB. Cancer-preventive foods and ingredients. *Arthritis Rheum* 1992; 25:1510.
83. Setchell KD. Soy isoflavones-benefits and risks from nature's selective estrogen receptor modulators (SERMs). *J Am Coll Nutr* 2001; 20:354S-362S.
84. Harris J. The anti-inflammatory and anti-carcinogenic effects mediated by Genistein in breast cancer, Doctoral dissertation, Cardiff Metropolitan University; 2017.
85. Tse G, Eslick GD. Soy and isoflavone consumption and risk of gastrointestinal cancer: a systematic review and meta-analysis. *Eur J Nutr* 2016; 55:63-73.
86. Jung H, Ahn S, Kim BS, Shin SY, Lee YH, Lim Y. Isoflavones as modulators of adenosine monophosphate-activated protein kinase. *Appl Biol Chem* 2016; 59:217-225.
87. Salti GI, Grewal S, Mehta RR, Gupta TD, Boddie Jr AW, Constantinou AI. Genistein induces apoptosis and topoisomerase II-mediated DNA breakage in colon cancer cells. *Eur J Cancer* 2000 1;36:796-802.
88. Kwok CS, Boekholdt SM, Lentjes MA, Loke YK, Luben RN, Yeong JK, et al. Habitual chocolate consumption and risk of cardiovascular disease among healthy men and women. *Heart* 2015; 101:1279-1287.
89. Di Lorenzo A, Curti V, Tenore GC, Nabavi SM, Daglia M. Effects of tea and coffee consumption on cardiovascular diseases and relative risk factors: an update. *Curr Pharm Des* 2017; 23:2474-2487.
90. Rechner AR, Kroner C. Anthocyanins and colonic metabolites of dietary polyphenols inhibit platelet function. *Thromb Res* 2005; 116:327-334.
91. Xia M, Ling W, Zhu H, Wang Q, Ma J, Hou M, et al. Anthocyanin prevents CD40-activated proinflammatory signaling in endothelial cells by regulating cholesterol distribution. *Arterioscler Thromb Vasc Biol* 2007; 27:519-524.
92. Haseeb S, Alexander B, Baranchuk A. Wine and cardiovascular health: A comprehensive review. *Circulation* 2017; 136:1434-1448.
93. Dauchet L, Kesse-Guyot E, Czernichow S, Bertrais S, Estaquio C, Peneau S, et al. Dietary patterns and blood pressure change over 5-y follow-up in the SU.VI.MAX cohort. *Am J Clin Nutr* 2007; 85:1650-1656.
94. Ramdath DD, Padhi EM, Sarfaraz S, Renwick S, Duncan AM. Beyond the cholesterol-lowering effect of soy protein: a review of the effects of dietary soy and its constituents on risk factors for cardiovascular disease. *Nutrients* 2017; 9:324.
95. Hughes LA, Arts IC, Ambergen T, Brants HA, Dagnelie PC, Goldbohm RA, et al. Higher dietary flavone, flavonol, and catechin intakes are associated with less of an increase in BMI over time in women: a longitudinal analysis from the Netherlands Cohort Study. *Am J Clin Nutr* 2008; 88:1341-1352.
96. Shenoy SF, Kazaks AG, Holt RR, Chen HJ, Winters BL, San Khoo C, et al. The use of a commercial vegetable juice as a practical means to increase vegetable intake: a randomized controlled trial. *Nutr J* 2010; 9:38-48.
97. Hung HC, Joshipura KJ, Jiang R, Hu FB, Hunter D, Smith-Warner SA, et al. Fruit and vegetable intake and risk of major chronic disease. *J Natl Cancer Inst* 2004; 96:1577-1584.
98. Krauss RM, Eckel RH, Howard B, Appel LJ, Daniels SR, Deckelbaum RJ, et al. AHA scientific statement: AHA dietary guidelines Revision 2000: A statement for healthcare professionals from the nutrition committee of the American Heart Association. *J Nutr* 2001; 131:132-146.
99. Anisimov VN. Life span extension and cancer risk: myths and reality. *Exp Gerontol* 2001; 36:1101-1136.
100. Ferrari CK. Functional foods, herbs and nutraceuticals: towards biochemical mechanisms of healthy aging. *Biogerontology* 2004; 5:275-289.
101. Koltover V. Antioxidant biomedicine: from free radical chemistry to systems biology mechanisms. *Russ Chem Bull* 2010; 59:37-42.
102. Williamson G, Holst B. Dietary reference intake (DRI) value for dietary polyphenols: are we heading in the right direction? *Br J Nutr* 2008; 99:55-58.
103. Grosso G, Estruch R. Nut consumption and age-related disease. *Maturitas* 2016; 84:11-16.
104. Jang S, Johnson RW. Can consuming flavonoids restore old microglia to their youthful state? *Nutr Rev* 2010; 68:719-728.
105. Shukitt-Hale B, Lau FC, Joseph JA. Berry fruit supplementation and the aging brain. *J Agric Food Chem* 2008; 56:636-641.
106. Shukitt-Hale B, Carey A, Simon L, Mark DA, Joseph JA. Effects of Concord grape juice on cognitive and motor deficits in aging. *Nutrition* 2006; 22:295-302.
107. Joseph JA, Shukitt-Hale B, Denisova NA, Bielinski D, Martin A, McEwen JJ, et al. Reversals of age-related declines in neuronal signal transduction, cognitive, and motor behavioral deficits with blueberry, spinach, or strawberry dietary supplementation. *J Neurosci* 1999; 19:8114-8121.
108. Joseph JA, Denisova NA, Arendash G, Gordon M, Diamond D, Shukitt-Hale B, et al. Blueberry supplementation enhances signaling and prevents behavioral deficits in an Alzheimer disease model. *Nutr Neurosci* 2003; 6:153-162.
109. Sutherland BA, Rahman RM, Appleton I. Mechanisms of action of green tea catechins, with a focus on ischemia-induced neurodegeneration. *J Nutr Biochem* 2006; 17:291-306.
110. Milerova J, Cerovska J, Zamrazil V, Bilek R, Lapcik O, Hampl



- R. Actual levels of soy phytoestrogens in children correlate with thyroid laboratory parameters. *Clin Chem Lab Med* 2006; 44:171-174.
111. Chandra AK, De N. Goitrogenic/antithyroidal potential of green tea extract in relation to catechin in rats. *Food Chem Toxicol* 2010; 48:2304-2311.
112. Egert S, Rimbach G. Which sources of flavonoids: complex diets or dietary supplements? *Adv Nutr* 2011; 2:8-14.
113. Mateos R, Lecumberri E, Ramos S, Goya L, Bravo L. Determination of malondialdehyde (MDA) by high-performance liquid chromatography in serum and liver as a biomarker for oxidative stress: Application to a rat model for hypercholesterolemia and evaluation of the effect of diets rich in phenolic antioxidants from fruits. *J Chromatogr B Analyt Technol Biomed Life Sci* 2005; 827:76-82.
114. Medina-Remón A, Barrionuevo-González A, Zamora-Ros R, Andres-Lacueva C, Estruch R, Martínez-González MA, et al. Rapid Folin-Ciocalteu method using microtiter 96-well plate cartridges for solid phase extraction to assess urinary total phenolic compounds, as a biomarker of total polyphenols intake. *Anal Chim Acta* 2009; 634:54-60.
115. Schatzkin A. Dietary change as a strategy for preventing cancer. *Cancer Metastasis Rev* 1997; 16:377-392.
116. Ferguson LR. Nutrigenomics approaches to functional foods. *J Am Diet Assoc* 2009; 109:452-458.
117. Bigliardi B, Galati F. Innovation trends in the food industry: the case of functional foods. *Trend Food Sci Technol* 2013; 31:118-129.
118. Feeney MJ. Fruits and the prevention of lifestyle-related diseases. *Clin Exp Pharmacol Physiol* 2004; 31:11-13.
119. Weisburger JH. Chemopreventive effects of cocoa polyphenols on chronic diseases. *Exp Biol Med (Maywood)* 2001; 226:891-897.
120. Duthie GG, Gardner PT, Kyle JA. Plant polyphenols: are they the new magic bullet?. *Proc Nutr Soc* 2003; 62:599-603.
121. Gardner E, Ruxton C, Leeds A. Black tea—helpful or harmful? A review of the evidence. *Euro J Clin Nutr* 2007; 61:3-18.
122. Nurtjahja-Tjendraputra E, Ammit AJ, Roufogalis BD, Tran VH, Duke CC. Effective anti-platelet and COX-1 enzyme inhibitors from pungent constituents of ginger. *Thromb Res* 2003; 111:259-265.
123. Bellavia A, Larsson SC, Bottai M, Wolk A, Orsini N. Fruit and vegetable consumption and all-cause mortality: a dose-response analysis. *Am J Clin Nutr* 2013; 98:454-459.
124. Zhang X, Shu XO, Xiang YB, Yang G, Li H, Gao J. Cruciferous vegetable consumption is associated with a reduced risk of total and cardiovascular disease mortality. *Am J Clin Nutr* 2011;94: 240-246.
125. Leenders M, Sluijs I, Ros MM, Boshuizen HC, Siersema PD, Ferrari P. Fruit and vegetable consumption and mortality: European prospective investigation into cancer and nutrition. *Am J Epidemiol* 2013;178:590-602.
126. Andersen LF, Jacobs DR, Carlsen MH, Blomhoff R. Consumption of coffee is associated with reduced risk of death attributed to inflammatory and cardiovascular diseases in the Iowa Women's Health Study. *Am J Clin Nutr* 2006; 83:1039-1046.
127. Kuriyama S, Shimazu T, Ohmori K, Kikuchi N, Nakaya N, Nishino Y, et al. Green tea consumption and mortality due to cardiovascular disease, cancer, and all causes in Japan: the Ohsaki study. *JAMA* 2006; 296:1255-1265.
128. Nagura J, Iso H, Watanabe Y, Maruyama K, Date C, Toyoshima H, et al. Fruit, vegetable and bean intake and mortality from cardiovascular disease among Japanese men and women: the JACC Study. *Br J Nutr* 2009; 102:285-292.
129. Sahyoun NR, Jacques PF, Zhang XL, Juan W, McKeown NM. Whole-grain intake is inversely associated with the metabolic syndrome and mortality in older adults. *Am J Clin Nutr* 2006; 83:124-131.
130. Sauvaget C, Nagano J, Hayashi M, Spencer E, Shimizu Y, Allen N. Vegetables and fruit intake and cancer mortality in the Hiroshima/Nagasaki Life Span Study. *Br J Cancer* 2003; 88:689-694.
131. Genkinger JM, Platz EA, Hoffman SC, Comstock GW, Helzlsouer KJ. Fruit, vegetable, and antioxidant intake and all-cause, cancer, and cardiovascular disease mortality in a community-dwelling population in Washington County, Maryland. *Am J Epidemiol* 2004; 160:1223-1233.
132. Tucker KL, Hallfrisch J, Qiao N, Muller D, Andres R, Fleg JL. The combination of high fruit and vegetable and low saturated fat intakes is more protective against mortality in aging men than is either alone: the Baltimore Longitudinal Study of Aging. *J Nutr* 2005; 135:556-561.
133. Bazzano LA, He J, Ogden LG, Loria CM, Vupputuri S, Myers L, et al. Fruit and vegetable intake and risk of cardiovascular disease in US adults: the first National Health and Nutrition Examination Survey Epidemiologic Follow-up Study. *Am J Clin Nutr* 2002; 76:93-99.
134. Strandhagen E, Hansson PO, Bosaeus I, Isaksson B, Eriksson H. High fruit intake may reduce mortality among middle-aged and elderly men. *The Study of Men Born in 1913*. *Eur J Clin Nutr* 2000; 54:337-341.
135. Rissanen TH, Voutilainen S, Virtanen JK, Venho B, Vanharanta M, Mursu J, et al. Low intake of fruits, berries, and vegetables is associated with excess mortality in men: the Kuopio Ischaemic Heart Disease Risk Factor (KIHD) Study. *J Nutr* 2003; 133:199-204.