Synthesis, Spectroscopic Study and Radical Scavenging Activity of Kaempferol Derivatives: Enhanced Water Solubility and Antioxidant Activity

Sui-Ping Deng 1,*,†, Yi-Li Yang 1,†, Xing-Xing Cheng 1, Wen-Rong Li 1 and Ji-Ye Cai 1,2

1 Department of Chemistry, Jinan University, Guangzhou 510632, China; yyl0420@stu2016.jnu.edu.cn (Y.-L.Y.); 17817717548@stu2016.jnu.edu.cn (X.-X.C.); 18312736162@163.com (W.-R.L.); tjycai@jnu.edu.cn (J.-Y.C.)
2 State Key Laboratory of Quality Research in Chinese Medicines, Macau University of Science and Technology, Macau 000853, China
* Correspondence: tspdeng@jnu.edu.cn; Tel.: +86-20-8522-3569
† These authors contributed equally to this work.

Received: 29 December 2018; Accepted: 19 February 2019; Published: 23 February 2019

Abstract: Kaempferol (Kae) is a natural flavonoid with potent antioxidant activity, but its therapeutic use is limited by its low aqueous solubility. Here, a series of Kae derivatives were synthesized to improve Kae dissolution property in water and antioxidant activity. These compounds included sulfonated Kae (Kae-SO$_3$), gallium (Ga) complexes with Kae (Kae-Ga) and Kae-SO$_3$ (Kae-SO$_3$-Ga). The compound structures were characterized by high-resolution mass spectrometry (HRMS), nuclear magnetic resonance (NMR) spectroscopy, ultraviolet-visible (UV-Vis) spectroscopy, Fourier transform infrared (FT-IR) spectroscopy and thermal methods (TG/DSC). The results showed that a sulfonic group (-SO$_3$) was successfully tethered on the C3’ of Kae to form Kae-SO$_3$. And in the metal complexation, 4-CO and 3-OH of the ligand participated in the coordination with Ga(III). The metal-to-ligand ratio 1:2 was suggested for both complexes. Interestingly, Kae-SO$_3$-Ga was obviously superior to other compounds in terms of overcoming the poor water-solubility of free Kae, and the solubility of Kae-SO$_3$-Ga was about 300-fold higher than that of Kae-Ga. Furthermore, the evaluation of antioxidant activities in vitro was carried out for Kae derivatives by using α,α-diphenyl-β-picrylhydrazyl (DPPH) and 2,2’-azino-bis(3-ethylbenzo-thiazoline-6-sulfonic acid) diaminonium salt (ABTS) free radical scavenging. The results showed that Kae-SO$_3$-Ga was also optimal for scavenging free radicals in a dose-dependent manner. These data demonstrate that sulfonate kaempferol-gallium complex has a promising future as a potential antioxidant and as a potential therapeutic agent for further biomedical studies.

Keywords: kaempferol; gallium complex; water solubility; antioxidant activity; flavonoid

1. Introduction

In general, the most discussed feature of polyphenols and plant phenolics is their capability to scavenge reactive oxygen species (ROS) [1–3]. However, the phenolics cannot be synthesized by human beings, but can be gained through the diet. Vegetables and fruits are the major indispensable sources of dietary phenolics [4,5]. Excess free radicals generated in body metabolism can be sufficiently decreased by regular intake of enough vegetables and fruits [6].

Flavonoids, which are phenolic compounds, extensively spread in the plant kingdom. The four major types of flavonoids include 4-oxoflavonoids (flavones and flavonols), isoflavones, flavan-3-ol derivatives (catechin and tannins) and anthocyanins [7]. Flavonoids are composed of two aromatic rings (rings A and B) linked through an oxygenated heterocycle (ring C). This ring C is the most representative core feature of each flavonoid subfamily [8].
Flavonoids exert a wide range of benefits to human health [9,10]. They are considered as one of irreplaceable antioxidants in our daily diet because they function as singlet oxygen scavengers, free radical quencher, or chelators of metal ions, which catalyze the oxidative reactions [11]. It was suggested that free radicals and lipid peroxidation were involved in several human diseases and age-related pathologies [12]. Flavonoids can directly scavenge superoxide and peroxynitrite in an effective way, block the reaction of free radicals in the nitric oxide signaling pathway in different classes of cells [13], and hinder injury caused by free radicals.

Interestingly, the antioxidant activity of flavonoids is affected by the number of their hydroxyl groups. The performance of propyl gallate and other gallates with longer chain containing polyhydroxy groups is more preferable than the monohydric antioxidants. However, the antioxidant activity starts to go down to a lower level in structures having more than three hydroxy groups [14]. Moreover, the poor water solubility of flavonoids restricted their use. Hence, new strategies need to be developed to improve the activities of polyhydroxy flavonoids.

Flavonoids are ideal antioxidants not only due to their radical scavenging ability but also their power to chelate with metal ions [15–17]. Their metal complexes are usually found to be more active when compared with free ligands [18]. For instance, Jabeen et al. [15] studied the antioxidant activity of Cu$^{2+}$ and Fe$^{3+}$ complexes with three flavonoids (morin, quercetin and 5-hydroxyflavone) by using ultraviolet-visible (UV-Vis) spectroscopy and cyclic voltammetry. The results showed these complexes were much better antioxidants than corresponding parent flavonoids.

During the past few decades, the development of flavonoid-metal chelation has aroused a lot of interest, and continual efforts have been directed in synthesizing and characterizing novel flavonoid-metal ion complexes because of their potential applications in various fields [19]. Ravishankar et al. [20] synthesized ruthenium-conjugated chrysin analogues, which exactly displayed a 4-fold inhibition of platelet function and thrombus formation in vitro than chrysin. Medina et al. [21] synthesized apigenin-oxidovanadium compound and the complex showed moderate anti-cancer activity on lung A549 and cervix HeLa cell lines.

Gallium (Ga) demonstrated potential biomedical applications due to its interesting physical and chemical properties. Ga was reactive in electron transfer process when it was placed in electrochemical environments. Evaluation of redox reaction between gallium-indium-stannum alloy and redox active species showed that Ga was more easily oxidized by losing electrons than other metals [22]. When Ga was coordinated with a redox-active ligand, it was able to regulate the electronic nature of the ligand [23]. $^{67}$Ga (half life 78 h) and $^{68}$Ga (half life 68 min) has been successfully used as imaging agents to assess certain types of tumors, inflammation, and other lesions [24–27].

Kaempferol (Kae), well known for its prominent antioxidative activity [28], is a typical natural flavonol (Figure 1a). It is distributed in 80% of plant-based foods, including strawberries, grapes, apples, beans, kale, leek, broccoli, tomato, cabbage and tea, etc. [29]. It has been reported to have antioxidative, anti-microbial, anti-inflammatory, anti-diabetic and anti-cancer activities, as well as other pharmacologic activities against atherosclerosis and hyperlipidemia [30–32]. However, neither structural analyses nor biological activities of Ga(III) complex with Kae (Kae-Ga) have been described in detail. In this work, Kae was sulfonated in order to improve its water solubility. Besides, the coordination sites of Kae and sulfonated kaempferol (Kae-SO$_3$) with Ga(III) cation were confirmed by UV-Vis spectroscopy, Fourier transform infrared (FT-IR) spectroscopy. The complexes were characterized by $^1$H and $^{13}$C nuclear magnetic resonance (NMR) spectroscopies, mass spectrometry and thermal methods. Moreover, 2,2-diphenyl-1-picrylhydrazyl (DPPH) and 2,2’-azino-bis (3-ethylbenzo-thiazoline-6-sulfonic acid) diammonium salt (ABTS) assays were used to evaluate antioxidant activities of the Kae derivatives.
2. Results and Discussion

2.1. Sulfonation of Kae on Ring B

Kae was sulfonated in order to improve its water-solubility. After sulfonation reaction, $^{13}$C and $^1$H NMR spectra of Kae and Kae-SO$_3$ were obtained in DMSO-$d_6$ solvent, respectively. In the NMR studies, all the protons and carbons resonate at their expected frequency ranges in DMSO-$d_6$ at room temperature, properly assigned by the help of corresponding NMR experiments. A possible sulfonation site at Kae could be found by comparing the chemical shifts of carbon atoms in Table 1. The results indicated that the chemical shift of C3' of Kae and Kae-SO$_3$ was 115.91 ppm and 127.63 ppm, respectively. It showed the largest difference of about 12 ppm between Kae and Kae-SO$_3$, and it might indicate that the chemical environment of C3' changed greatly after Kae sulfonation. In addition, chemical shifts of other carbon atoms on ring B was also changed, but to a less extent about 2 ppm. Because the sulfonic group was a strong electron withdrawing group, it could affect the electron cloud density around the carbon atoms on ring B. However, the chemical shift of carbon atoms on ring A was nearly unchanged. Therefore, a sulfonic group was expected to replace the hydrogen atom of C3' (Figure 1b).

Table 1. $^{13}$C nuclear magnetic resonance (NMR) chemical shifts (ppm) for Kae and Kae-sulfonate. The differences resulting from sulfonation $\Delta = \delta$(Kae-SO$_3$) – $\delta$(Kae) are also given below.

|      | Kae   | Kae-SO$_3$ | $\Delta$ |
|------|-------|------------|--------|
| C2   | 156.65| 155.35     | −1.3   |
| C3   | 136.13| 136.56     | 0.43   |
| C4   | 176.38| 176.42     | 0.04   |
| C5   | 164.36| 164.52     | 0.16   |
| C6   | 98.67 | 98.76      | 0.09   |
| C7   | 161.18| 161.2      | 0.02   |
| C8   | 93.95 | 93.9       | −0.05  |
| C9   | 147.29| 146.36     | −0.93  |
| C10  | 103.52| 103.35     | 0.03   |
| C1'  | 122.14| 122.01     | −0.13  |
| C2'  | 129.98| 131.08     | 1.1    |
| C3'  | 115.91| 127.63     | 11.72  |
| C4'  | 159.66| 156.65     | −3.01  |
| C5'  | 115.91| 117.32     | 1.41   |
| C6'  | 129.98| 129.18     | −0.8   |
Furthermore, there was a lack of a signal of H_3 proton in $^1$H NMR spectra of Kae-SO$_3$ and Ga(III) complex with Kae-SO$_3$ (Kae-SO$_3$-Ga) in Table 2, respectively. It proved the substitution of H_3 proton by a sulfonic group in ring B of Kae molecule. Meanwhile, chemical shifts of H_2 and 4'-OH protons also changed a little after the sulfonation reaction. Because the sulfonic group was a strong electron withdrawing group, it influences electronic cloud density of nearby hydrogen atoms.

**Table 2.** Chemical shifts of hydrogen atoms in $^1$H NMR spectra for Kae, Kae-Ga, Kae-SO$_3$ and Kae-SO$_3$-Ga, respectively.

| Compound  | 3-OH  | 7-OH  | 5-OH  | 4'-OH | H$_2$  | H$_2'$ | H$_3$  | H$_3'$ | H$_8$  | H$_6$  |
|-----------|-------|-------|-------|-------|-------|-------|-------|-------|-------|-------|
| Kae       | 12.49 | 10.80 | 10.12 | 9.42  | 8.06  | 8.03  | 6.95  | 6.92  | 6.44  | 6.19  |
| Kae-Ga    | —     | 10.32 | 10.23 | 7.92  | 7.80  | 7.77  | 6.84  | 6.81  | 6.55  | 6.36  |
| Kae-SO$_3$| 12.50 | 10.99 | 10.89 | 9.61  | 8.06  | 8.42  | 6.96  | —     | 6.46  | 6.20  |
| Kae-SO$_3$-Ga | — | 11.04 | 8.55  | 8.08  | 7.79  | 7.40  | 7.24  | —     | 7.07  | 6.87  |

2.2. **UV-Vis Spectra Analysis**

The UV–Vis spectra of Kae, Kae-SO$_3$, Kae-Ga and Kae-SO$_3$-Ga in ethanol were analyzed, respectively (Figure 2). The data is listed in Table 3. It was obvious that the corresponding profiles for Kae and Kae-SO$_3$ were similar in Figure 2. The same pattern was also observed when comparing with the profiles for Kae-Ga and Kae-SO$_3$-Ga. This UV pattern similarity supported the idea that -SO$_3$ group has little effect on Kae’s UV absorption, and the chelation ability of Kae with Ga(III) was hardly affected by introducing a sulfonic functional group in the basic flavonoid skeleton.

**Figure 2.** Ultraviolet-visible (UV-Vis) spectra of (a) Kae, (b) Kae-Ga, (c) Kae-SO$_3$ and (d) Kae-SO$_3$-Ga, respectively.

**Table 3.** Ultraviolet-visible (UV-Vis) spectra data of Kae, Kae-Ga, Kae-SO$_3$ and Kae-SO$_3$-Ga, respectively.

| Compound  | Peak I (nm) | Peak II (nm) |
|-----------|-------------|--------------|
| Kae       | 368         | 266          |
| Kae-Ga    | 430         | 271          |
| Kae-SO$_3$| 368         | 270          |
| Kae-SO$_3$-Ga | 426 | 269          |

For Kae, two major characteristic bands, i.e. two absorption peaks, were observed at 368 nm and 266 nm for band I and band II, respectively (Figure 2a), which was consistent with the previous study [33]. The absorption peak at 266 nm (band II) was considered to be annotated for the absorption involving benzoyl system (ring A). The absorption peak at 368 nm (band I) was closely related to the
cinnamoyl system (ring B) of Kae. After Kae chelating with a Ga(III) ion, band II moved slightly from 266 nm to 271 nm (band IV) by red shift, which suggested that no considerable change arose in benzoyl system in the chelating reaction. However, the interaction between Kae and Ga(III) caused a significant change to the peak at 368 nm (band I). It was displaced by the peaks at ~430 nm of Kae-Ga (band III) and Kae-SO$_3$-Ga (band VII), respectively (Figure 2). A bathochromic shift of about 62 nm was clearly appeared. It confirmed that complex formation took place between Ga(III) ion and Kae ligand.

Moreover, according to the Kae structure (Figure 1a), it could be inferred that the 3-hydroxy-4-oxo part of Kae might be the initial sites involved in the metal coordination process due to the acidic nature of 3-OH and appropriate location of 4-CO. It was well known that high delocalization of oxygen electrons of 3-OH group could prompt the π electrons delocalization [17,34]. After 3-OH group binding with a Ga(III) ion, a big outstretched π bond system could be developed subsequently via electronic redistribution. Thus, an enhanced conjugative effect occurred in association with 3-OH and 4-CO in ring C [35], and it brought a direct consequence—a new ring was formed in the Ga(III) complexes with Kae-Ga and Kae-SO$_3$-Ga (Figure 1c,d). This newly-formed ring could enable these complexes to obtain additional molecular stabilization of lower energy state.

### 2.3. FT-IR Spectra Analysis

Figure 3 showed FT-IR spectra of Kae, Kae-Ga, Kae-SO$_3$ and Kae-SO$_3$-Ga, respectively. Data is listed in Table 4. For free Kae, the characteristic C=O stretching band was found at 1659.5 cm$^{-1}$. It shifted to 1615.7 cm$^{-1}$ and 1617.6 cm$^{-1}$ in the case of Kae-Ga and Kae-SO$_3$-Ga, respectively. This was because the metal coordination had a significant impact on 4-CO carbonyl group of ring C, and it confirmed the Ga(III) had bonded to 4-CO. In the spectra of Kae-Ga and Kae-SO$_3$-Ga, the presence of unique vibration at 604.0 cm$^{-1}$ and 629.9 cm$^{-1}$, respectively, represented a Ga-O stretching band, which also indicated the formation of metal complex. However, free Kae and its sulfonate exhibited no such band. What is more, a slight increase in vibrational frequency of ν(C–OH) between Kae and Kae-Ga (1380.1 cm$^{-1}$ to 1384.8 cm$^{-1}$), as well as that between Kae-SO$_3$ and Kae-SO$_3$-Ga (1378.9 cm$^{-1}$ to 1384.4 cm$^{-1}$), suggested the involving of ν(C–OH) vibration of phenolic hydroxyl, which revealed that the metal complexation almost had no impact on phenolic hydroxyl. It confirmed the Ga(III) had bonded to 3-OH of ring C. The above outcomes indicated that the Ga(III) ion probably coordinated with Kae and Kae-SO$_3$ at 3-OH and 4-CO of ring C. The presence of peaks at 1087.4 cm$^{-1}$ and 1090.0 cm$^{-1}$ belonged to S=O deformation band in the spectra of Kae-SO$_3$ and Kae-SO$_3$-Ga, respectively. This confirmed that -SO$_3$ group had bonded to Kae. The broad ν(O–H) vibrational bands near 3400 cm$^{-1}$ in all samples suggested the existence of water.

![Figure 3](image_url)  
**Figure 3.** Fourier transform infrared (FT-IR) spectra of (a) Kae, (b) Kae-Ga, (c) Kae-SO$_3$ and (d) Kae-SO$_3$-Ga, respectively.
Table 4. Frequencies of characteristic absorption bands in Fourier transform infrared (FT-IR) spectra of Kae, Kae-Ga, Kae-SO$_3$ and Kae-SO$_3$-Ga, respectively.

| Compound       | –OH | C=O  | C=C  | C–OH (Phenolic Hydroxyl) | C–O–C | O–M | –SO$_3$Na |
|----------------|-----|------|------|--------------------------|-------|-----|-----------|
| Kae            | 3386.1 | 1659.5 | 1507.3 | 1380.1            | 1176.4 | —   | —         |
| Kae-Ga         | 3434.9 | 1615.7 | 1533.3 | 1384.8            | 1170.0 | 604.0 | —         |
| Kae-SO$_3$     | 3438.9 | 1654.7 | 1502.2 | 1378.9            | 1174.2 | —   | 1087.4    |
| Kae-SO$_3$-Ga | 3438.9 | 1617.6 | 1513.9 | 1384.4            | 1197.2 | 629.9 | 1090.0    |

2.4. High-Resolution Mass Spectrometry (HRMS) Analysis

The refinement and upgrading of mass spectrometry techniques have given a considerable impetus to determine the composition, molecular weight distribution and structure of complicated materials [36]. In order to more accurately determine the structure of Ga(III) complexes, the coordination products were identified by HRMS using the positive ionization mode. In Figure 4a, the species with $m/z = 639.0034$ was assigned to a 1:2 (metal-to-ligand ratio (M:L)) complex for Kae-Ga, which was a consequence of the loss of two hydrogen atoms from two Kae molecules, and then chelating with one Ga(III) cation. In Figure 4b, the peak at $m/z = 798.9646$ represented two Kae-SO$_3$ molecules and one Ga(III) ion, which also depicted a 1:2 (M:L) complex for Kae-SO$_3$-Ga.

![Figure 4. The high-resolution mass spectra (HRMS) of the kaempferol-gallium complexes. (a) 1:2 (metal-to-ligand ratio (M:L)) Kae-Ga, (b) 1:2 (M:L) Kae-SO$_3$-Ga.](image)

2.5. $^1$H NMR Spectrometry Analysis

Roy et al. [37] successfully synthesized a luteolin-vanadium(II) complex. The results showed that a vanadium(II) cation was complexed with 4-CO and 5-OH sites of luteolin. This hydroxyl chelation site was different from that in Kae-Ga and Kae-SO$_3$-Ga, respectively. In the present work, however, the Ga(III)-binding sites on Kae were further confirmed by $^1$H NMR study (Table 2). Compared to the free Kae, the signal of 3-OH proton was absent in the metal complexes, including Kae-Ga and Kae-SO$_3$-Ga (Table 2). However, the other three hydroxyl group protons (4’-OH, 5-OH and 7-OH) remained after chelation. This indicated that the Ga(III) ion combined with Kae through the 3-OH group.
After the complexes were formed, $^1$H NMR data showed that chemical shifts of hydrogen atoms on the 3-OH had changed obviously, while those on the other hydroxyl groups changed slightly. It was probably attributed to the increase of the conjugation effect caused by the coordination when the complex was formed, and the subsequent increase of flavonoid planarity [30]. This result provided evidence that Kae successfully chelated with Ga(III) ion via 3-OH and 4-CO groups, and in the same way Kae-SO$_3$ combined with Ga(III) ion.

### 2.6. Thermal Study of the Kae-Ga Complex

With the utilization of a simultaneous thermal analyzer, the Kae-Ga sample was detected under dynamic inert atmosphere (nitrogen) and the data of differential scanning calorimetry (DSC) and thermal gravity (TG) could be simultaneously obtained. Figure 5 showed the thermal analysis (TG/DSC) of Kae-Ga with the heating rate of 20 °C·min$^{-1}$. TG and derivative thermogravimetric (DTG) plots showed that Kae-Ga exhibited a three-step degradation process. First, a slight weight loss (5.48%) was observed at 34–128 °C, and it was suggested that the complex contained two water molecules, which was 5.32% in calculation. Second, a significant weight loss (30.16%) could be seen at 297 °C, which denoted an exothermic peak. The Kae-Ga complex underwent decomposition, and was converted into carbon oxides and water. Third, in the DTG curve, the next peak in the temperature range of 844–1033 °C was related to the complete decomposition of the complex. The residue eventually turned out to be gallium oxide and remained stable.

![Figure 5. The thermogravimetric and differential scanning calorimeter (TG/DSC) curves of melting process of Kae-Ga complex in N$_2$.](image)

Through comprehensive analysis of the above three curves, the Kae-Ga complex exhibited a different degradation over temperatures of 800 °C when compared with other flavonoid-metal complexes in previous studies [38]. Kae-Ga complex decomposed finally over 900 °C, while other flavonoid complexes often below 800 °C. It might probably provide a clue that the Kae-Ga complex possessed a better stability in thermodynamics.

### 2.7. Enhanced Water-Solubility of Kae-SO$_3$ and Its Complex

Though Kae has many pharmacological activities, poor water solubility and dissolution of Kae has led to its low bioavailability, and subsequently few clinical applications. Certain techniques were thus used in previous studies to improve the aqueous solubility of Kae, such as nanoparticle and liposome delivery systems. Nanoparticle systems recently became a effective technique to significantly enhance dissolution and bioavailability of some poorly water-soluble drugs by reducing the size of the compounds from micron- to nanoscale, usually between 10 nm to 1000 nm. Lin et al. [39] demonstrated that the dissolution percentage of Kae nanoparticle (KAEN) (about 88 nm) was 139-fold higher than that of the pure Kae (about 6.2 μm) in a pH 1.2 hydrochloric acid buffer solution, and KAEN showed
better antioxidant activity than Kae in water. Dissolution enhancement was related to the reduced particle size, high encapsulation efficiency and crystal-to-amorphous transformation of Kae, as well as hydrogen bonding between Kae with Eudragit E100 (EE100) and polyvinyl alcohol (PVA) as nontoxic excipients [39].

In addition, β-cyclodextrin (β-CD) and sulfobutylether β-CD (SBE-β-CD) were useful in overcoming poor water solubility of the flavonoids by inclusion complexation with the flavonoids in their hydrophobic holes, such as has been shown for Kae, luteolin and myricetin [40]. β-CD has a hydrophilic outer surface, and sulfobutylether is a negatively charged group. These factors led to understand that SBE-β-CD was more effective in increasing the water solubility and antioxidant activity of the flavonoids when compared with β-CD. Jung et al. [40] reported that solubility of the flavonoids with SBE-β-CD was about 10- to 50-fold higher than that of flavonoids without CD.

Moreover, flavonoid sulphation or sulfonation was also an effective strategy to improve poor aqueous solubility of flavonoids by increasing flavonoid polarity. Sulphated flavonoids, such as persicarin and quercetin 3-sulphate, were often found in some specific plants. The flavonoids converted to their sulphate esters by enzymatic reactions in plant metabolism [41]. These substituent sulphate groups were negatively charged, which indicated that sulphated flavonoids had increasing polarity and water-solubility than their parent flavonoids. In fact, sulphated flavonoids showed several pharmacological effects, such as anti-inflammatory, anticoagulant and antitumor activities [41].

In the present study, however, sulfonated Kae and its complexes were successfully synthesized in vitro using a simple and quick preparation process. The solubility of kaempferol-based compounds in water at 30 °C is listed in Table 5. The aqueous solubility of free Kae and Kae-Ga was $3.95 \times 10^{-4}$ mol·L$^{-1}$ (0.0113 g/100 g soln.) and $6.22 \times 10^{-4}$ mol·L$^{-1}$ (0.0398 g/100 g soln.), respectively. The results proved the poor water solubility of free Kae and Kae-Ga, and these solubilities were of the same order of magnitude. However, the solubility of Kae-SO$_3$ was obviously elevated to $6.42 \times 10^{-2}$ mol·L$^{-1}$ (2.49 g/100 g soln.). It was almost enhanced 220-fold when compared with that of Kae. Moreover, the solubility of Kae-SO$_3$-Ga was significantly increased to $1.70 \times 10^{-1}$ mol·L$^{-1}$ (14.34 g/100 g soln.), which was almost enhanced 360-fold when compared with that of Kae-Ga. Furthermore, Kae-SO$_3$-Ga had the highest solubility among the four compounds. The change of solubility showed that the solubility of Kae increased after transforming to its sulfonate, and a coordinated Ga(III) moiety to Kae-SO$_3$ could greatly enhance water solubility further. Aromatic sulfonation has been recognized as an electrophilic substitution reaction. Sulfonate is prized for its ability to be easily soluble in water [42]. The sulfonic acid group is negatively-charged; therefore, this negatively-charged group could offer an increasing molecular polarity for Kae. The fact that Kae-SO$_3$ was more polar than Kae allowed Kae-SO$_3$ to obtain greater water solubility and dissolution. Besides, the coordinated trivalent Ga(III) imparted the complex inner part to a mono-charged ion; hence, the complex inner part still maintained polarity. In addition, Ga(III) might tune the electronic nature of the ligand to assist in improving water solubility. Thus, it could be inferred that solubility improvement of poor water-soluble flavonoids could be achieved by sulfonation of flavonoids, followed by metal complexation.

| Compound | Solubility (g/100 g soln.) | Concentration (mol·L$^{-1}$) |
|----------|---------------------------|-----------------------------|
| Kae      | 0.0113 ± 0.0026           | $3.95 \times 10^{-4}$       |
| Kae-SO$_3$ | 2.49 ± 0.20             | $6.42 \times 10^{-2}$       |
| Kae-Ga   | 0.0398 ± 0.0040           | $6.22 \times 10^{-4}$       |
| Kae-SO$_3$-Ga | 14.34 ± 0.34   | $1.70 \times 10^{-1}$       |

2.8. Antioxidant Activities of Kae, Kae-SO$_3$ and Their Complexes

The metal ion complexation with the flavonoids may affect the chemical properties of flavonoid molecules, and therefore lead to changes in the antioxidant activities. Roy et al. [37] reported
that both luteolin and luteolin-vanadium(II) complex showed antioxidant activities in a time- and dose-dependent manner by using DPPH, ABTS and ferric reducing antioxidant power (FRAP) methods. However, luteolin-vanadium(II) complex showed better antioxidant activity than luteolin. The compounds could donate electron or hydrogen atom, and react with free radicals or terminate the chain reaction.

In the present study, the results were in agreement with the previous findings that Ga(III) complexation affected the antioxidant activity of the ligands. There is not a universal method to measure antioxidant activity. Hence, two different measurements have been used to assess the antioxidant activity of Kae-based compounds.

DPPH, a stable free radical, is used widely to assess the antioxidant capacity of hydrogen donors or free radical scavengers such as plant extracts and food matrix [43]. In the radical scavenging reaction, antioxidants donate a hydrogen radical or an electron to DPPH\(^-\) and transform it into a diamagnetic molecule; thus, the antioxidative activity primarily relies on hydrogen- or electron-donating capacity. According to previous studies, Kae was a flavonoid well known for its good radical scavenging activity and antioxidant activity [44]. Therefore, DPPH radical scavenging activities of Kae, Kae-Ga, Kae-SO\(_3\), Kae-SO\(_3\)-Ga and L-Ascorbic acid (Vc) with different concentrations were measured, respectively. Water-soluble Kae-SO\(_3\), Kae-SO\(_3\)-Ga and Vc were dissolved in phosphate buffered saline (PBS) solution, while water-insoluble Kae and Kae-Ga in ethanol. The results are shown in Figure 6a and Table 6a.

![Figure 6](image)

**Figure 6.** Effects of Kae-based compounds on (a) \(\alpha,\alpha\)-diphenyl-\(\beta\)-picrylhydrazyl (DPPH) radical scavenging activity and (b) 2,2'-azino-bis (3-ethylbenzo-thiazoline-6-sulfonic acid) diammonium salt (ABTS) radical scavenging activity. L-Ascorbic acid (Vc) as standard radical scavenger. Water-soluble Kae-SO\(_3\), Kae-SO\(_3\)-Ga and Vc were dissolved in phosphate buffered saline (PBS) solution, respectively, while others in ethanol. Data was expressed as means ± SD obtained in three independent experiments. Significant difference was found from Kae derivative groups and Vc group to the control (\(p < 0.05\)).

It was observed that DPPH radical scavenging activity was increased significantly for all compounds with increase of their concentrations in the range of 0–75 \(\mu\text{mol}\cdot\text{L}^{-1}\) when compared with the control (\(p < 0.05\)). It was attributed to the scavenging ability of the Kae, Kae-Ga, Kae-SO\(_3\), Kae-SO\(_3\)-Ga and Vc as standard radical scavenger. At the concentration of 75 \(\mu\text{mol}\cdot\text{L}^{-1}\), Kae showed about 66% radical scavenging activity towards DPPH, while Kae-SO\(_3\)-Ga about 82%, the highest value than other three Kae-based compounds. Thus, the results indicated that Kae-SO\(_3\)-Ga had the best radical scavenging activity, and these modifications in Kae appeared to offer a workable way to improve the free radical scavenging activity.

In order to confirm the radical scavenging activity, the absorption of active ABTS assay at 734 nm in the presence of these Kae-based compounds was observed. Figure 6b showed the ABTS** radical scavenging abilities of Kae-based compounds at different concentrations, respectively. Compared with the DPPH radical scavenging ability, Kae-based compounds showed a higher ABTS** radical quenching capacity, which might be due to the different reaction mechanisms of the used methods [45]. Among the Kae-based compounds, Kae-SO\(_3\)-Ga was the most effective compound for scavenging...
ABTS$^{•+}$ radical, which was almost closed to Vc, the standard material. For instance, at a concentration of 50 µmol·L$^{-1}$, Kae scavenged free radical about 58%, while Kae-SO$_3$-Ga about 84%, the highest value than other three Kae-based compounds (Table 6b).

| Concentration (×10$^{-6}$ mol·L$^{-1}$) | Kae (%) | Kae-Ga (%) | Kae-SO$_3$ (%) | Kae-SO$_3$-Ga (%) | Vc (%) |
|----------------------------------------|--------|------------|----------------|-------------------|-------|
| 5                                     | 5.07   | 6.24       | 8.60           | 11.53             | 15.29 |
| 15                                    | 12.85  | 16.26      | 17.51          | 20.85             | 36.79 |
| 30                                    | 27.34  | 30.63      | 33.89          | 36.75             | 45.44 |
| 45                                    | 41.84  | 43.90      | 48.48          | 53.92             | 63.93 |
| 60                                    | 54.13  | 56.36      | 63.90          | 65.68             | 78.65 |
| 75                                    | 65.75  | 72.77      | 80.85          | 82.47             | 88.37 |

The distinct antioxidant activity of Kae-SO$_3$-Ga could be due to its obviously improving water-solubility and enhancing electron-donating capacity through a chelating Ga(III) ion with ring C. Moreover, the active hydroxyl groups of Kae were retained in Kae-SO$_3$ and Kae-SO$_3$-Ga after sulfonation. Thus, Kae-SO$_3$-Ga obtained an enhancing ability to stabilize unpaired electrons and then scavenge free radicals. It was thus suggested that Kae-SO$_3$-Ga was more suitable as an antioxidant than the parent Kae due to its potential to scavenge free radicals more effectively.

It is well known that flavonoids have many beneficial pharmacological and biological functions, such as antioxidant, anticoagulant, immunomodulatory, anti-inflammatory, anti-microbial, etc. [41,46]. However, some types of flavonoids showed cytotoxicity against normal healthy cells. Wangchuk et al. [46] reported that luteolin exhibited considerable cytotoxicity on a human normal cholangiocyte cell line. Kae, quercetin and isoquercitrin showed hemolytic potential on human erythrocytes [47]. In our preliminary study about cytotoxicity, interestingly, Kae and its complexes showed low cytotoxicity against normal epithelial cells, but were reasonably toxic on breast cancer cells. Further studies would be carried out on their potential biological activities.

3. Conclusions

Based on flavonoid Kae, three compounds including sulfonated Kae, the Ga(III) complexes with Kae and sulfonated Kae were successfully synthesized and characterized by means of UV-Vis and FT-IR spectroscopies, $^1$H- and $^{13}$C-NMR spectrometry, mass spectrometry and thermogravimetric analysis. Sulfonated Kae and sulfonated Kae-Ga(III) complex showed greater water-solubility than Kae, revealing that the poor water-solubility of flavonoids was overcome by sulfonation, which was related to the increasing molecular polarity by negatively-charged sulfonic group and the complex inner shell. The DPPH and ABTS radical scavenging activity by using UV–Vis spectroscopy under imitated physiological conditions revealed that Kae-SO$_3$-Ga exhibited the best antioxidant activity than Kae, Kae-Ga and Kae-SO$_3$. Both flavonoid sulfonation and metal ion have impacts on the antioxidative ability of free flavonoid, revealing that the sulfonated flavonoid-metal complex obtains
optimal antioxidant activity through enhanced water-solubility and electron-donating capacity. It
could be inferred that sulfonated flavonoid-metal complexes might be potential antioxidants for
therapeutic studies. However, further studies are required to understand their fate in the biological
system as well as their mechanisms of action in health and disease.

4. Materials and Methods

4.1. Materials and Instrument

Kae (98%) was purchased from Meilun Biotechnology Co. Ltd. (Dalian, China). Gallium nitrate
(Ga(NO)₃), potassium persulfate (K₂S₂O₈), DPPH and ABTS assays were delivered by Aladdin
Biochemical Co. Ltd. (Shanghai, China). Ethanol was obtained from Tianjin Damao Chemical
Reagent Factory (Tianjin, China). The stock solutions of Kae and Ga salt were prepared in ethanol.

The UV-Vis spectra were recorded by a Cary 5000 UV-Vis-NIR spectrometer (Varian, Palo Alto,
CA, USA) with a wavelength range of 200–800 nm. The FT-IR spectra were recorded in the range of
400–4000 cm⁻¹ using a Nicolet 6700 art visible spectrometer (Thermo Scientific, Madison, WI, USA)
and samples were prepared in the solid state (KBr pellets). TG/DSC analysis of the complex was
carried out by a thermal analyzer of TGA/DSC 3+ (Mettler Toledo, Schwerzenbach, Switzerland).

4.2. Synthesis of Kae Derivatives

Kae-Ga: Kae-Ga was synthesized according to a previous study with appropriate
modifications [48]. Kae and Ga(NO)₃ were dissolved in ethanol. The two solutions were mixed
and refluxed. The solvent was then removed under reduced pressure. The dark brown residues
were washed three times with ethanol and dried in air, and its yield was 81.29%.

Kae-SO₃⁻: Kae was added to concentrated sulfuric acid. The mixture was stirred at 60 °C, then
dispersed into saturated sodium chloride solution and filtered to remove the solvent under reduced
pressure. The filtered solids were dissolved in distilled water and appropriate sodium chloride
was added to it. Floculent deposits were found after a period of 12 h of static culture. Lastly, the
precipitate was filtered and dried in air, presenting as light yellow snowflake microcrystalline and its
yield was 73.12%.

Kae-SO₃⁻-Ga: Kae-SO₃⁻ was dissolved in 20 mL distilled water, followed by introducing 10 mL
aqueous solution of Ga(NO)₃. The mixture was refluxed at 70 °C. The solid brownish precipitate
was rinsed carefully by distilled water, then evaporated rotationally and dried in air, and its yield
was 65.92%.

4.3. UV-Vis Spectra, FT-IR Spectra and TG/DSC Measurements

UV-Vis spectra of Kae and its Ga(III) complexes were obtained by a double beam
spectrophotometer. The dynamics of the development of the complexes and the interactions between
Kae and Ga(III) ion were detected by monitoring the changes in UV-Vis spectrum. Infrared spectra
were recorded in KBr pellets with a spectrometer in the range of 400–4000 cm⁻¹. The TG/DSC analysis
of the complex was carried out by a thermal analyzer of TGA/DSC 3+ at a heating rate of 20 °C min⁻¹
between 25–1000 °C and under N₂ atmosphere (flow rate: 10 mL min⁻¹).

4.4. NMR Spectrometry

¹H NMR and ¹³C NMR experiments were carried out by a Bruker Avance 300 MHz NMR
spectrometer in DMSO-d₆ using tetramethyl silane (TMS) as internal standard, and the chemical shift
values of hydrogen atoms and carbon atoms were reported in parts per million (ppm), respectively.

4.5. HRMS

Positive ion mode of detection was employed in HRMS by using the X500R QTOF system
(AB SCIEX, Framingham, MA, USA). Sample solutions were continually infused via a syringe pump
at a flow rate of 0.3 µL·min\(^{-1}\). The results of mass spectrometry confirmed that the Kae metal complexes were successfully synthesized due to peaks at \(m/z = 639.0034\) and 798.9646 for Kae-Ga and Kae-SO\(_3\)-Ga, respectively.

4.6. Measurements of Antioxidant Activity by DPPH and ABTS Radicals

The DPPH radical scavenging assay was used according to the previous method with some modifications [49]. Briefly, Kae and Kae-Ga were pre-dissolved in ethanol, while Kae-SO\(_3\) and Kae-SO\(_3\)-Ga in PBS solution. Then different concentrations of Kae-based compounds (5, 15, 30, 45, 60, 75 × 10\(^{-6}\) mol·L\(^{-1}\)) was added to DPPH solution (5 × 10\(^{-5}\) mol·L\(^{-1}\)), respectively, at room temperature. The reaction was kept in the dark for 40 min. Then the absorbance of samples (A) at 517 nm was recorded and compared with that of control (A\(_0\)), which was prepared in an analogous method with no addition of the Kae-based compounds. Vc was used as a standard radical scavenger for comparison. The suppression ratio for DPPH was calculated from the following expression: Scavenging ratio (%) = \([(A_0 - A)/A_0]\) × 100%.

The ABTS\(^{+}\) radical scavenging activity of Kae-based compounds was measured using the previous method with some modifications [50]. The radical cation was prepared by mixing a 7.4 mmol·L\(^{-1}\) ABTS\(^{+}\) stock solution with 2.6 mmol·L\(^{-1}\) K\(_2\)S\(_2\)O\(_8\) (1:1, v/v), and the mixture was incubated for 14 h in darkness at 4 °C. The ABTS\(^{+}\) solution was diluted to the absorbance approximately 0.7 at 734 nm to gain an ABTS\(^{+}\) working solution. Sample solutions (5, 10, 20, 30, 40, 50 × 10\(^{-6}\) mol·L\(^{-1}\)) and ABTS\(^{+}\) working solution were mixed together, respectively, and placed at room temperature for 5 min before detection. A solution without sample was set as a control. The absorbance of samples (A) and that of control (A\(_0\)) was recorded at 734 nm. ABTS\(^{+}\) scavenging activity was calculated by the following formula: Radical scavenging ratio (%) = \((A_0 - A/A_0)\) × 100%.

4.7. Statistical Analysis

All experiments were repeated at least three times. Data was analyzed using one-way analysis of variance (ANOVA) and Tukey’s test by SPSS software. \(p < 0.05\) was considered significant.

Author Contributions: Conceptualization, S.-P.D.; Data curation, S.-P.D. and Y.-L.Y.; Formal analysis, S.-P.D., Y.-L.Y., X.-X.C., W.-R.L. and J.-Y.C.; Funding acquisition, S.-P.D. and J.-Y.C.; Investigation, Y.-L.Y.; Supervision, S.-P.D.; Validation, S.-P.D.; Visualization, S.-P.D.; Writing—original draft, S.-P.D. and Y.-L.Y.; Writing—review & editing, S.-P.D.

Acknowledgments: This work was supported by Guangdong Natural Science Foundation, China (No. 2016A030313078), the Fundamental Research Funds for the Central Universities, China (No. 21617430), Foundation for Distinguished Young Talents in Higher Education of Guangdong, China (No. LYM11020), and Macau Science and Technology Development Fund (No. 028/2014/A1).

Conflicts of Interest: The authors declare no conflicts of interest.

References

1. Quideau, S.; Deffieux, D.; Douat-Casassus, C.; Pouységuy, L. Plant polyphenols: Chemical properties, biological activities, and synthesis. Angew. Chem. Int. Ed. 2011, 50, 586–621. [CrossRef] [PubMed]
2. Martins, N.; Barros, L.; Ferreira, I.C.F.R. In vivo antioxidant activity of phenolic compounds: Facts and gaps. Trends Food Sci. Technol. 2016, 48, 1–2. [CrossRef]
3. Anandam, S.; Selvamuthukumar, S. Fabrication of cyclodextrin nanosponges for quercetin delivery: Physicochemical characterization, photostability, and antioxidant effects. J. Mater. Sci. 2014, 49, 8140–8153. [CrossRef]
4. Balducci, V.; Incerpi, S.; Stano, P.; Tofani, D. Antioxidant activity of hydroxytyrosyl esters studied in liposome models. Biochim. Biophys. Acta 2018, 1860, 600–610. [CrossRef] [PubMed]
5. Guo, R.; Chang, X.; Guo, X.; Brennan, C.S.; Li, T.; Fu, X.; Liu, R.H. Phenolic compounds, antioxidant activity, antiproliferative activity and bioaccessibility of Sea buckthorn (Hippophae rhamnoides L.) berries as affected by in vitro digestion. Food Funct. 2017, 8, 4229–4240. [CrossRef] [PubMed]
6. Grzesik, M.; Naparà, O.K.; Bartosz, G.; Sadowska-Bartosz, I. Antioxidant properties of catechins: Comparison with other antioxidants. *Food Chem.* **2018**, *241*, 480–492. [CrossRef] [PubMed]

7. Syed, D.N.; Adhami, V.M.; Khan, N.; Khan, M.I.; Mukhtar, H. Exploring the molecular targets of dietary flavonoid fisetin in cancer. *Semin. Cancer Biol.* **2016**, *40–41*, 130–140. [CrossRef] [PubMed]

8. Oteiza, P.J.; Fraga, C.G.; Mills, D.A.; Taft, D.H. Flavonoids and the gastrointestinal tract: Local and systemic effects. *Mol. Asp. Med.* **2018**, *61*, 41–49. [CrossRef] [PubMed]

9. Souza, R.F.V.D.; Giovani, W.F.D. Synthesis, spectral and electrochemical properties of Al(III) and Zn(II) complexes with flavonoids. *Spectrochim. Acta Part A Mol. Biomol. Spectrosc.* **2005**, *61*, 1985–1990. [CrossRef] [PubMed]

10. Cao, H.; Chen, X.; Jassbi, A.R.; Xiao, J. Microbial biotransformation of bioactive flavonoids. *Biotechnol. Adv.* **2015**, *33*, 214–223. [CrossRef] [PubMed]

11. Cömert, E.D.; Gökmen, V. Evolution of food antioxidants as a core topic of food science for a century. *Food Res. Int.* **2018**, *105*, 76–93. [CrossRef] [PubMed]

12. Chandrasekaran, A.; Idelchik, M.D.; Melendez, J.A. Redox control of senescence and age-related disease. *Redox Biol.* **2017**, *11*, 91–102. [CrossRef] [PubMed]

13. Chanput, W.; Krueyos, N.; Ritthiruangdej, P. Anti-oxidative assays as markers for anti-inflammatory activity of flavonoids. *Int. Immunopharmacol.* **2016**, *40*, 170–175. [CrossRef] [PubMed]

14. Varatharajan, K.; Pushparani, D.S. Screening of antioxidant additives for biodiesel fuels. *Renew. Sustain. Energy Rev.* **2018**, *82*, 2017–2028. [CrossRef]

15. Jabeen, E.; Janjua, N.K.; Ahmed, S.; Murtaza, I.; Ali, T.; Hameed, S. Radical scavenging propensity of Cu2+, Fe3+ complexes of flavonoids and in-vivo radical scavenging by Fe3+-primuletin. *Spectrochim. Acta Part A Mol. Biomol. Spectrosc.* **2017**, *171*, 432–438. [CrossRef] [PubMed]

16. Hofer, T.; Jørgensen, T.O.; Olsen, R.L. Comparison of food antioxidants and iron chelators in two cellular free radical assays: Strong protection by luteolin. *J. Agric. Food Chem.* **2014**, *62*, 8402–8410. [CrossRef] [PubMed]

17. Tu, L.Y.; Pi, J.; Jin, H.; Cai, J.Y.; Deng, S.P. Synthesis, characterization and anticancer activity of kaempferol-zinc(II) complex. *Biorg. Med. Chem. Lett.* **2016**, *26*, 2730–2734. [CrossRef] [PubMed]

18. Samsonowicz, M.; Regulska, E. Spectroscopic study of molecular structure, antioxidant activity and biological effects of metal hydroxyflavonol complexes. *Spectrochim. Acta Part A Mol. Biomol. Spectrosc.* **2017**, *173*, 757–771. [CrossRef] [PubMed]

19. Sharma, R.K. Synthesis and characterization of a biologically active lanthanum(III)–catechin complex and DNA binding spectroscopic studies. *Spectrosc. Lett.* **2009**, *42*, 178–185.

20. Ravishankar, D.; Salamah, M.; Attina, A.; Potli, R.; Vallance, T.M.; Javed, M.; Williams, H.F.; Alzahrani, E.M.S.; Kabova, E.; Vaiyapuri, R. Ruthenium-conjugated chrysin analogues modulate platelet activity, thrombus formation and haemostasis with enhanced efficacy. *Sci. Rep.* **2017**, *7*, 5738. [CrossRef] [PubMed]

21. Medina, J.J.M.; Naso, L.G.; Sadowska-Bartosz, I. Antioxidant properties of catechins: Comparison with other antioxidants. *Food Chem.* **2018**, *241*, 480–492. [CrossRef] [PubMed]

22. Uehara, T.; Yokoyama, M.; Suzuki, H.; Hanaoka, H.; Arano, Y. A gallium-67/68–labeled antibody fragment (pyNO2GaCl: A Redox-Active Gallium Complex. *Inorganochem.* **2018**, *6*, 50. [CrossRef]

23. Kirsh, J.; Woodside, A.; Manor, B.; Carroll, P.; Rablen, P.; Graves, C. Synthesis and Characterization of (pyNO–Η2GaCl): A Redox-Active Gallium Complex. *Inorganochem.* **2018**, *6*, 50. [CrossRef]

24. Ūehara, T.; Yokoyama, M.; Suzuki, H.; Hanaoka, H.; Arano, Y. A gallium-67/68–labeled antibody fragment for immuno-SPECT/PET shows low renal radioactivity without loss of tumor uptake. *Clin. Cancer Res.* **2018**, *24*, 3309–3316. [CrossRef] [PubMed]

25. Sasikumar, A.; Joy, A.; Pillai, M.R.; Nanabala, R.; Anees, K.M.; Jayaprakash, P.G.; Madhavan, J.; Nair, S. Diagnostic Value of 68Ga PSMA-11 PET/CT Imaging of Brain Tumors—Preliminary Analysis. *Clin. Nucl. Med.* **2017**, *42*, e41–e48. [CrossRef] [PubMed]

26. Tarkin, J.M.; Joshi, F.R.; Evans, N.R.; Chowdhury, M.M.; Figg, N.L.; Shah, A.V.; Starks, L.T.; Martin-Garrido, A.; Manavaki, R.; Yu, E.; et al. Detection of atherosclerotic Inflammation by 68Ga-DOTATATE PET compared to [18F]FDG PET imaging. *J. Am. Coll. Cardiol.* **2017**, *69*, 1774–1791. [CrossRef] [PubMed]
27. Eiber, M.; Weirich, G.; Holzapfel, K.; Souvatzoglou, M.; Haller, B.; Rauscher, I.; Beer, A.J.; Wester, H.J.; Gschwend, J.; Schwaiger, M.; et al. Simultaneous 68Ga-PSMA HBED-CC PET/MRI improves the localization of primary prostate cancer. *Eur. Urol.* 2016, 70, 829–836. [CrossRef] [PubMed]

28. Ueda, T.; Inden, M.; Shirai, K.; Sekine, S.; Masaki, Y.; Kurita, H.; Ichihara, K.; Inuzuka, T.; Hozumi, I. The effects of Brazilian green propolis that contains flavonols against mutant copper-zinc superoxide dismutase-mediated toxicity. *Sci. Rep.* 2017, 7, 2882. [CrossRef] [PubMed]

29. Kashyap, D.; Sharma, A.; Tuli, H.S.; Sak, K.; Punia, S.; Mukherjee, T.K. Kaempferol—A dietary anticancer molecule with multiple mechanisms of action: Recent trends and advancements. *J. Funct. Foods* 2017, 30, 203–219. [CrossRef]

30. Selvaraj, S.; Krishnaswamy, S.; Devashya, V.; Sethuraman, S.; Krishnan, U.M. Flavonoid-metal ion complexes: A novel class of therapeutic agents. *Med. Res. Rev.* 2014, 34, 677–702. [CrossRef] [PubMed]

31. Tu, L.Y.; Bai, H.H.; Cai, J.Y.; Deng, S.P. The mechanism of kaempferol induced apoptosis and inhibited proliferation in human cervical cancer SiHa cell: From macro to nano. *Scanning* 2016, 38, 644–653. [CrossRef]

32. Ayasa, O.; Shingo, M.; Masamori, I.; Makoto, S.; Jun, I.; Ryuichiro, S. Kaempferol stimulates gene expression of low-density lipoprotein receptor through activation of Sp1 in cultured hepatocytes. *Sci. Rep.* 2016, 6, 24940.

33. Guo, M.; Perez, C.; Wei, Y.; Rapoza, E.; Su, G.; Bouabdallah, F.; Chasteen, N.D. Iron-binding properties of plant phenolics and cranberry’s bio-effects. *Dalton Trans.* 2007, 4951–4961. [CrossRef] [PubMed]

34. Dong, H.; Yang, X.; He, J.; Cai, S.; Xiao, K.; Zhu, L. Enhanced antioxidant activity, antibacterial activity and hypoglycemic effect of luteolin by complexation with manganese(II) and its inhibition kinetics on xanthine oxidase. *RSC Adv.* 2017, 7, 53385–53395. [CrossRef]

35. Gao, L.G.; Wang, H.; Song, X.L.; Cao, W. Research on the chelation between luteolin and Cr(III) ion through infrared spectroscopy, UV-vis spectrum and theoretical calculations. *J. Mol. Struct.* 2013, 1034, 386–391. [CrossRef]

36. Apicella, B.; Bruno, A.; Wang, X.; Spinelli, N. Fast Fourier Transform and autocorrelation function for the analysis of complex mass spectra. *Int. J. Mass Spectrom.* 2013, 338, 30–38. [CrossRef]

37. Roy, S.; Mallick, S.; Chakraborty, T.; Ghosh, N.; Singh, A.K.; Manna, S.; Majumdar, S. Synthesis, characterisation and antioxidant activity of luteolin-vanadium(II) complex. *Food Chem.* 2015, 173, 1172–1178. [CrossRef] [PubMed]

38. Tan, M.; Zhu, J.; Pan, Y.; Chen, Z.; Liang, H.; Liu, H.; Wang, H. Synthesis, cytotoxic activity, and DNA binding properties of copper (II) complexes with hesperetin, naringenin, and apigenin. *Bioinorg. Chem. Appl.* 2009, 2009, 347872. [CrossRef] [PubMed]

39. Kwon, Y.; Kim, H.; Park, S.; Jung, S. Enhancement of solubility and antioxidant activity of some flavonoids based on the inclusion complexation with sulfobutylether β-cyclodextrin. *Bull. Korean Chem. Soc.* 2010, 31, 3035–3037. [CrossRef]

40. Teles, Y.C.F.; Souza, M.S.R.; Souza, M.D.F.V.D. Sulphated flavonoids: biosynthesis, structures, and biological activities. *Molecules* 2018, 23, 480. [CrossRef] [PubMed]

41. Han, T.; Cheng, G.; Liu, Y.; Yang, H.; Hu, Y.T.; Huang, W. In vitro evaluation of tectoridin, tectorigenin and tectorigenin sodium sulfonate on antioxidant properties. *Food Chem. Toxicol.* 2012, 50, 409–414. [CrossRef] [PubMed]

42. Loizzo, M.R.; Tundis, R.; Bonesi, M.; Menichini, F.; Mastellone, V.; Avallone, L.; Menichini, F. Radical scavenging, antioxidant and metal chelating activities of *Annona cherimola* Mill. (cherimoya) peel and pulp in relation to their total phenolic and total flavonoid contents. *J. Food Compos. Anal.* 2012, 25, 179–184. [CrossRef]

43. Pu, W.; Wang, D.; Zhou, D. Structural characterization and evaluation of the antioxidant activity of phenolic compounds from *astragalus taipaishanensis* and their structure-activity relationship. *Sci. Rep.* 2015, 5, 13914. [CrossRef] [PubMed]

44. Dejian, H.; Boxin, O.; Prior, R.L. The chemistry behind antioxidant capacity assays. *J. Agric. Food Chem.* 2005, 53, 1841–1856.
46. Wangchuk, P.; Apte, S.H.; Smout, M.J.; Groves, P.L.; Loukas, A.; Doolan, D.L. Defined small molecules produced by Himalayan medicinal plants display immunomodulatory properties. *Int. J. Mol. Sci.* **2018**, *19*, 3490. [CrossRef] [PubMed]

47. Vellosa, J.C.R.; Regasini, L.O.; Khalil, N.M.; Bolzani, V.D.S.; Khalil, O.A.K.; Manente, F.A.; Netto, H.P.; Oliveira, O.M. Antioxidant and cytotoxic studies for kaempferol, quercetin and isoquercitrin. *Eclética Quím.* **2011**, *36*, 7–20. [CrossRef]

48. Vimalraj, S.; Rajalakshmi, S.; Raj, D.P.; Vinoth, S.K.; Deepak, T.; Gopinath, V.; Murugan, K.; Chatterjee, S. Mixed-ligand copper(II) complex of quercetin regulate osteogenesis and angiogenesis. *Mater. Sci. Eng. C Mater. Biol. Appl.* **2018**, *83*, 187–194. [CrossRef] [PubMed]

49. Bukhari, S.B.; Memon, S.; Mahroof-Tahir, M.; Bhanger, M.I. Synthesis, characterization and antioxidant activity copper-quercetin complex. *Spectrochim. Acta Part A Mol. Biomol. Spectrosc.* **2009**, *71*, 1901–1906. [CrossRef] [PubMed]

50. Tian, X.; Liu, Y.; Feng, X.; Khaskheli, A.A.; Xiang, Y.; Huang, W. The effects of alcohol fermentation on the extraction of antioxidant compounds and flavonoids of pomelo peel. *LWT-Food Sci. Technol.* **2018**, *89*, 763–769. [CrossRef]

© 2019 by the authors. Licensee MDPI, Basel, Switzerland. This article is an open access article distributed under the terms and conditions of the Creative Commons Attribution (CC BY) license (http://creativecommons.org/licenses/by/4.0/).