

# Ethnopharmacological, chemical, and pharmacological aspects of *Halenia elliptica*: A comprehensive review

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Submitted: 22-08-2014

Revised: 12-09-2014

Published: 04-08-2015

## ABSTRACT

Traditional Tibetan medicine (TTM) is an old traditional medical system, which is an effective and natural method of improving physical and mental health, and has been widely spread in the western part of China for centuries. *Halenia elliptica* (*H. elliptica*) D. Don, known as “Jiadiranguo” (Tibetan medicine name) is one of the most important herbal medicine in TTM that is from the genus *Halenia* (family: Gentianaceae). The whole herb can be used as a medicine to treat hepatobiliary diseases and xeransis, and possesses many biological and pharmacological activities including heat clearing, bile benefiting, liver soothing, digestion promoting, blood nursing, detoxification activities, and so on. In modern research, *H. elliptica* can be used to treat acute or chronic hepatitis, especially hepatitis B. In addition, the chemical compounds of the herb have potent antihepatitis B virus (anti-HBV) activity *in vitro*. As an important TTM, further studies on *H. elliptica* can lead to the development of new drugs and therapeutics for various diseases, and more attention should be paid on the aspects of how to utilize it better.

**Key words:** Chemistry, ethnopharmacology, *Halenia elliptica*, pharmacology

## INTRODUCTION

As a well-known traditional medicine, *Halenia elliptica* (*H. elliptica*) D. Don (family: Gentianaceae) is customarily used to treat hepatitis and cholecystitis, which is widely distributed across the Qinghai-Tibetan Plateau and the western part of China. It has been officially recorded in “Tibetan medicine standards” since 1995 and more than 100 prescriptions containing *H. elliptica* have been used to treat all kinds of diseases in Tibet and Qinghai.<sup>[1]</sup> Modern pharmacological studies have shown that *H. elliptica* and its active constituents have a wide range of pharmacological properties including hepatoprotective, anticancer, cardiovascular, and antidiabetic activities, most of which seriously matched with

its traditional uses. Meanwhile, it was also served as tea for the people in the Qinghai-Tibetan Plateau.<sup>[2]</sup>

In this review, the advances in ethnopharmacological, phytochemical, biological, and pharmacological activities; and the toxicology of *H. elliptica* will be revealed, and the increasing data supports further exploration of *H. elliptica* and its active constituents.

## Botany and ethnopharmacology

### Botany

According to the description of the flora of China, *H. elliptica* is about 15-90 cm tall. The stems are erect, subquadrangular, striate, simple, or branched. The petiole of the basal leaves are flattened and 1-3 cm in length. The stem leaves are sessile or have a short petiole; the leaf blades are oblong, elliptic, ovate-lanceolate, or ovate with a dimension of 1.5-7 cm × 0.5-3.5 cm. The number of leaf veins is five. The calyx lobes are elliptic to ovate in shape and the apex is acuminate. The corolla is campanulate, varies from blue to purple in color, 1-2.5 cm in length, basal spurs are 5-14 mm, lobes elliptic to ovate, apex obtuse and apiculate [Figure 1]. According to the size of the basal spurs, two varieties of species were separated from *H. elliptica*, namely, *H. elliptica* var. *elliptica* and *H. elliptica* var. *grandiflora*. However, due to their similar appearance they are called by the same name, “Jiadiranguo” in traditional Tibetan medicine (TTM).

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10.4103/0973-7847.162114



**Figure 1:** The habitat and its flower of *Halenia elliptica* D. Don

### Resource distribution

There are about 100 species of plants of *Halenia* genus around the world, mainly distributed in Northern Hemisphere and South America. In China, there are two species including *H. elliptica* and *H. Corniculata* (L.) Cornaz, found in asymmetric geographical distribution. *H. elliptica* has a huge reserve and is widely distributed over Tibet, Sichuan, Yunnan, Guizhou, Qinghai, Xinjiang, Hunan, and Hubei while *H. Corniculata* (L.) Cornaz is mainly distributed in North China such as Inner Mongolia, Hebei, Liaoning, Jilin, and Heilongjiang.<sup>[3]</sup> Plants of *Halenia* genus grow in regions with altitudes between 800 and 4600 meters above sea level, mostly in the mountains having an altitude of 1200-3500 m and seldom in areas with elevation higher than 3500m. They mainly inhabit in hillside meadows, roadsides, and thickets, and even in waterside or arbor forests.<sup>[4]</sup>

### Ethnopharmacology

*H. elliptica* was first recorded in *Tibetan Medical Thangka of The Four Medical Tantras*, a classic Tibetan medical book.<sup>[5]</sup> It was also documented in the *Jing Zhu Ben Cao* that “Jiadiranguo” grows in waterside marshes with iron chopstick-like stems, glaucous leaves, and slightly curly base. The herbs are bitter in taste and can be used in the treatment of pathopyretic ulcer.<sup>[6]</sup> The *Yue Wang Yao Zhen* recorded that it can clear liver heat and gallbladder heat.<sup>[7]</sup> The *Gan Lu Ben Cao Ming Jing* stated that it tastes bitter and acrid, and is a cool-natured herb with the ability to clear pulse heat and treat blood and gallbladder diseases.<sup>[8]</sup> The *Zang Yao Pei Fang Xin Bian* recorded that it is a bitter, cool-natured herb, and effective in treating gallbladder heat, headache, stomach heat, and liver heat. *H. elliptica* has also been used in Mongolian medicine for a long time, which is called “Huridige” and belongs to *Rosette saxifrage*. *Tu Jian* accounted that “Huridige” grows in the shade of forest, possesses round and thick leaves, short stems clustered as trotters, and light yellow flowers in closed form. It is a bitter, cool-natured herb, neutral in nature, and effective in clearing heat.<sup>[9]</sup>

## PHYTOCHEMISTRY RESEARCH

More than 30 compounds have been isolated and identified from the whole plant of *H. elliptica*, which abounds with xanthenes and

xanthone glycosides, chromones as well as flavonoids, secoiridoid glycosides, and triterpenoid alkaloids.<sup>[10]</sup> Some of them display many bioactivities *in vivo* or *in vitro*; and the different chemical compositions of *H. elliptica* serve its different pharmacological activities.

### Xanthenes and xanthone glycosides

The main chemical components of *H. elliptica* are xanthenes and xanthone glycosides. Sun isolated five kinds of free liposoluble xanthenes of *H. elliptica* D. Don plants such as 1,7-dihydroxy-2,3,4,5-tetramethoxyxanthone (I), 1,5-dihydroxy-2,3,7-trimethoxyxanthone (II), 1,2-dihydroxy-3,4,5-trimethoxyxanthone (III), 1,5-dihydroxy-2,3-dimethoxyxanthone (IV), 1,7-dihydroxy-2,3-dimethoxyxanthone (V) were obtained.<sup>[11]</sup> Three new xanthone dual glucosides, 1-O- $[\beta$ -D-xylopyranose-(1-6)- $\beta$ -D-glucopyranose]-2,3,5,7-tetramethoxyxanthone (VI), 1-O- $[\beta$ -D-xylopyranose-(1-6)- $\beta$ -D-glucopyranose]-2,3,5-trimethoxyxanthone (VII), and 1-O- $[\beta$ -D-xylopyranose-(1-6)- $\beta$ -D-glucopyranose]-2,3,4,5-tetramethoxyxanthone (VIII) were obtained from a water extract of *H. elliptica* D. Don herbs.<sup>[4]</sup> VI and VII, named haleniaside and demethoxyhaleniaside, respectively, are the two main active components of antihepatitis. Xanthone-O-glucosides isolated from *H. elliptica* D. Don plants by Dhasmana were identified as 2,3,7-trimethoxyxanthone-1-O-glucoside (IX) and 2,3,5-trimethoxyxanthone-1-O-glucoside (X).<sup>[12,13]</sup> Shi got access to two xanthenes, 1-hydroxy-2,3,7-trimethoxyxanthone and 1-hydroxy-2,3,4,7-tetramethoxyxanthone.<sup>[14]</sup> For the first time, Zhang isolated 1-hydroxy-3,7,8-trimethoxyxanthone (XIII) and 1,7-dihydroxy-3,8-dimethoxyxanthone (XIV) from *H. elliptica* D. Don plants.<sup>[15]</sup> Gao got 1,7-dihydroxy-2,3,5-trimethoxyxanthone (XV) from the ethanol extract of *H. elliptica* D. Don plants;<sup>[16]</sup> and another new xanthone, 1,5-dihydroxy-2,3,4-trimethoxyxanthone, was isolated in 2011 (XVI).<sup>[17-21]</sup> The structures of xanthenes and xanthone glycosides listed above are shown in Figure 2.

### Chromones

Nine water-soluble compounds were isolated from the water soluble parts of *H. elliptica*, six of which were elucidated structurally as chromone derivatives (1-6, halenic acid A-C, halenichromone A-C) and three were known as 2-methylchromones (7-9). Compounds 7-9 were isolated from natural sources for the first time.<sup>[22]</sup> The structures of compounds 7-9 are shown in Figure 3.

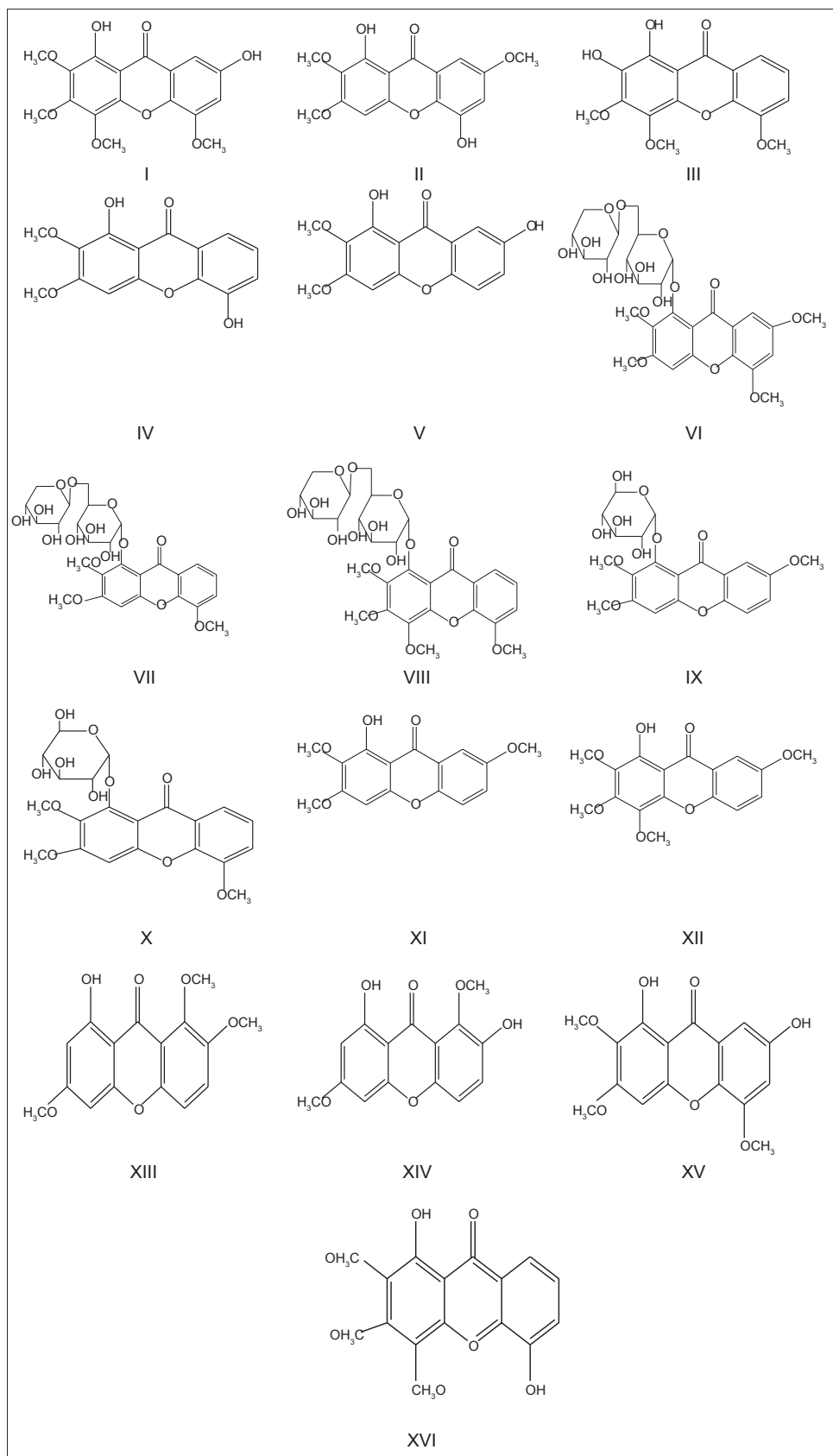
### Other ingredients

Some flavonoids have been obtained from the whole plant of *H. elliptica* including luteolin-O- $\beta$ -D-glucopyranoside, 7-O-primeverosylluteolin, 7-O-glucosylluteolin, luteolin, and apigenin.<sup>[19]</sup> Dhasmana isolated fat-soluble ingredients with antihepatitis activity from the plants such as pentacyclic triterpenoid oleanolic acid and sitosterol glucoside.<sup>[20]</sup> Gao obtained secoiridoid glycoside, swertiamarin and sweroside.<sup>[23]</sup>

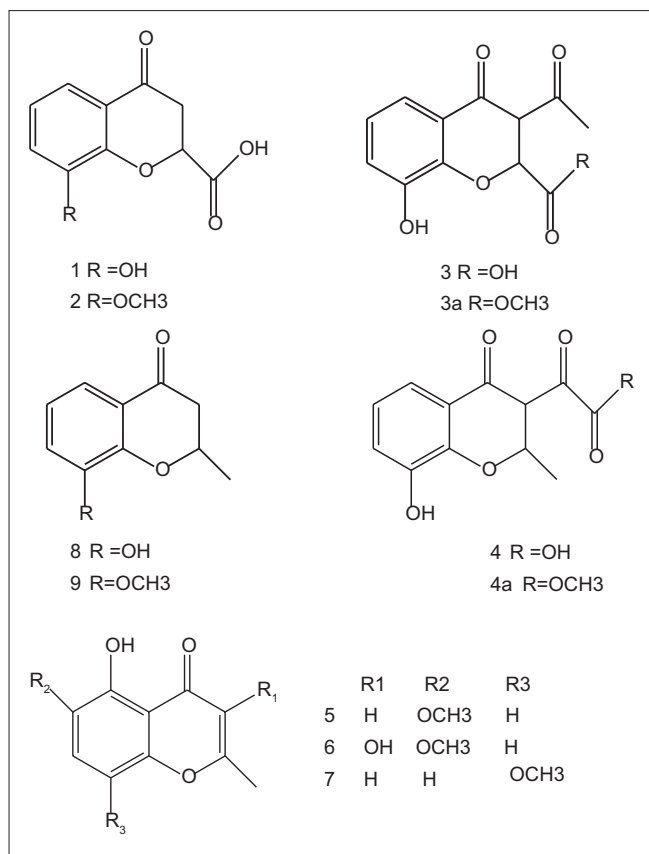
### Pharmacological effects

#### Protective effects against liver injury

Hepatoprotective activity, as another important property of *H. elliptica*, has been comprehensively studied. The hepatoprotective



**Figure 2:** Xanthenes and xanthone glycosides structures from *H. elliptica*



**Figure 3:** Chromones structures from *H. elliptica*

effects against carbon tetrachloride ( $\text{CCl}_4$ )-induced liver toxicity in rats was observed by Zhang.<sup>[24]</sup> The rats were randomly divided into four groups, three experimental groups and a normal control group. The experimental groups were intraperitoneally injected *Halenia* decoction (0.5 mg/g amount to crude drug 0.5 g), total xanthone glycosides (5 mg/100 g), and haleniaside (3 mg/100 g), respectively, and the normal control group was intraperitoneally given water for 11 days. All the groups except the normal control group were injected with  $\text{CCl}_4$  to induce acute hepatic injury. Before injecting the drugs, the outline of liver lobules was vague, hepatic cells around the central vein were obviously necrotic, and the cytoplasm of liver lobule reduced extremely or even disappeared in the test groups. After injecting the drugs, most of the liver lobule outline was distinct, the necrotic area around the central vein decreased, and integral karyon in the central area increased. From the liver histochemistry of the treated groups it was found that the RNA content increased, especially in the haleniaside group. It suggested that haleniaside may be the main active ingredient to increase the RNA content.

Gao compared the inhibition action of six xanthones in  $\text{Fe}^{+2}$ -cystine-induced rat liver microsomal lipid peroxidation at concentrations of 1  $\mu\text{g}/\text{mL}$ , 10  $\mu\text{g}/\text{mL}$ , and 100  $\mu\text{g}/\text{mL}$ , and found that  $\text{C}_1$ -hydroxyl-substituted xanthones have no antioxidant activity that may be caused by the stable hydrogen bonds between the  $\text{C}_1$ -hydroxyl group and the carbonyl group in the structure, which prevents the dissociation of the hydrogen

ion. A stronger antioxidation of  $\text{C}_7$ -hydroxyl-substituted compound demonstrated that  $\text{C}_7$  hydroxyl group is an important part of the activity, so the antioxidant role of 1,7-dihydroxy-2,3,4,5-tetramethoxyxanthone is especially powerful; it is stronger than the classic antioxidant vitamin E. The presence of active hydrogen in the structure of xanthones is an important factor of the hepatoprotective action of *H. elliptica*.<sup>[16]</sup>

### Nonspecific immunomodulatory effects

Zhang has researched the immunopharmacology of *H. elliptica* dry extract to clarify their effect on phagocytic function of the mononuclear macrophage on the basis of their influence on charcoal clearance rate in mice.<sup>[25]</sup> The results showed that the dry extract could enhance the mononuclear macrophage phagocytic function. But the dry extract had no effect on the hemolytic activity of serum hemolysin and splenocyte in normal mice. In the kidney-yang deficiency model, the dry extract can improve the humoral immunity of the hydrocortisone-induced mice.

### Antioxidant properties<sup>[26,27]</sup>

The antioxidant properties of different extracts of *H. elliptica* were investigated by several established *in vitro* and *in vivo* models. The methanol extract of *H. elliptica* revealed strong antioxidant activity *in vitro*. The  $\text{CCl}_4$ -induced liver toxicity experiment showed that rats treated with the methanol extract of *H. elliptica* (100 mg/kg and 200 mg/kg) and silymarin (50 mg/kg) as the standard treatment, had lower levels of alanine transaminase (ALT), aspartate aminotransferase (AST), alkaline phosphatase (ALP), and total bilirubin than those of the  $\text{CCl}_4$  group ( $P < 0.01$ ). The results of the methanol extract at 100 mg/kg were comparable to those of silymarin at 50 mg/kg ( $P > 0.05$ ). The methanol extract did not show any mortality at doses up to 2000 g/kg body weight. These results seem to support the traditional use of *H. elliptica* in pathologies involving hepatotoxicity, and the possible mechanism of this activity may be due to the strong free radical-scavenging and the antioxidant activities of the methanol extract.

### Vasodilation effect<sup>[28,29]</sup>

Xanthones from *H. elliptica* are considered to be vasoactive substances, which exhibit either endothelium-dependent or endothelium-independent mechanisms in rat coronary artery. Six xanthones including 1-Hydroxy-2,3,5-trimethoxyxanthone (1), 1-hydroxy-2,3,4,7-tetramethoxyxanthone (2), 1-hydroxy-2,3,4,5-tetramethoxyxanthone (3), 1,7-dihydroxy-2,3,4,5-tetramethoxyxanthone (4), 1,5-dihydroxy-2,3-dimethoxyxanthone (5), and 1,7-dihydroxy-2,3-dimethoxyxanthone (6) caused vasodilation in the coronary artery precontracted with 1  $\mu\text{M}$  5-hydroxytryptamine. Removal of endothelium of the coronary artery led to decreases in the vasorelaxant effects of 1 and 6 but not of 2, 3, 4, and 5. The mechanism of the vasorelaxant effects of these xanthones may be relevant to the structure-activity differences in the level and the position of the substituent groups with the primary xanthone structure.

### Antihepatitis B virus (anti-HBV) activity<sup>[22]</sup>

Chromone derivatives isolated from *H. elliptica* have been tested for their potencies toward inhibiting the secretion of HBV antigens



in the human HBV-transfected liver cell line HepG2. Lamivudine was used as a positive control, which can suppress HbsAg secretion by 20.1% and HBeAg secretion by 19.7%, at 100 µg/mL (436 µM). The results demonstrated that 2-methylchromones compounds exhibited strong anti-HBV activities, inhibiting HBsAg secretion by 36.8% at a noncytotoxic concentration of 50 µg/mL (284 µM) for 8-hydroxy-2-methyl-4H-1-benzopyran-4-one and by 70.9% at a noncytotoxic concentration of 100 µg/mL (526 µM) for 8-methoxy-2-methyl-4H-1-benzopyran-4-one. While halenic acid and halenia chromone were slightly active or totally inactive at low concentrations. Further investigations are necessary to explore the values of 2-methylchromones as anti-HBV agents.

### Other effects

*H. elliptica* also showed an inhibition effect on contraction of frog heart *in vivo* and *in vitro*. The chloroform extract of the whole plant has antiameoba effect.<sup>[10]</sup>

### Metabolic pathways of xanthone

Feng has studied the *in vitro* metabolic pathways of 1-hydroxyl-2,3,5-trimethoxyxanthone, the main constituent purified from *H. elliptica*, identified three metabolites (M1-M3), which demonstrated that demethylation and hydroxylation were the major phase I metabolic reactions for 1-hydroxyl-2,3,5-trimethoxyxanthone in human liver microsomes; and illustrated that CYP3A4 and CYP2C8 were the primary CYP450 isoforms responsible for its metabolism, and CYP1A2, CYP2A6, CYP2B6, CYP2C9, and CYP2C19 were also involved, especially in the formation of M3.<sup>[30]</sup> 1-hydroxyl-2,3,5-trimethoxyxanthone revealed moderate inhibitory effects on CYP1A2 (IC<sub>50</sub> = 1.06 µM) and CYP2C9 (IC<sub>50</sub> = 3.89 µM), minimal inhibition on CYP3A4 (IC<sub>50</sub> = 11.94 µM), and no inhibition on CYP2D6 (dextromethorphan) and probe substrates CYP2E1 (chlorzoxazone).<sup>[31]</sup> *In vitro* metabolic transformations of five xanthenes of *H. elliptica* have been evaluated by metabolic transformation in rat liver microsomes *in vitro*. The results showed that the metabolic transformation occurred mainly at 2-, 4-, 5-, and 7-carbonic positions on their structures. The metabolites could be considered as the new vasoactive substances.<sup>[32]</sup>

### Toxicology

The results of acute toxicity test show that LD<sub>50</sub> of *H. elliptica* decoction intraperitoneally injected in mice is 27.4 g/kg, and no death is observed in mice after intragastric administration at dose of 100 g/kg. The subacute oral toxicity of *H. elliptica* was investigated in rats. Comparison between the normal control group and the treated group two months after oral administration revealed that the latter were more active with increased food consumption and had more lustrous fur. There were no significant differences in body weight, hemogram, hepatic function, activity of glutamic pyruvic transaminase (GPT), and histological observations of the heart, the liver, and the kidney between the two groups. The results indicated that the medicine was safe in the given dose range.<sup>[25]</sup>

### Future perspectives

*H. elliptica* is one of the most representative Tibetan medicine, which have been recorded in many traditional ancient books. In recent years, phytochemical and pharmacological studies of *H. elliptica* have attracted considerable interest. Large amount of extracts and active constituents have been isolated and proved to have hepatoprotective, antiviral, and antioxidant properties, and enhance the immune response effects, etc. However, many challenges such as poor quality control and failed development of *H. elliptica* still exist. In the future, it is necessary to carry out further study on the structure-activity relationships and action mechanisms of major xanthenes. Research should pay attention to the anti-HBV and other antihepacivirus activities of *H. elliptica* *in vitro/in vivo*. Further studies on *H. elliptica* can lead to the development of new drugs.

### ACKNOWLEDGMENTS

The research was supported by the funds of the Basic Research for Application of Sichuan province of China (2014JY0233), Macao Science and Technology Development Fund (052/2013/A2), Macau University of Science and Technology Fund (0321), and Youth Foundation of Chengdu University (2014XQCKS10).

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**How to cite this Article:** Zhang Z, Bian Q, Luo P, Sun W. Ethnopharmacological, chemical, and pharmacological aspects of *Halenia elliptica*: A comprehensive review. *Phcog Rev* 2015;9:114-9.

**Source of Support:** Nil, **Conflict of Interest:** None declared