Review

Chemical Structures and Pharmacological Profiles of Ginseng Saponins

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Abstract: Ginseng is a group of cosmopolitan plants with more than a dozen species belonging to the genus Panax in the family Araliaceae that has a long history of use in traditional Chinese medicine (TCM). Among the bioactive constituents extracted from ginseng, ginseng saponins are a group of natural steroid glycosides and triterpene saponins found exclusively throughout the plant. Studies have shown that these ginseng saponins play a significant role in exerting multiple therapeutic effects. This review covers their chemical structure and classification, as well as their pharmacological activities, including their regulatory effects on immunomodulation, their anticancer effects, and their functions in the central nervous and cardiovascular systems. The general benefits of ginseng saponins for boosting physical vitality and improving quality of life are also discussed. The review concludes with fruitful directions for future research in the use of ginseng saponins as effective therapeutic agents.

Keywords: ginseng saponins; chemical structure; pharmacological action

1. Introduction

Archaeological evidence indicates that the pharmacological use of plants originated in the Paleolithic Age some 60,000 years ago. In the ancient Orient, the use of herbal medicines can be traced back at least 5000 years. Legendary Emperor, Shennong, classified hundreds of medicinal and poisonous herbs in Shennong’s Herbal Classic (The Classic of Herbal Medicine), the oldest pharmacopoeia in the world [1]. Ever since, humankind has been using a variety of plants as nutrients, beverages, cosmetics, dyes, and medicines to maintain health status and improve a good quality of life.

Over the centuries, botanical medicines have been widely used to treat numerous medical conditions owing to their perceived effectiveness, fewer side effects and relatively low cost. To date, traditional Chinese medicines (TCMs) are currently attracting attention worldwide as alternative and supplemental medicines [2], with ginseng, as the ‘King of all herbs’, receiving particular attention. Especially in East Asia, ginseng is deemed to be one of the most precious plants in herbal medicine.

Ginseng is a group of cosmopolitan plants with more than a dozen species belonging to the genus Panax in the family Araliaceae. Etymologically, the word Panax is derived from the Greek word ‘panacea’, meaning ‘all-healing’ and ‘longevity’. For thousands of years, ginseng has been used as a restorative of body vitality and as an adaptogen to relieve mental and physical stress and fatigue [3]. A detailed description of the medical applications of ginseng can be found in the aforementioned Shennong’s Herbal Classic, compiled around 200 AD. In summary, ginseng is one of the world’s most popular and nourishing medicinal herbs, and can be consumed for a long period of time without causing harm. Among the 17 different species assigned to the genus Panax, Panax ginseng (P. ginseng), Panax notoginseng (P. notoginseng), and Panax quinquefolius (P. quinquefolius) are the most frequently used
as medicines owing to the presence within them of ginseng saponins, which have been shown to have multiple pharmacological efficacies [4].

Ginseng has become one of the most widely consumed herbal nutritional products and alternative/complementary medicine worldwide, with a total production volume of over 80,000 tons and total sales of approximately $2.1 billion in 2013. Global demand for ginseng products, at the end of 2016 was an estimated to total 5.2 million kilograms, and the global ginseng market is also predicted to be worth some $7.51 billion by 2025 [5].

Ginseng contains various bioactive monomers, including ginseng saponins, fatty acids, polysaccharides, and mineral oils. Among its diverse constituents of ginseng, its pharmacological activities can be attributed primarily to the triterpene saponins, generally known as ginsenosides [6]. Up to now, over 100 ginseng saponins have been identified from the 11 different Panax species since their first description in the 1960s by Shibata’s group [7]. Ginseng saponins were first isolated by thin layer chromatography (TLC) and named alphabetically according to how far they migrated on the plate in a given system, e.g., Rb1, Rb2, Re, Rc, and so on [8].

Recent years have seen renewed interest in investigating the pharmacological benefits of ginseng in treating mammalian diseases. This review first describes the chemical structure of common ginseng saponins and their differing carbohydrate moieties, and then summarizes the therapeutic properties of the key saponins in terms of boosting physical vitality, as well as their regulatory effects on immunomodulation, their anticancer effects, and their functions in the central nervous and cardiovascular systems.

2. Chemical Structure and Classification of Ginseng Saponins

The basic structure of ginseng saponins comprises a hydrophobic, steroidal four-ring system [9] with a trans relationship and diverse carbohydrate moieties (i.e., glucose [glc], rhamnose [rha], xylose [xyl], and arabinose [ara]) attached to the carbon-3 (C3), carbon-6 (C6), and carbon-20 (C20) positions [10]. Table 1 summarizes the different carbohydrate moieties of the key ginseng saponins. Each ginseng saponin can be further classified as belonging to the dammarane type, including the panaxadiol-type (PPD) group, which contains a hydrogen atom at C6 (i.e., Rb1, Rb2, Rb3, Rc, Rd, Rg3, Rh2, Rs); the panaxatriol-type (PPT) group, which contains a C6 sugar side-chain (i.e., Re, Rf, Rg1, Rg2, Rh); or the oleanane type, which has two minor classes, the oleanolic acid group (i.e., Ro) and ocotillo group (i.e., F11), based on their chemical structures [11]. Figure 1 summarizes the chemical structures of the three basic ginseng saponin types.

However, ginseng saponins are not easily absorbed by the body through the intestines because of their hydrophilicity, although a small amount can be absorbed in the gastrointestinal tract after orally administration. Reports suggest that the intestinal flora can degrade saponins into secondary glycosides or aglycones through hydrolysis of the glycosidic bonds and regulate their pharmacokinetics after oral intake, which in turn further determines their pharmacological efficacies [12]. As shown in Figure 2, upon their oral consumption, ginseng saponins are partly transformed into PPD and PPT through a series of deglycosylation procedures via acid hydrolysis and intestinal bacterial actions [13]. All the metabolites, such as compound K (CK), are nonpolar compared to the parental components, which can easily be absorbed in the gastrointestinal tract and express biological actions.
Table 1. Summary of ginseng saponin carbohydrates.

| Formula   | Carbohydrate Moieties | R1 (C3) | R2 (C6) | R3 (C20) |
|-----------|-----------------------|---------|---------|----------|
| Protopanaxadiol | C_{30}H_{52}O_{3}  | H       | H       | H        |
| Ra1       | C_{26}H_{58}O_{12}  | Glc-Glc | H       | Glc-Ara(p)-Xyl |
| Ra2       | C_{26}H_{58}O_{12}  | Glc-Glc | H       | Glc-Ara(f)-Xyl |
| Ra3       | C_{26}H_{60}O_{12}  | Glc-Glc | H       | Glc-Glc-Xyl |
| Rh1       | C_{23}H_{55}O_{11}  | Glc-Glc | H       | Glc-Glc |
| Rh2       | C_{23}H_{55}O_{11}  | Glc-Glc | H       | Glc-Glc |
| Rh3       | C_{23}H_{55}O_{11}  | Glc-Glc | H       | Glc-Xyl |
| Rc        | C_{23}H_{55}O_{11}  | Glc-Glc | H       | Glc-Ara(f) |
| Rd        | C_{45}H_{85}O_{15}  | Glc-Glc | H       | Glc |
| Rg3       | C_{45}H_{85}O_{20}  | Glc-Glc | H       | H |
| Rh2       | C_{45}H_{85}O_{20}  | Glc     | H       | H |
| Rs1       | C_{45}H_{85}O_{20}  | Glc-Glc-Ac | H | Glc-Ara(p) |
| Rs2       | C_{45}H_{85}O_{20}  | Glc-Glc-Ac | H | Glc-Ara(f) |
| Rs3       | C_{45}H_{85}O_{20}  | Glc-Glc-Ac | H | H |
| Compound K | C_{36}H_{52}O_{8}  | H       | H       | Glc |

Protopanaxatriol

Re       | C_{36}H_{52}O_{4}  | H       | H       | H |
Re       | C_{36}H_{52}O_{4}  | H       | Glc-Rha | Glc |
Rf       | C_{36}H_{52}O_{4}  | H       | Glc-Glc | H |
Rg1      | C_{36}H_{52}O_{4}  | H       | Glc     | H |
Rg2      | C_{36}H_{52}O_{4}  | H       | Glc-Rha | H |
Rh1      | C_{36}H_{52}O_{4}  | H       | Glc     | H |
F1       | C_{36}H_{52}O_{4}  | H       | H       | Glc |
F3       | C_{36}H_{52}O_{4}  | H       | H       | Glc-Ara(p) |

Oleanolic acid

Ro       | C_{48}H_{74}O_{19} | GlcUA-Glc | Glc |

Ac: acetyl; Ara(f): α-L-arabinofuranosyl; Ara(p): α-L-arabinopyranosyl; Glc: β-D-glucopyranosyl; GlcUA: β-D-guluronic acid; Rha: α-L-rhamnopyranosyl; Xyl: β-D-xylopyranosyl

Figure 1. Chemical structures of three ginseng saponin types: (a) protopanaxadiol type (PPD); (b) protopanaxatriol type (PPT); and (c) oleanolic acid type.
Figure 2. Cont.
Figure 2. Main metabolic processes of (a) 20(S)-protopanaxadiol- and (b) 20(S)-protopanaxatriol-type ginseng saponins. The changed carbohydrate moieties are highlighted in red. For the daughter ginseng saponin comes from two different parental ginseng saponins, one of the changed carbohydrate moieties is highlighted in red and the other is highlighted in blue.
3. Key Saponins Found in Ginseng

Of the more than 100 different ginseng saponins, Rb1, Rb2, Rc, Rd, Re, Rf, and Rg1 are the most abundant in raw ginseng roots. Given that they comprise more than 90% of the total saponins in P. ginseng [14,15], they are also the most frequently studied. In one prior study, the crude saponin measured in 4-year-old dried ginseng accounted for over 2% of the entire plant, and the Rb1, Rf, and Rg1 content in 30 piece-graded curved ginseng samples was 2.2–4.7, 0.4–1.3, and 1.6–4.0 mg/g, respectively [16].

Of the two major ginseng saponin groups, the dammarane type dominates the oleanane type in both amount and structural varieties. The dammarane type includes Rb1 and Rg1 whose genuine aglycones, are protopanaxadiol (PPD) and protopanaxatriol (PPT), respectively, whereas the oleanane type is represented by only one minor saponin, i.e., Ro, whose aglycone is oleanolic acid.

According to a DESI-tandem mass spectrometry-based study, Rg1 and Rf are highly concentrated within the outer bark and inner core areas of the ginseng root, whereas Rd, Re, and Ra1 are distributed at high concentrations in the bark and at low concentrations in the center [17]. Rb1 is abundant in the roots, rhizomes, and root hairs of ginseng, compared with the stem and leaves [14].

4. Pharmacological Profiles and Mechanisms of Ginseng Saponins

A great number of studies have examined the various mechanisms underlying the efficacy of ginseng saponins. Some of their physiological effects may confer general health benefits. For example, there are reports of the general bolstering of the immune system, anti-inflammatory effects, anti-hepatotoxicity effects, and protective actions against mammalian tumor cell lines and non-organ-specific cancers [18]. In addition, ginseng saponins have been shown to exert various pharmacological effects with newly emerging mechanisms, including cardiovascular protective activity and anti-inflammatory, antiviral, and immunoregulatory effects. The next section provides a brief introduction to these effects and activities, and Figure 3 summarizes the pharmacological profiles and related mechanisms of ginseng saponins.

![Figure 3. Pharmacological profiles and mechanisms of ginseng saponins.](image)

Their hydrophobic steroid backbones allow ginseng saponins to intercalate into the hydrophobic interior of the lipid bilayer and further interact with the polar heads of the membrane phospholipids, which may in turn allow them to readily enter cells by simple diffusion and modulate cellular functions by binding to cytoplasmic receptors [19]. The rapid non-genomic action caused by steroids has recently been widely recognized [20]. Several mechanisms for these effects have been proposed, including changes in the membrane fluidity and activity of steroid hormones on plasma membrane receptors.
Ginseng saponins may also modify the membrane protein structure by changing the membrane dynamics and modulating the activity of ion channels, membrane-bound receptors, and enzymes. Consequently, a single ginseng saponin may be capable of reflecting its pharmacological profiles via multiple mechanisms.

4.1. General Effects of Ginseng Saponins

The classical beneficial effects of ginseng are to replenish vital energy and increase longevity. Therefore, since ancient times, ginseng has been considered a panacea that can provide eternal youth, revitalize the body and mind, and enhance physical strength and vigor. Contemporary pharmacological studies have revealed the adaptogenic activities of ginseng and its bioactive molecules acting against stress and fatigue, as well as its non-specific resistance to various pathological factors in helping to rejuvenate the body. For instance, ginseng saponins have been shown to stimulate the formation of blood vessels and improve blood circulation in the brain, thereby enhancing memory and cognitive abilities [21]. 20(R)-Rg3 may have beneficial effects against fatigue by increasing the residence time of Rg3 in the nasal cavity and enhancing the absorption through the nasal mucosa [22]. Moreover, PPD (10 mg/kg) and PPT (5 and 10 mg/kg) administration has been reported to exert anti-fatigue and anti-stress effects in mice model [23].

4.2. Effects on the Central Nervous System (CNS)

Ginseng saponins have been reported to boost memory and learning, and recent studies have shown them to play a role in the treatment of neurological diseases. For example, animal studies have demonstrated that Rb1, CK [24] and Rg1 can prevent scopolamine-induced memory deficits. In a C57BL/6 mice model, CK reversed memory impairment by inducing translocation of the nuclear factor erythroid 2-related factor (Nrf2), which further enhanced the cell stress response. Rb1 has also been found to increase the uptake of choline in the central cholinergic nerve endings [25], which have been implicated in mediating memory and learning.

The use of a combination use of ginseng saponins was associated with a prolonged drug induced sleeping time in a mouse model [26], suggesting a central nervous system (CNS)-depressing property caused by the mixture. Other studies have indicated that ginseng saponins also have neuroprotective effects. In one study, for example, Rb1 treatment inhibited GSK-3β-mediated C/EBP homologous protein (CHOP) signaling, thereby preventing neuronal apoptosis due to endoplasmic reticulum stress [27]. In another, Rg3 pretreatment attenuated Hcy-induced neurotoxicity by reducing the intracellular Ca2+ elevation via N-methyl-D-aspartate receptor (NMDAR) activation [28]. Rg3 was also demonstrated to protect the vascular endothelial cells via the estrogen receptor (ER), which is similar to the effect exerted by 17-β-estradiol. Furthermore, Rg1 also exerts an estrogenic effect by activating extracellular regulated protein kinases (ERK) and Akt signaling in an ERα phosphorylation-dependent manner, which results in memory improvement [29]. Hence, ginseng saponins may play a CNS modulatory function by interacting with hormone receptors.

Several common pharmaceuticals and supplements with antioxidant properties have been investigated as therapeutic agents for various diseases. Ginseng saponins have been proven to exert a CNS protection effect that is attributed to their antioxidant properties, which operate by increasing internal antioxidant enzymes. For example, Rb1 exerts antioxidant effects in the ischemic hippocampal neurons [30].

4.3. Anticarcinogenic Activities

The significant antitumor properties of ginseng saponins are known to be the result of their anti-inflammatory, anti-proliferation, anti-metastasis, and anti-angiogenesis effects, and reversal of multi-drug resistance. In addition, their low toxicities and few side effects make them a promising prospect for anticancer research. Ginseng saponins exert anticarcinogenic effects both in vitro and in vivo through various mechanisms, including direct cytotoxicity, growth suppression, differentiation induction, and metastasis inhibition.
For example, Rh2 has been shown to suppress tumor cell growth in breast cancer [31], prostate cancer, and leukemia [32], and to arrest melanoma cell cycle progression in the G1 phase. Orally administered and subcutaneously injected Rh2 was reported to inhibit the growth of human ovarian cancer cells transplanted into nude mice, and significantly prolong the overall survival time. Rg3 treatment in lung cancer cells was found to suppress NF-κB activity to a significant degree, thereby inhibiting tumor progression [33], and 20(S)-Rg3 to reduce human gallbladder cell (GBC) progression to a remarkable extent by activating the p53-mediated apoptotic pathway [34]. In another study, Rg3 displayed antitumor activity in LNCaP cell line via the down-regulation of the androgen receptor (AR) and other prostate cancer biomarkers [35]. Rb1 has been shown to inhibit angiogenesis via induction of the anti-angiogenic modulator pigment epithelium-derived factor (PEDF) [36]. Similarly, Rb2 and Rg3 were found to suppress angiogenesis and metastasis both in vitro and in vivo [37].

Compound K (CK), an important active metabolite of PPD-type ginseng saponin bio-transformed by intestinal microorganisms after oral administration, is drawing increasing attention recently owing to its potent anticancer benefits [38]. Research has shown that CK inhibited angiogenesis in human umbilical vein endothelial cell (HUVEC) by suppressing sphingosine-1-phosphate (S1P) induced cell migration via sphingosine kinase-d1 (SphK1) modulation [39]. Besides, CK was found to decreases basic fibroblast growth factor (FGF)-induced angiogenesis by down-regulating the phosphorylation of p38-mitogen-activated protein kinase (p38-MAPK) and Akt. A recent finding documented by our group was that CK can reduce resistance to chemotherapeutic drugs and inhibit the epithelial-to-mesenchymal transition via the Wnt/β-catenin pathway in ovarian cancer stem cells (CSC), indicating that it may be a good candidate for CSC weakening [40].

Ginseng saponins have also demonstrated strong chemoprotective and chemotherapeutic properties in a wide range of experimental studies both in vitro and in vivo. It seems that a large-scale, controlled clinical study is needed to validate these results.

4.4. Immunomodulatory Effects

The immunomodulatory and anticarcinogenic activities of ginseng saponins are often discussed together. For example, Rg1 has been reported to increase both humoral and cell-mediated immune responses [41]. Furthermore, an enhanced antitumor immunological response contributed to preventing tumor recurrence, i.e., Rh2, when used as an adjuvant, was found to trigger CD4+/CD8a+ T-lymphocyte infiltration in tumor tissues and to increase T-lymphocyte cytotoxicity with cancer chemotherapeutic drugs [42]. CK and PPT type saponins have also been shown to promote the differentiation of peripheral blood monocytes into dendritic cells (DC), suggesting a potential use in cancer immunotherapy [43]. In addition, one in vivo study found that three days of Rg1 (10 mg/kg) administration in mice prior to immunization increased the number of spleen plaque-forming cells and antigen-reactive T-lymphocytes. The ratio of T-helper cells and the natural killer activity of splenocytes were also enhanced [41].

Ginseng saponins can also be considered good drug candidates or supplements for transplantation and autoimmune disorders owing to their immunosuppression ability. For example, Rd and CK up-regulate regulatory T cell (Treg) differentiation in vitro by increasing the production of immunosuppressive cytokines TGF-β and IL-35. Moreover, it has also been reported that CK can reduce adjuvant-induced arthritis by up-regulating Treg cells and suppressing T cell activation in rats [44].

To sum up, there is growing evidence of the immunomodulatory properties of ginseng saponins. Further investigation is needed to uncover the underlying mechanism of their cancer-related immunotherapy in the tumor microenvironment.

4.5. Cardiovascular System

Cardiovascular disease (CVD) is the leading cause of death globally. In recent years, the effects of ginseng saponins on CVDs have been researched extensively because of their intrinsic properties of controlling reactive oxygen species (ROS) and nitric oxide (NO) production, and the ability to activate various receptors in endothelial cells. For example, Rb1 has been shown to preserve both
endothelial NO production in HUVEC from the toxic effects of oxidized low-density-lipoproteins and the endothelium-dependent relaxation of porcine coronary arteries exposed to homocysteine [45]. In another study, 20(S)-Rg3 enhanced endothelial NO synthase (eNOS) production by rapidly inducing the ERK/Akt signaling pathway via the activation of peroxisome proliferator-activated receptor-γ (PPAR-γ) and induced HUVEC proliferation [46]. Similarly, Rb1 acutely up-regulated eNOS by phosphorylating Ser1177 and increased NO production in human aortic endothelial cells [47].

Ginseng saponins have also been documented to exert an ion channel-regulating effect, with one study reporting Rb1 to exert an antihypertrophic effect by releasing NO and to attenuate the expression of NFAT3 and GATA4 transcription factors in cardiomyocytes by reducing the calcineurin signal transduction pathway [48]. Another study proved that Rg1 treatment to directly suppress the Ca\(^{2+}\) channel to inhibit Ca\(^{2+}\) overload in cardiomyocytes and improve the production of antioxidant enzymes in cardiac myocytes during myocardial ischemia [49]. In ischemia reperfusion treatment, Re effectively activates the cardiac K\(^{+}\) channels via eNOS activation by means of an ER\(\alpha\) related non-genomic pathway [50].

In addition, ginseng saponins have a protective effect on tissue damage, rendering them a novel therapeutic for heart failure. For example, Re has been shown to protect cardiomyocytes from oxidative injury by scavenging hydrogen peroxide and hydroxyl radicals [51]. Saponins from \(P.\) notoginseng have also been found to protect the heart against doxorubicin induced cardiotoxicity and to block monocrotaline-induced cardiac hypertrophy in rats [52].

### 4.6. Diabetes Mellitus

Owing to their effects on the endocrine system, ginseng saponins are widely used in the adjuvant treatment of diabetes and diabetic complications. For instance, Rb1 has a significant antihyperglycemic effect and increases insulin sensitivity, and is thus used clinically to treat diabetes mellitus (DM) [53]. Furthermore, Rb1 is also useful for treating obesity, which is mediated in part by reducing food intake and body weight in addition to lowering glucose [54]. The acute intraperitoneal injection of Rb1 (10 mg/kg) was demonstrated to significantly suppress food intake in rats, and chronic Rb1 treatment was found to possess no deleterious side effects in obese rats [55].

In addition, Rb1 was reported to improve insulin resistance in a high-fat diet-induced type 2 DM mouse model by reducing blood glucose [56]. Oral Re intake has also been shown to significantly decrease fasting blood glucose levels and improve glucose tolerance and systemic insulin sensitivity without affecting body weight in mice [57]. These findings indicate that ginseng saponins may be useful for improving glucose tolerance and insulin resistance in patients with type 2 diabetes.

Diabetic cardiomyopathy manifests primarily as myocardial dysfunction in the absence of other heart disease, and may eventually progress to heart failure [58]. In vitro experiments showed that Rh2 activated PPAR\(\delta\) in cardiomyocytes cultured in high glucose, in turn by inhibiting the expression of STAT3, reducing cardiac fibrosis, and protecting against diabetic cardiomyopathy [59]. In a rat model, Rh2 improved heart function in streptozotocin-induced type 1 diabetes.

### 4.7. Clinical Application

Owing to their incomplete pharmacokinetic parameters and unknown toxicities [60], individual ginseng saponins have not yet been investigated in clinical studies. However, ginseng and ginseng extract have yet have been widely used in clinical practice for thousands of years.

For example, in the context of Alzheimer’s dementia (AD), ginseng-treated patients exhibited a clinical improvement in cognitive function at 4 weeks, with continuing effects documented at 12 weeks. According to a more recent study, treatment with relatively large amounts of ginseng extract for more than 12 weeks improved the frontal assessment battery, an indicator of frontal cortical activity, in elderly patients with AD [61]. Oral administration of a ginseng saponin at 4.5g per day in AD patients had been shown to have a positive effect on the frontal lobe, which could enhance memory function [62]. Epidemiological studies of ginseng intake in 4600 cancer patients showed those who had taken ginseng
to have a lower risk of developing certain cancers (i.e., stomach, liver, and lung) than those who had not. Moreover, increased intake leads to a lower danger ratio, proving ginseng’s usefulness for primary prevention [63].

5. Other Active Ingredients in Ginseng

In addition to saponins, ginseng also comprises several other active constituents, including polysaccharides, phenolic compounds, and alkaloids. The more biologically active carbohydrates isolated from ginseng are acidic polysaccharides, known as ginsenan, which have a typical pectin structure. Polysaccharides are the most abundant components of ginseng. This class of compounds was first isolated and documented in the 1960s [64]. To date, about 80 polysaccharides isolated from ginseng have demonstrated promising pharmacological activities, including antitumor, immunomodulation, and anti-fatigue effects. However, few studies have tested them relative to ginseng saponins, and the latter also have more significant effects. A clinical trial comparing ginseng saponins and polysaccharides revealed the saponin fraction of red ginseng to decrease the augmentation index in arterial stiffness patients, but no effects were observed for the polysaccharides [65].

More than 10 phenolic compounds have been identified in P. ginseng, including elemicin and dauricine, and been found to exert various biological properties, including antitumor, antioxidant, and anti-inflammatory activities. Flavonoids are a group of polyphenolic compounds that consist of two phenyl rings and a heterocyclic ring and are universally present in plants. Flavonoids are believed to have health promoting properties due to their antioxidant activities [66]. Kaempferol is the representative flavonoid in P. ginseng.

Alkaloids, such as fumarse and girinimbi, are a minor non-saponin component of P. ginseng. Three β-carboline alkaloids were isolated from the ginseng root for the first time in 1987 [67], and two additional β-carboline alkaloids were reported the following year.

6. Conclusion and Future Prospects

Ginseng saponins are believed to inhibit ROS production, enhance immune function and CNS function, and prevent CVD. Moreover, their ability to kill cancer cells with relatively little toxicity to normal cells makes ginseng saponins an attractive candidate for drug development. However, as with other natural compounds derived from plants, their poor water solubility, permeability, absorption, and bioavailability have restricted their application. It is, therefore, hoped that more studies directed at designing and synthesizing novel derivatives and developing suitable drug delivery systems for ginseng saponins will be carried out in future.

Natural compounds are genetically encoded by their producing organisms, which means that they are the products of natural selection directed at benefiting their producers. The advantages of those compounds are thus the result of their effective engagement with their biologically relevant targets and receptors [68]. Botanic natural compounds are well-tolerated and do not exert toxic effects even at high doses. Moreover, some natural compounds may increase the efficacy of conventional therapy by mutually reinforcing the combined therapeutic actions [40,69]. Ergo, the use of phytomedicine has been valued throughout human history. Although ginseng has been widely used for thousands of years, demonstrating various health benefits for a wide range of physiological ailments, there is surprisingly little scholarly knowledge of the molecular mechanisms of action for its bioactive ingredients. There is increasing evidence to suggest that ginseng saponins are involved in the regulation of hormone receptors, which are linked to a diverse array of signaling pathways. Thus, research identifying the cellular targets responsible for the activities of ginseng saponins and characterizing their activities are warranted, which is essential for the development of multi-target phytomedicine.

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References
1. Yun, T.K. Brief Introduction of Panax Ginseng C.A. Meyer. *J. Korean Med. Sci.* **2001**, *16*, 16–18. [CrossRef] [PubMed]

2. Choi, J.G.; Eom, S.M.; Kim, J.; Kim, S.H.; Huh, E.; Kim, H.; Lee, Y.; Lee, H.; Oh, M.S.A. Comprehensive Review of Recent Studies on Herb-drug Interaction: A focus on Pharmacodynamic Interaction. *J. Altern. Complement. Med.* **2016**, *22*, 262–279. [CrossRef] [PubMed]

3. Patel, S.; Rauf, A. Adaptogenic Herb Ginseng (Panax) as Medical Food: Status Quo and Future Prospects. *Biomed. Pharmacother.* **2017**, *85*, 120–127. [CrossRef] [PubMed]

4. Xiang, Y.Z.; Shang, H.C.; Gao, X.M.; Zhang, B.L. A Comparison of the Ancient Use of Ginseng in Traditional Chinese Medicine with Modern Pharmacological Experiments and Clinical Trials. *Phyther. Res.* **2008**, *22*, 851–858. [CrossRef] [PubMed]

5. Baeg, I.H.; So, S.H. The World Ginseng Market and the Ginseng (Korea). *J. Ginseng Res.* **2013**, *37*, 1–7. [CrossRef] [PubMed]

6. Shin, B.K.; Kwon, S.W.; Park, J.H. Chemical Diversity of Ginseng Saponins from Panax Ginseng. *J. Ginseng Res.* **2015**, *39*, 287–298. [CrossRef] [PubMed]

7. Shibata, S.; Tanaka, O.; Soma, K.; Iida, Y.; Ando, T.; Nakamura, H. Studies on Saponins and Sapogenins of Ginseng the Structure of Panaxatriol. *Tetrahedron Lett.* **1965**, *42*, 207–213. [CrossRef]

8. Kaku, T.; Miyata, T.; Uruno, T.; Sako, I.; Kinoshita, A. Chemico-pharmacological Studies on Saponins of *Panax Ginseng* C. A. Meyer; I. Chemical part. *Arzneimittelforschung* **1975**, *25*, 343–347. [PubMed]

9. Lu, J.M.; Jiang, J.; Jamaluddin, M.S.; Liang, Z.; Yao, Q.; Chen, C. Ginsenoside Rb1 Blocks Ritonavir-induced Oxidative Stress and ENOS Downregulation Through Activation of Estrogen Receptor-beta and Upregulation of SOD in Human Endothelial Cells. *Int. J. Mol. Sci.* **2019**, *20*, 294. [CrossRef]

10. Shin, K.C.; Oh, D.K. Classification of Glycosidases that Hydrolyze the Specific Positions and Types of Sugar Moieties in Ginsenosides. *Crit. Rev. Biotechnol.* **2016**, *36*, 1036–1049. [CrossRef]

11. Wong, A.S.T.; Che, C.M.; Leung, K.W. Recent Advances in Ginseng as Cancer Therapeutics: A Functional and Mechanistic Overview. *Nat. Prod. Rep.* **2015**, *32*, 256–272. [CrossRef] [PubMed]

12. Wan, J.Y.; Wang, C.Z.; Liu, Z.; Zhang, Q.H.; Musch, M.W.; Bissonnette, M.; Chang, E.B.; Li, P.; Qi, L.W.; Yuan, C.S. Determination of American Ginseng Saponins and their Metabolites in Human Plasma, Urine and Feces Samples by Liquid Chromatography Coupled with Quadrupole Time-of-flight Mass Spectrometry. *J. Chromatogr. B Anal. Technol. Biomed. Life Sci.* **2016**, *15*, 1015–1016. [CrossRef]

13. Yue, P.Y.K.; Mak, N.K.; Cheng, Y.K.; Leung, K.W.; Ng, T.B.; Fan, D.T.P.; Yeung, H.W.; Wong, R.N.S. Pharmacogenomics and the Yin/Yang Actions of Ginseng: Anti-tumor, Angiomodulating and Steroid-like Activities of Ginsenosides. *Chinon Med.* **2007**, *2*, 6. [CrossRef] [PubMed]

14. Mohanan, P.; Subramaniyan, S.; Mathiyalagan, R.; Yang, D.C. Molecular Signaling of Ginsenosides Rb1, Rg1, and Rg3 and their Mode of Actions. *J. Ginseng Res.* **2018**, *42*, 123–132. [CrossRef] [PubMed]

15. Kim, I.S.; Choi, D.K.; Cho, D.Y.; Jakaria, M.; Haque, M.E.; Kim, J. Active Ginseng Components in Cognitive Impairment: Therapeutic Potential and Prospects for Delivery and Clinical Study. *Oncotarget* **2018**, *9*, 33601–33620. [CrossRef]

16. Cho, C.W.; Kim, Y.C.; Kang, J.H.; Rhee, Y.K.; Choi, S.Y.; Kim, K.T.; Lee, Y.C.; Hong, H. Do Characteristic Study on the Chemical Components of Korean Curved Ginseng Products. *J. Ginseng Res.* **2013**, *37*, 347–354. [CrossRef] [PubMed]

17. Xu, J.; Chu, Y.; Liao, B.; Xiao, S.; Yin, Q.; Bai, R.; Su, H.; Dong, L.; Li, X.; Qian, J.; et al. Panax Ginseng Genome Examination for Ginsenoside Biosynthesis. *Gigascience* **2017**, *6*, 1–15. [CrossRef]

18. Ong, Y.C.; Yong, E.L. Panax (Ginseng)-Panacea or Placebo? Molecular and Cellular Basis of its Pharmacological Activity. *Ann. Acad. Med. Singap.* **2000**, *29*, 42–46.

19. Attele, A.S.; Wu, J.A.; Yuan, C.S. Ginseng Pharmacology: Multiple Constituents and Multiple Actions. *Biochem. Pharmacol.* **1999**, *58*, 1685–1693. [CrossRef]

20. Losel, R.; Wehling, M. Nongenomic Actions of Steroid Hormones. *Nat. Rev. Mol. Cell Biol.* **2003**, *4*, 46–56. [CrossRef]
21. Oliynyk, S.; Oh, S. Actoprotective Effect of Ginseng: Improving Mental and Physical Performance. *J. Ginseng Res.* 2013, 37, 144–166. [CrossRef] [PubMed]

22. Zhang, W.; Wang, X.; Zhang, M.; Xu, M.; Tang, W.; Zhang, Y.; Lu, L.; Gao, J.; Gao, S. Intranasal Delivery of Microspheres Loaded with 20(R)-ginsenoside Rg3 Enhances Anti-fatigue Effect in Mice. *Curr. Drug Deliv.* 2017, 14, 867–874. [CrossRef] [PubMed]

23. Oh, H.A.; Kim, D.E.; Choi, H.J.; Kim, N.J.; Kim, D.H. Anti-fatigue effects of 20(S)-protopanaxadiol and 20(S)-protopanaxatriol in mice. *Biol. Pharm. Bull.* 2015, 38, 1415–1419. [CrossRef] [PubMed]

24. Seo, J.Y.; Ju, S.H.; Oh, J.; Lee, S.K.; Kim, J.S. Neuroprotective and Cognition-enhancing Effects of Compound K Isolated from Red Ginseng. *J. Agric. Food Chem.* 2016, 64, 2855–2864. [CrossRef] [PubMed]

25. Rokot, N.T.; Kairupan, T.S.; Cheng, K.C.; Runtuwene, J.; Kapantow, N.H.; Amitani, M.; Morinaga, A.; Amiani, H.; Asakawa, A.; Inui, A. A role of Ginseng and its Constituents in the Treatment of Central Nervous System Disorders. *Evid.-Based Complement. Altern. Med.* 2016, 2016, 2614742. [CrossRef] [PubMed]

26. Cha, H.Y.; Park, J.H.; Hong, J.T.; Yoo, H.S.; Song, S.; Hwang, B.Y.; Eun, J.S.; Oh, K.W. Anxiolytic-like Effects of Ginsenosides on the Elevated Plus-maze Experiment in Mice. *Biol. Pharm. Bull.* 2005, 28, 1621–1625. [CrossRef] [PubMed]

27. Liu, D.; Zhang, H.; Gu, W.; Liu, Y.; Zhang, M. Ginsenoside Rb1 Protects Hippocampal Neurons from High Glucose-induced Neurotoxicity by Inhibiting GSK3β-mediated CHOP Induction. *Mol. Med. Rep.* 2014, 9, 1434–1438. [CrossRef]

28. Kim, J.H.; Cho, S.Y.; Lee, J.H.; Jeong, S.M.; Yoon, I.S.; Lee, B.H.; Lee, J.H.; Pyo, M.K.; Lee, S.M.; Chung, K.S.; Cho, S.H.; Shin, J.S.; Kim, D.H.; Choi, J.H.; Choi, S.Y.; Rhee, Y.K.; Hong, H.D.; Lee, K.T. Ginsenoside Rb1 Inhibits Tube-like Structure Formation of Endothelial Cells by Regulating TGF-β Pigment Epithelium-derived Factor Through the Oestrogen Receptor-receptor Signaling to MAPK/ERK and P13K/Akt. *Biochim. Biophys. Acta Gen. Subj.* 2012, 1820, 453–460. [CrossRef]

29. Leung, K.W.; Cheung, L.W.T.; Pon, Y.L.; Wong, R.N.S.; Mak, N.K.; Fan, T.P.P.; Au, S.C.L.; Tombran-Tink, J.; Wong, K. Ginsenoside Rb1 Inhibits Tube-like Structure Formation of Endothelial Cells by Regulating Pigment Epithelium-derived Factor Through the Oestrogen β receptor. *Br. J. Pharmacol.* 2007, 152, 207–215. [CrossRef] [PubMed]

30. Mochizuki, M.; Yoo, Y.C.; Matsuzawa, K.; Sato, K.; Saito, I.; Tono-oaka, S.; Samukawa, K.; Azuma, I. Inhibitory Effect of Tumor Metastasis in Mice by Saponins, Ginsenoside-Rb2, 20(R)- and 20(S)-ginsenoside-Rg3, of Red Ginseng. *Biol. Pharm. Bull.* 1995, 18, 1197–1202. [CrossRef]

31. Shi, C.; Zheng, D.D.; Fang, L.; Wu, F.; Kwong, W.H.; Xu, J. Ginsenoside Rg1 Promotes Nonamylloidgenic Cleavage of APP Via Estrogen Receptor Signaling to MAPK/ERK and P13K/Akt. *Biochim. Biophys. Acta Gen. Subj.* 2012, 1820, 453–460. [CrossRef]

32. Choi, S.; Kim, T.W.; Singh, S.V. Ginsenoside Rh2-mediated G1 Phase Cell Cycle Arrest in Human Breast Cancer Cells is Caused by P15INK4B and P27INK4A-dependent Inhibition of Cyclin-dependent Kinases. *Pharm. Res.* 2009, 26, 2280–2288. [CrossRef] [PubMed]

33. Seo, J.Y.; Ju, S.H.; Oh, J.; Lee, S.K.; Kim, J.S. Neuroprotective and Cognition-enhancing Effects of Compound K Isolated from Red Ginseng. *J. Agric. Food Chem.* 2016, 64, 2855–2864. [CrossRef] [PubMed]

34. Liu, D.; Zhang, H.; Gu, W.; Liu, Y.; Zhang, M. Ginsenoside Rb1 Protects Hippocampal Neurons from High Glucose-induced Neurotoxicity by Inhibiting GSK3β-mediated CHOP Induction. *Mol. Med. Rep.* 2014, 9, 1434–1438. [CrossRef]

35. Ong, W.Y.; Farooqui, T.; Koh, H.L.; Farooqui, A.A.; Ling, E.A. Protective Effects of Ginseng on Neurological Disorders. *Front. Aging Neurosci.* 2015, 7, 129. [CrossRef]

36. Chung, K.S.; Shin, J.S.; Kim, D.H.; Choi, H.J.; Choi, S.Y.; Rhee, Y.K.; Hong, H.D.; Lee, K.T. Ginsenoside Rh2 Induces Cell Cycle Arrest and Differentiation in Human Leukemia Cells by Upregulating TGF-β Expression. *Carcinogenesis* 2013, 34, 331–340. [CrossRef] [PubMed]

37. Liu, D.; Zhang, H.; Gu, W.; Liu, Y.; Zhang, M. Ginsenoside Rb1 Protects Hippocampal Neurons from High Glucose-induced Neurotoxicity by Inhibiting GSK3β-mediated CHOP Induction. *Mol. Med. Rep.* 2014, 9, 1434–1438. [CrossRef]

38. Wang, L.; Li, X.; Song, Y.M.; Wang, B.; Zhang, F.; Yang, R.; Wang, H.Q.; Zhang, G.J. Ginsenoside Rg3 Sensitizes Human Non-small Cell Lung Cancer Cells to γ-radiation by Targeting the Nuclear Factor-xB Pathway. *Mol. Med. Rep.* 2015, 12, 609–614. [CrossRef] [PubMed]

39. Chen, X.J.; Zhang, X.J.; Shui, Y.M.; Wan, J.B.; Gao, J.L. Anticancer Activities of Protopanaxadiol- and Protopanaxatriol-type Ginsenosides and their Metabolites. *Evid.-Based Complement. Altern. Med.* 2016, 2016, 1–19. [CrossRef]

40. Shin, K.O.; Seo, C.H.; Cho, H.H.; Oh, S.; Hong, S.P.; Yoo, H.S.; Hong, J.T.; Oh, K.W.; Lee, Y.M. Ginsenoside Compound K Inhibits Angiogenesis Via Regulation of Sphingosine Kinase-1 in Human Umbilical Vein Endothelial Cells. *Arch. Pharm. Res.* 2014, 37, 1183–1192. [CrossRef]
40. Deng, S.; Wong, C.K.C.; Lai, H.-C.; Wong, A.T.S. Ginsenoside-Rb1 Targets Chemotherapy-resistant Ovarian Cancer Stem Cells Via Simultaneous Inhibition of WNT/β-catenin Signaling and Epithelial-to-Mesenchymal Transition. *Oncotarget* **2016**, *8*, 25897–25914. [CrossRef]

41. Kenarova, B.; Neychev, H.; Hadjiiivanova, C.; Petkov, V.D. Immunomodulating Activity of Ginsenoside Rg1 from Panax ginseng. *Jpn. J. Pharmacol.* **1990**, *54*, 447–454. [CrossRef] [PubMed]

42. Wang, M.; Yan, S.J.; Zhang, H.T.; Li, N.; Liu, T.; Zhang, Y.L.; Li, X.X.; Ma, Q.; Qiu, X.C.; Fan, Q.Y.; et al. Ginsenoside Rh2 Enhances the Antitumor Immunological Response of a Melanoma Mice Model. *Onco. Lett.* **2017**, *13*, 681–685. [CrossRef] [PubMed]

43. Takei, M.; Tachikawa, E.; Umeyama, A. Dendritic Cells Promoted by Ginseng Saponins Drive a Potent Th1 Polarization. *Biomark. Insights* **2013**, *3*, 269–286. [CrossRef]

44. Chen, J.; Wu, H.; Wang, Q.; Chang, Y.; Liu, K.; Song, S.; Yuan, P.; Fu, J.; Sun, W.; Huang, Q.; et al. Ginsenoside Rb1 Targets Chemotherapy-resistant Ovarian Cancer Stem Cells Via Simultaneous Inhibition of WNT/β-catenin Signaling and Epithelial-to-Mesenchymal Transition. *Oncotarget* **2016**, *8*, 25897–25914. [CrossRef]

45. He, F.; Guo, R.; Wu, S.L.; Sun, M.; Li, M. Protective Effects of Ginsenoside Rb1 on Human Umbilical Vein Endothelial Cells in vitro. *J. Cardiovasc. Pharmacol.* **2007**, *50*, 314–320. [CrossRef] [PubMed]

46. Kwok, H.H.; Guo, G.L.; Lau, J.K.; Cheng, Y.K.; Wang, J.R.; Jiang, Z.H.; Keung, M.H.; Mak, N.K.; Yue, P.Y.; Wong, R.N. Stereoisomers Ginsenosides-20(S)-Rg3 and -20(R)-Rg3 Differentially Induce Angiogenesis Through Peroxisome Proliferator-activated Receptor-gamma. *Biochem. Pharmacol.* **2012**, *83*, 893–902. [CrossRef]

47. Yu, J.; Eto, M.; Akishita, M.; Kaneko, A.; Ouchi, Y.; Okabe, T. Signaling Pathway of Nitric Oxide Production Induced by Ginsenoside Rb1 in Human Aortic Endothelial Cells: A Possible Involvement of Androgen Receptor. *Biochem. Biophys. Res. Commun.* **2007**, *353*, 764–769. [CrossRef]

48. Jiang, Q.S.; Huang, X.N.; Dai, Z.K.; Yang, G.Z.; Zhou, Q.X.; Shi, J.S.; Wu, Q. Inhibitory Effect of Ginsenoside Rb1 on Cardiac Hypertrophy Induced by Monocrotaline in Rat. *J. Ethnopharmacol.* **2007**, *111*, 567–572. [CrossRef]

49. Zhu, D.; Wu, L.; Li, C.R.; Wang, X.W.; Ma, Y.J.; Zhong, Z.Y.; Zhao, H.B.; Cui, J.; Xun, S.F.; Huang, X.L.; et al. Antioxidant and Intracellular Calcium Homeostasis. *J. Cell. Biochem.* **2010**, *108*, 117–124. [CrossRef]

50. Furukawa, T.; Bai, C.X.; Kaibara, A.; Ozaki, E.; Kawano, T.; Nakaya, Y.; Awais, M.; Sato, M.; Umezawa, Y.; Kurokawa, J. Ginsenoside Re, a Main Phytosterol of Panax Ginseng, Activates Cardiac Potassium Channels Via a Nongenomic Pathway of Sex Hormones. *Mol. Pharmacol.* **2006**, *70*, 1916–1924. [CrossRef]

51. Xie, J.T.; Shao, Z.H.; Vanden Hoek, T.L.; Chang, W.T.; Li, J.; Mendendale, S.; Wang, C.Z.; Hsu, C.W.; Becker, L.B.; Yin, J.J.; et al. Antioxidant Effects of Ginsenoside Re in Cardiomyocytes. *Eur. J. Pharmacol.* **2006**, *532*, 201–207. [CrossRef] [PubMed]

52. Qin, N.; Gong, Q.H.; Wei, L.W.; Wu, Q.; Huang, X.N. Total Ginsenosides Inhibit the Right Ventricular Hypertrophy Induced by Monocrotaline in Rat. *Biol. Pharm. Bull.* **2008**, *31*, 1530–1535. [CrossRef] [PubMed]

53. Hwang, Y.P.; Jeong, H.G. Ginsenoside Rb1 Protects Against 6-hydroxydopamine-induced Oxidative Stress by Increasing Heme Oxygenase-1 Expression Through an Estrogen Receptor-related PI3K/Akt/Nrf2-dependent Pathway in Human Dopaminergic Cells. *Toxicol. Appl. Pharmacol.* **2010**, *242*, 18–28. [CrossRef] [PubMed]

54. Shen, L.; Haas, M.; Wang, D.Q.H.; May, A.; Lo, C.C.; Obici, S.; Tso, P.; Woods, S.C.; Liu, M. Ginsenoside Rb1 Increases Insulin Sensitivity by Activating AMP-activated Protein Kinase in Male Rats. *Physiol. Rep.* **2015**, *3*, 12543. [CrossRef]

55. Xiong, Y.; Shen, L.; Liu, K.J.; Tso, P.; Xiong, Y.; Wang, G.; Woods, S.C.; Liu, M. Antiobesity and Anti-hyperglycemic Effects of Ginsenoside Rb1 in Rats. *Diabetes* **2010**, *59*, 2505–2512. [CrossRef] [PubMed]

56. Zhou, P.; Xie, W.; He, S.; Sun, Y.; Meng, X.; Sun, G.; Sun, X. Ginsenoside Rb1 as an Anti-diabetic Agent and its Underlying Mechanism Analysis. *Cells* **2019**, *8*, 204. [CrossRef] [PubMed]

57. Xie, J.T.; Mendendale, S.R.; Li, X.; Quigg, R.; Wang, X.; Wang, C.Z.; Ji, A.W.; Aung, H.H.; Rue, P.A.; Bell, G.L.; et al. Anti-diabetic Effect of Ginsenoside Re in Ob/Ob mice. *Biochim. Biophys. Acta-Mol. Basis Dis.* **2005**, *1740*, 319–325. [CrossRef] [PubMed]

58. Lee, W.S.; Kim, J. Diabetic Cardiomyopathy: Where We are and Where We are Going. *Korean J. Intern. Med.* **2017**, *32*, 404–421. [CrossRef] [PubMed]

59. Lo, S.H.; Hsu, C.T.; Niu, H.S.; Niu, C.S.; Cheng, J.T.; Chen, Z.C. Ginsenoside Rh2 Improves Cardiac Fibrosis Via PPARδ-STAT3 Signaling in Type 1-like Diabetic rats. *Int. J. Mol. Sci.* **2017**, *18*, 1364. [CrossRef]
60. Liu, P.; Xu, Y.; Yin, H.; Wang, J.; Chen, K.; Li, Y. Developmental Toxicity Research of Ginsenoside Rb1 Using a Whole Mouse Embryo Culture Model. *Birth Defects Res. Part B-Dev. Reprod. Toxicol.* 2005, 74, 207–209. [CrossRef]

61. Kim, H.J.; Jung, S.W.; Kim, S.Y.; Cho, I.H.; Kim, H.C.; Rhim, H.; Kim, M.; Nah, S.Y. Panax Ginseng as an Adjuvant Treatment for Alzheimer’s Disease. *J. Ginseng Res.* 2018, 42, 401–411. [CrossRef] [PubMed]

62. Heo, J.H.; Park, M.H.; Lee, J.H. Effect of Korean Red Ginseng on Cognitive Function and Quantitative EEG in Patients with Alzheimer’s Disease: A Preliminary Study. *J. Altern. Complement. Med.* 2015, 22, 280–285. [CrossRef] [PubMed]

63. Kamangar, F.; Gao, Y.T.; Shu, X.O.; Kahkeshani, K.; Ji, B.T.; Yang, G.; Li, H.L.; Rothman, N.; Chow, W.H.; Zheng, W. Ginseng Intake and Gastric Cancer Risk in the Shanghai Women’s Health Study Cohort. *Cancer Epidemiol. Biomark. Prev.* 2007, 16, 629–630. [CrossRef] [PubMed]

64. Ovodov, Y.S.; Solov’eva, T.F. Polysaccharides of Panax ginseng. *Chem. Nat. Compd.* 1966, 2, 242–245. [CrossRef]

65. Jovanovski, E.; Jenkins, A.; Dias, A.G.; Peeva, V.; Sievenpiper, J.; Arnason, J.T.; Rahelic, D.; Josse, R.G.; Vuksan, V. Effects of Korean Red Ginseng (Panax Ginseng C.A. Mayer) and its Isolated Ginsenosides and Polysaccharides on Arterial Stiffness in Healthy Individuals. *Am. J. Hypertens.* 2010, 23, 469–472. [CrossRef] [PubMed]

66. Kumar, S.; Pandey, A.K. Chemistry and Biological Activities of Flavonoids: An Overview. *Sci. World J.* 2013, 2013, 162750. [CrossRef]

67. Han, Y.N.; Ryu, S.Y.; Han, B.H.; Woo, L.K. Spinacine From Panax Ginseng. *Arch. Pharm. Res.* 1987, 10, 258. [CrossRef]

68. Wright, G.D. Unlocking the Potential of Natural Products in Drug Discovery. *Microb. Biotechnol.* 2019, 12, 55–57. [CrossRef]

69. Zhou, Y.; Tang, J.; Du, Y.; Ding, J.; Liu, J.Y. The Green Tea Polyphenol EGCG Potentiates the Antiproliferative Activity of Sunitinib in Human Cancer Cells. *Tumor Biol.* 2016, 37, 8555–8566. [CrossRef]