

REVIEW

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Hosta plantaginea (Lam.) Aschers (Yuzan): an overview on its botany, traditional use, phytochemistry, quality control and pharmacology

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Hosta plantaginea (Lam.) Aschers, as a traditional folk medicine, has been widely used both as a single herb and in prescriptions in Asia mainly due to its anti-inflammatory and analgesic effects. A total of 101 compounds including steroids, flavonoids, alkaloids and others have been isolated from *H. plantaginea*. Modern pharmacology has revealed that *H. plantaginea* possesses various therapeutic effects such as anti-inflammatory, analgesic and antibacterial effects both *in vitro* and *in vivo*. Although a number of reports on the chemical constituents and pharmacological activities of this plant are available, there is limited research on the bioactive constituents and the mechanism of the biological activities of *H. plantaginea*. Thus, it is essential to strengthen the research on bioactive constituents and their mechanisms as well as their structure–function relationships in *H. plantaginea*. Up to now, only three compounds have been established for the quality control of *H. plantaginea*. However, a comprehensive review on the botany, traditional use, phytochemistry, quality control and pharmacology information about this plant has not been reported so far; thus, a systematic and comprehensive review is very necessary. Therefore, this paper provided a comprehensive overview on the botany, traditional use, phytochemistry, quality control and pharmacology of *H. plantaginea* and also provided evidence for its further research and clinical applications.

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1. Introduction

Hosta Tratt. is a genus in the family Liliaceae, which comprises about 43 species mainly distributed in the temperate and sub-tropical zones of Asia, particularly in Japan and China.^{1–6} *Hosta* plants are commonly used ornationally or for medicine and are thus widely cultivated in parks and botanical gardens throughout the region.^{2,7} Only four native species have been found in China, namely, *H. plantaginea*, *H. ventricosa* (Salisb.) Stearn, *H. ensata* F. Maekawa and *H. albofarinosa* D. Q. Wang, which are widely cultivated in parks and/or commonly used as folk medicine in China.^{2,8,9} Among them, *H. plantaginea* is a perennial herb widely cultivated in China; it is used as a traditional Chinese medicine (TCM) and is known as Yu zan (Chinese name 玉簪). The dried whole plants, leaves, roots and flowers of this plant have been used as local and traditional medicine in China, Japan and South Korea.^{10–12} Chinese people call the flowers of *H. plantaginea* yu-zan-hua, and they are also named *bai-e-hua*, *bai-he-hua* and *nei-xiao-hua*; they have long been commonly used in traditional Mongolian medicine for the

treatment of inflammatory and painful diseases, such as sore throat, muteness, lung heat and toxic heat.^{10,13–16} Moreover, *yuzan-hua* was the main herb in the prescriptions of the *Yuzan qingyan shiwei* pill, *Yuzan qingyan shiwei* powder and *Qinyan liuwei* powder (Fig. 1).

H. plantaginea is enriched in multiple structurally diverse and biologically important steroids and flavonoids. Modern pharmacology has revealed that *H. plantaginea* has anti-inflammatory, analgesic, antibacterial, antifungal, anti-cancer, antioxidant and other biological activities.^{17–19} Up to now, a large number of studies focused on the phytochemistry and pharmacology of *H. plantaginea* have been published. However, a comprehensive review on the botany, traditional use,



Fig. 1 (a) Whole plants; (b) fresh flowers; (c) dried flowers; (d–f) three prescriptions containing the flowers of *H. plantaginea*.

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phytochemistry, quality control and pharmacology information about this plant has not been reported; thus, a systematic and comprehensive review is very necessary. In this review, we systematically summarized the progress in the botany, traditional use, phytochemistry and quality control studies of *H. plantaginea* over the past decades, with all the elucidated compounds listed. The biological characterization of the extracts and constituents isolated from this plant were also discussed. It is hoped that the information presented in this paper will be useful for the full utilization of *H. plantaginea* for new drug development and pharmaceutical applications.

2. Botany

2.1 Morphology

As a perennial herbaceous plant, *H. plantaginea* grows to 40–80 centimeter. The calyx has several to a dozen flowers: the flowers are sessile, ovate or lanceolate, 2.5–7 cm long, and 1–1.5 cm wide; the inner sepals are very small. Also, the flowers are solitary or sometimes in clusters of 2 or 3 and fragrant; the pedicel is about 1 cm. The perianth is white, funnel form, and 10–13 cm in size. The stamens are slightly shorter than or subequaling perianth; the filaments are adnate to the perianth tube near base. The capsule is cylindrical, about 6 × 1 cm, and 3-angled. The flowering stages range from July to August. The rhizome is 1.5–2 cm thick and stout. The petiole grows from 20 to 40 cm in length; the leaf blade is ovate-cordate, -orbicular, or ovate, 14–25 × 8–16 cm, glabrous, veins in 6–10 pairs, base cordate, margin slightly undulate, and apex abruptly acute. Scape 40–80 cm. Raceme: several to more than 10-flowered, bracts 2 subtending each flower, outer one ovate or lanceolate, 2.5–7 × 1–1.5 cm; the inner one is very small.^{2,9}

2.2 Geographical distribution of *H. plantaginea*

H. plantaginea is chiefly distributed in underwood, grassy slope, or rocky regions at lower altitudes below 2000 meters from Sichuan, Hubei, Hunan, Jiangsu, Anhui, Zhejiang and other provinces of China. The requirements for growing the plants are not stringent, and they need little sunshine and appropriate temperatures. Moreover, this plant is mostly cultivated in parks as an ornamental plant.^{2,9}

3. Traditional use

3.1 Medicinal use

H. plantaginea was listed for medicinal use first in “Ben Cao Pin Hui Jing Yao” during the Ming Dynasty more than five hundred years ago. Because of their versatile biological and pharmacological activities, the *H. plantaginea* plants have been traditionally used in China, Japan and South Korea. In traditional Chinese medicine, the flowers were used as an oral medicine for sore throat, muteness, lung heat, and toxic heat. The whole plants or leaves were applied to inflammatory mass, hemorrhoids and snake bite, and the roots were taken orally for inflammatory mass, vomiting blood and osteophytes.^{10,13,14} In the TCM culture, *H. plantaginea* is described as acrid in taste,

a little cold in nature and attributable to the stomach, lung and liver meridians. According to Bencao Gangmu and Zhong's previous reports,^{20,21} the roots of *H. plantaginea* were used as *Sinopodophyllum emodi* (Wall) Ying to treat cancer in Japan because of their very similar morphology and efficacy. Meanwhile, the roots were applied to dermatitis in South Korea.²²

The prescriptions related to the flowers of *H. plantaginea* recorded in TCM were the *Yuzan qingyan shiwei* pill, *Yuzan qingyan shiwei* powder and *Qinyan liuwei* powder, which are used for the treatment of sore throat, muteness, lung heat and toxic heat. Studies on the side effects and safety evaluations of this plant are very limited although it is widely used in TCM.

3.2 Non-medicinal use

In addition to its use in medicine, the flowers, young leaves and buds of *H. plantaginea* are used as a daily food material to eat or drink in some cities of China.^{23,24} The flowers of *H. plantaginea* are very beautiful with concentrated fragrance; thus, the plant is often placed at home or office.^{25,26} Moreover, *H. plantaginea* is commonly used ornamentally and thus widely cultivated in the parks and botanical gardens in China.^{3,4,6,27,28} Additionally, *H. plantaginea* is an ombrophyte and has the advantages of rapid reproduction, easy growth and strong resistance; thus, it is commonly used for vegetation protection.^{27,28}

4. Phytochemistry

Up to now, 101 chemical constituents have been isolated and identified from *H. plantaginea*, including steroids, flavonoids, alkaloids, phenylethanols, acetophenones, and others (Table 1). Among them, steroids and flavonoids are considered to be the primary bioactive constituents of this herbal medicine.

4.1 Steroids

Steroids were regarded as the major bioactive principle of *H. plantaginea*, and phytochemical researchers were focused on these species since 1997.¹¹ Until now, 30 steroid compounds have been isolated and identified from *H. plantaginea* (Fig. 2). The sugar moieties in the carbohydrate part of steroidal saponins are β -D-glucose, β -D-galactose, α -L-rhamnose and β -D-xylose; in contrast, in terms of aglycones as the core of these compounds, there are five different types, namely, spirostanes (**1–19**), stigmastanes (**20–23**), furostanes (**24** and **25**), C22 steroids (**26**), and C21 steriods (**27–30**), as shown in Fig. 2, where the sugar groups are usually attached to the C-3 position of aglycones. By comparison, it can be clearly seen that spirostanes are mainly found as glycosides in this plant, including four different subtypes: gitogenin (**1–11**), tigogenin (**12**), neo-gitogenin (**13**), manogenin (**14–19**).

4.2 Flavonoids

Flavonoids consist of a large group of polyphenolic compounds with benzo- γ -pyrone structures, which are ubiquitously present in plants; there is no exception for the *H. plantaginea* plants. Flavonoids are another major bioactive constituents in *H. plantaginea* and they are divided into three categories, namely,



Table 1 Secondary metabolites isolated from *H. plantaginea*

No. Secondary metabolites	Part	Ref.
Steroids		
1 Gitogenin	Flowers	16, 29 and 30
2 (25 <i>R</i>)-2 α ,3 β ,17 β ,24 β -Tetrahydroxy-5 α -spirostanane	Flowers	31
3 Gitogenin-3- β -D-galactopyranoside	Flowers	30
4 Gitogenin-3- β -D-glucopyranosyl-(1 \rightarrow 4)- β -D-galactopyranoside	Flowers, leaves	24, 30 and 32
5 Gitogenin-3- α -L-rhamnopyranosyl(1 \rightarrow 2)- β -D-galactopyranoside	Flowers, leaves	24 and 30
6 12-Hydroxy-gitogenin-3- α -L-rhamnopyranosyl(1 \rightarrow 2)- β -D-galactopyranoside	Leaves	24 and 32
7 Gitogenin-3- β -D-glucopyranosyl-(1 \rightarrow 4)-O-[α -L-rhamnopyranosyl(1 \rightarrow 2)]- β -D-galactopyranoside	Flowers	30
8 Gitogenin-3- β -D-glucopyranosyl-(1 \rightarrow 2)-O- β -D-glucopyranosyl-(1 \rightarrow 4)- β -D-galactopyranoside	Flowers, underground	11, 16, 29, 30 and 33
9 Gitogenin-3- β -D-glucopyranosyl-(1 \rightarrow 2)-O-[β -D-xylopyranosyl(1 \rightarrow 3)]-O- β -D-glucopyranosyl-(1 \rightarrow 4)- β -D-galactopyranoside	Flowers, underground	11, 16, 29, 30 and 33
10 Gitogenin-3- β -D-glucopyranosyl-(1 \rightarrow 2)-O-[O- α -L-rhamnopyranosyl(1 \rightarrow 4)- β -D-xylopyranosyl(1 \rightarrow 3)]-O- β -D-glucopyranosyl(1 \rightarrow 4)- β -D-galactopyranoside	Flowers, underground	11, 16, 29, 30 and 33
11 Gitogenin-3-O- β -D-xylopyranosyl-(1 \rightarrow 4)- β -D-glucopyranosyl-(1 \rightarrow 2)-[β -D-xylopyranosyl(1 \rightarrow 3)]-O- β -D-glucopyranosyl(1 \rightarrow 4)- β -D-galactopyranoside	Flowers	30
12 Tigogenin-3- β -D-glucopyranosyl(1 \rightarrow 4)-O-[α -L-rhamnopyranosyl(1 \rightarrow 2)]- β -D-galactopyranoside	Flowers, leaves	24 and 30
13 Hostaside IV	Leaves	24
14 Manogenin	Flowers	16
15 9-Dehydromanogenin	Flowers, leaves	31 and 32
16 (25 <i>R</i>)-2 α ,3 β ,24 β -Trihydroxy-5 α -spirost-9(11)-en-12-one	Flowers	31
17 Hostasaponin A	Rhizomes	34
18 Hostasaponin B	Rhizomes	34
19 9,11-Dehydromanogenin-3- β -D-glucopyranosyl-(1 \rightarrow 2)-O-[β -D-xylopyranosyl(1 \rightarrow 3)]-O- β -D-glucopyranosyl-(1 \rightarrow 4)- β -D-galactopyranoside	Leaves	24 and 32
20 β -Sitosterol	Flowers, leaves	32 and 33
21 β -Stigmasterol	Leaves	32
22 β -Sitosterol-3- β -D-glucopyranoside	Flowers, leaves	16, 29, 32, 33 and 35
23 Stigmasterol-3- β -D-glucoside	Flowers	35
24 (2 α ,3 β ,5 α ,25 <i>R</i>)-2,3-Dihydroxy-22-methoxyfurostan-26-yl- β -D-glucopyranoside	Underground	11
25 16,22-Oxido-26-hydroxycholest-4-en-3-one	Flowers	31
26 (2 α ,3 β ,5 α ,16 β)-Pregn-20-ene-20-carboxylic acid-2,16-dihydroxy- γ -lactone-3- β -D-glucopyranosyl-(1 \rightarrow 2)-O- β -D-glucopyranosyl(1 \rightarrow 4)- β -D-galactopyranosyl	Underground	11
27 2 α ,3 β -Dihydroxy-2 α -pregn-16-en-20-one	Rhizomes	34
28 Hostaside III	Leaves	24
29 Hostaside I	Leaves	24
30 Hostaside II	Leaves	24
Flavonoids		
31 Kaempferol	Flowers	29, 36 and 37
32 Astragalin	Flowers	37
33 Kaempferol-7- β -D-glucoside	Flowers	29 and 35-37
34 Kaempferol-3,7-di- β -D-glucoside	Flowers	37
35 Kaempferol-3- β -D-sophoroside	Flowers	37
36 Plantanone A	Flowers	37
37 Kaempferol-3-O-rutinoside	Flowers	29, 36 and 37
38 Kaempferol-3- β -D-glucopyranosyl-(1 \rightarrow 2)- β -D-glucopyranoside-7- β -D-glucopyranoside	Flowers	39
39 Hostaflavone A	Flowers	39
40 Kaempferol-3-O-rutinoside-7-O-glucoside	Flowers, leaves	32 and 37
41 Kaempferol-3-O-(2 β -D-glucopyranosyl)- β -D-rutinoside	Leaves	32
42 Kaempferol-3- β -D-glucopyranosyl-(1 \rightarrow 2)-[α -L-rhamnopyranosyl(1 \rightarrow 6)]- β -D-glucopyranoside	Flowers	37
43 Plantanone B	Flowers	37
44 Kaempferol-3- α -L-rhamnopyranosyl-(1 \rightarrow 6)- β -D-glucopyranosyl-(1 \rightarrow 2)- β -D-glucopyranoside	Flowers	37
45 Kaempferol-3-O-[β -D-glucopyranosyl(1 \rightarrow 2)-[α -L-rhamnopyranosyl(1 \rightarrow 6)]- β -D-glucopyranosyl]-7- β -D-glucopyranoside	Flowers	37
46 Plantanone C	Flowers	41
47 Quercetin	Flowers	36
48 Hostaflavanone A	Flowers	38
49 5,7-Dimethoxy-4'-hydroxyflavan	Flowers	40
50 5,7-Dimethoxy-8-methyl-4'-hydroxyflavan	Flowers	40
51 Epicatechin	Flowers	40
52 Catechin	Flowers	40



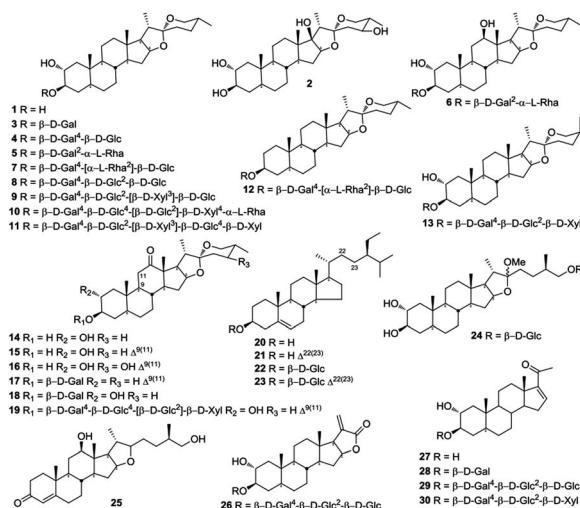
Table 1 (Contd.)

No.	Secondary metabolites	Part	Ref.
53	Epigallocatechin	Flowers	40
54	Gallocatechin	Flowers	40
Alkaloids			
55	Hostasinine A	Whole plants	42
56	Hostasine	Whole plants	43
57	8-Demethoxyhostasine	Whole plants	43
58	8-Demethoxy-10-O-methylhostasine	Whole plants	43
59	10-O-Methylhostasine	Whole plants	43
60	(+)-Haemanthamine	Whole plants	43
61	O-Demethylhaemanthamine	Whole plants	43
62	Haemanthidine	Whole plants	43
63	Yemenine C	Whole plants	43
64	Lycorine	Whole plants	43
65	Pseudolycorine	Whole plants	43
66	O-Methylcorenine	Whole plants	43
67	9-O-Demethyl-7-O-methyllycorenine	Whole plants	43
68	Albomaculine	Whole plants	43
69	7-Deoxy- <i>trans</i> -dihydronarciclasine	Whole plants	43
70	8-O-Demethylmaritidine	Whole plants	43
71	Ungermine	Whole plants	43
72	Norsanguinine	Whole plants	43
73	(1 <i>S</i> ,3 <i>S</i>)-1-Methyl-1,2,3,4-tetrahydro- β -carboline-3-carboxylic acid	Flowers	38
74	(1 <i>R</i> ,3 <i>S</i>)-1-Methyl-1,2,3,4-tetrahydro- β -carboline-3-carboxylic acid	Flowers	38
Phenylethanols and acetophenones			
75	Phenethyl- β -D-glucopyranoside	Flowers	38 and 40
76	Phenethanol- β -D-gentiobioside	Flowers	34
77	Phenethyl-O-rutinoside	Flowers	34
78	4-Hydroxylacetophenone	Flowers	40
79	Acetophenone-4-O- β -D-glucoside	Flowers	40
80	2-Hydroxyl-6-methoxyacetophenone-4-O- β -D-glucoside	Flowers	40
Phenylpropanoids			
81	<i>anti</i> -1-Phenylpropane-1,2-diol	Flowers	38
82	<i>anti</i> -1-Phenylpropane-1,2-diol-2-O- β -D-glucopyranoside	Flowers	38
83	Coumaric acid	Flowers	40
84	3-(4-Hydroxy-3-methoxyphenyl) acrylic acid methyl ester	Flowers	40
85	3,4-Dihydroxycinnamyl alcohol-3-O-glucoside	Flowers	40
Terpenoids			
86	Hoplanoside A	Flowers	44
87	Lomacarinoside A	Flowers	44
88	(4 <i>S</i>)-4-Hydroxy-3,5,5-trimethyl-2-cyclohexen-1-one,4-[3-(β -D-glucopyranosyloxy)butyl]	Flowers	45
89	Roseoside	Flowers	45
90	Plantaginoside	Flowers	45
Aliphatics			
91	Docosanol	Leaves	32
92	Arachic acid	Flowers	29 and 36
93	Hexadecanoic acid	Flowers	29 and 36
94	Hostacerebroside A	Flowers	46
91	Docosanol	Leaves	32
92	Arachic acid	Flowers	29 and 36
93	Hexadecanoic acid	Flowers	29 and 36
94	Hostacerebroside A	Flowers	46
95	1-O- β -D-Glucopyranosyl-(2 <i>S</i> ,3 <i>R</i> ,4 <i>E</i> ,8 <i>Z</i>)-2-[(2-hydroxytetradecanoyl)amido]-4,8-octadecadienyl-1,3-diol	Flowers	31
96	(2 <i>S</i>)-1-O-Linenoyl-3-O- β -D-galactopyranosylglycerol	Flowers	31
97	(2 <i>S</i>)-1-O-(10,13)-Octadecenoyl-3-O- β -D-galactopyranosyl glycerol	Flowers	31
Others			
98	4-Hydroxybenzaldehyde	Flowers	40
99	4-Hydroxy-3-methoxybenzene	Flowers	31



Table 1 (Contd.)

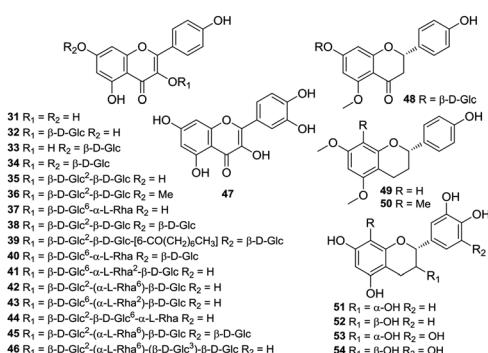
No.	Secondary metabolites	Part	Ref.
100	(2-Methyl) heptyl phthalate	Flowers	31
101	(2-Methylphenyl) (4-hydroxy-3-methoxyphenyl)-1,4-diene-3-pentanone	Flowers	31

Fig. 2 Chemical structures of steroids from *H. plantaginea*.

17 flavonols (31–47), one flavanone (48), and six flavans (49–54) (Fig. 3). Among them, kaempferol (31) and its derivatives (32–46) as well as quercetin (47) belong to flavonols, which are the main active ingredients of *H. plantaginea* and most of them are flavonoid glycosides with 3- and/or 7-linked glycans. In fact, 22 flavonoids were isolated and identified from the flowers of *H. plantaginea* by us and in Yu's previous work in 2017–2019.^{37–41} This is the first report on the isolation of flavonones and flavans from the Liliaceae family.³⁸

4.3 Alkaloids

The *H. plantaginea* plants are also rich in alkaloids, and some of them show notable inhibitory activities against acetylcholinesterase (AChE) and tobacco mosaic virus (TMV).⁴³ To date, 20

Fig. 3 Chemical structures of flavonoids from *H. plantaginea*.

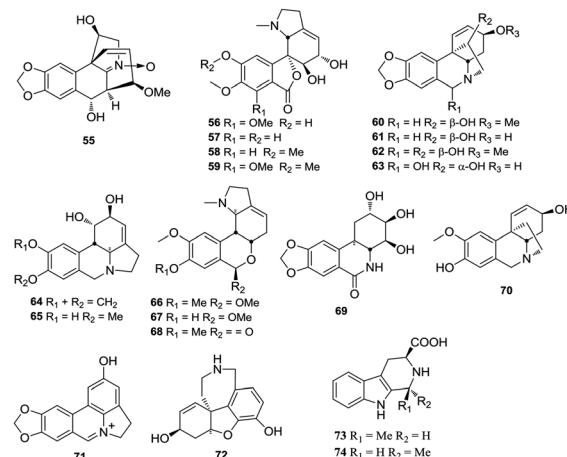
alkaloids have been isolated and identified from *H. plantaginea*. These alkaloids include 18 benzylphenethylamine alkaloids (55–72) and two beta-carboline alkaloids (73 and 74) (Fig. 4). Hostasinine A (55) is a benzylphenethylamine alkaloid with a novel skeleton featuring a C-4–C-6 linkage and a nitrone moiety. Moreover, other 17 benzylphenethylamine alkaloids (56–72) represent five skeletal types of alkaloids, namely, lycoreniene-type (56–59 and 66–68), isocarbstyryl-type (69), crinine-type (60–63 and 70), lycorine-type (64, 65 and 71), and galanthamine-type (72).

4.4 Phenylethanols and acetophenones

Phenylethanols and acetophenones represent a relatively small class of compounds in *H. plantaginea*. To date, only three phenylethanols, namely, phenethyl-O-beta-D-glucopyranoside (75), phenethanol-beta-D-gentiobioside (76) and phenethyl-O-rutinoside (77) have been obtained from the ethanolic extract of *H. plantaginea*.^{38,40} Moreover, three acetophenones, namely, 4-hydroxyacetophenone (78), acetophenone-4-O-beta-D-glucoside (79) and 2-hydroxyl-6-methoxyacetophenone-4-O-beta-D-glucoside (80) have also been isolated from *H. plantaginea* (Fig. 5).⁴⁰

4.5 Phenylpropanoids

Phenylpropanoids also represent a relatively small class of compounds in *H. plantaginea*. To date, only two phenylpropanols (81 and 82) and three phenylpropionic acids (83–85) have been obtained from the ethanolic extract of *H. plantaginea* (Fig. 6).^{38,40}

Fig. 4 Chemical structures of alkaloids from *H. plantaginea*.

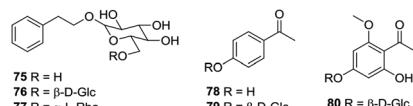


Fig. 5 Chemical structures of phenylethanols and acetophenones from *H. plantaginea*.

4.6 Terpenoids

To date, only two monoterpenes (86 and 87) and three megastigmanes (88–90) have been obtained from the ethanolic extract of *H. plantaginea* (Fig. 7).^{44,45}

4.7 Aliphatics

Some studies have been carried out to investigate the aliphatic compounds in *H. plantaginea*. Seven aliphatics (91–97) have been isolated from the flowers and leaves of *H. plantaginea* (Fig. 8).

4.8 Others

In addition to the above-mentioned main components, other components have also been found in the flowers of *H. plantaginea*, such as 4-hydroxybenzaldehyde (98),⁴⁰ 4-hydroxy-3-methoxybenzene (99),³¹ (2-methyl)heptyl phthalate (100),³¹ and (2-methylphenyl) (4-hydroxy-3-methoxyphenyl)-1,4-diene-3-pentanone (101)³¹ (Fig. 9).

5. Quality Control

Quality control is very important for the use of TCMs. Many rapid, sensitive and stable technologies such as UPLC-MS, HPLC and UV have been applied for the qualitative and quantitative analyses of *H. plantaginea*.^{47–52} To date, only three compounds, namely, kaempferol (31), kaempferol-7-O- β -D-glucoside (33) and kaempferol-3-O-rutinoside (37) have been used as quantitative markers by HPLC or UPLC-MS. Interestingly, the contents of compound 31 were found to be 0.0025%, 0.33–0.56%, 0.29–0.44% and 0.020% in the flowers of *H. plantaginea* by different research groups.^{47,49–51} Moreover, the contents of compounds 33 and 37 were found to be 0.0058% and 0.090% by UPLC-MS, respectively.⁴⁶ In addition, the content

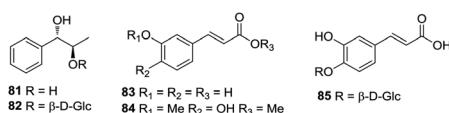


Fig. 6 Chemical structures of phenylpropanoids from *H. plantaginea*.

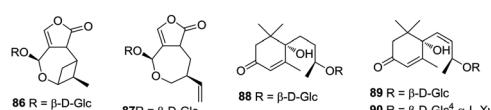


Fig. 7 Chemical structures of terpenoids from *H. plantaginea*.

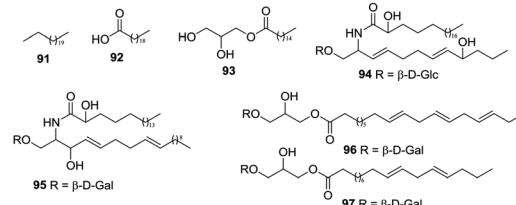


Fig. 8 Chemical structures of aliphatics from *H. plantaginea*.

of total saponins was found to be 1.61–4.70% in the flowers of *H. plantaginea* by ultraviolet-visible (UV) spectrophotometry.⁵² Furthermore, Li and co-authors established the fingerprint of the flowers of *H. plantaginea* by HPLC.⁴⁸ In fact, the harvest times, geographical locations, and other factors can affect the contents of the active compounds in the flowers of *H. plantaginea*, which should be considered when assessing their clinical efficacies. Therefore, there is an urgent need to determine other bioactive markers using various chromatographic and spectroscopic means to establish the comprehensive quality standards for this medicinal plant.

6. Pharmacology

H. plantaginea has long been used in China, Japan, Korea and other countries because of its various pharmacological effects. In recent years, research reports on the chemical constituents and pharmacological activities of *H. plantaginea* have shown an increasing trend. Modern pharmacology has revealed that *H. plantaginea* has anti-inflammatory, analgesic, antibacterial, antifungal, anti-cancer, antioxidant and other biological activities.

6.1 Anti-inflammatory and analgesic activities

In TCM, the whole plants, leaves, roots and flowers of *H. plantaginea* have been used as local and traditional ethnic medicine for the treatment of sore throat, inflammatory mass, dysuria, lung heat, snake bite and others. In agreement with the traditional usage of *H. plantaginea*, several studies have illustrated that this plant possesses anti-inflammatory and analgesic effects both *in vitro* and *in vivo*. The anti-inflammatory abilities against cyclooxygenase (COX)-1 and -2 enzymes were evaluated. Flavonoid compounds 31–40, 42–45 and 48 showed significant inhibitory activities against COX-1 and COX-2 at a concentration of 50 mM, with the inhibition ratios ranging from 53.00% to 80.55% for COX-1 and from 52.19% to 66.29% for COX-2. Further detailed testing showed that these compounds inhibited the COX-1 and COX-2 enzymes with the IC₅₀ values of 12.90–46.16 μ M in comparison with the positive control celecoxib with

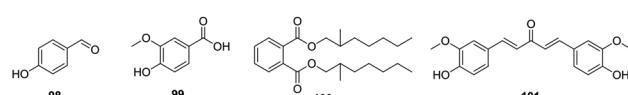


Fig. 9 Chemical structures of others from *H. plantaginea*.



the IC_{50} values of 9.00 μM for COX-1 and 1.04 μM for COX-2.³⁷⁻³⁹ Thus, flavonoids are supposed to contribute to the anti-inflammatory effect of *H. plantaginea*.

In vivo, at the doses of 1.0, 2.0 and 4.0 g kg^{-1} (raw herb), different concentrations of ethanol (50%, 65%, 80% and 95%) crude extracts obtained from the flowers of *H. plantaginea* showed moderate anti-inflammatory effects in a xylene-induced ear oedema mouse model.⁵³ Moreover, the crude extract, ethyl acetate fraction and *n*-BuOH fraction showed a significant anti-inflammatory effect in xylene-induced mouse ear oedema, acetic acid-induced writhing and carrageenan-induced oedema from different research groups.^{32,54,55} In addition, the leaves of *H. plantaginea*, Yu-Zan-Qing-Yan-Shi-Wu-Wei powder and Yu-Zan-Qing-Yan-Shi-Wu-Wei pill have good therapeutic effects on patients with acute and chronic pharyngitis.⁵⁶⁻⁵⁸

In 2010, the analgesic activity of *H. plantaginea* was studied *in vivo*.⁵⁹ The oral administration of a 50% ethanol extract of the *H. plantaginea* flowers (0.5 and 2.0 g kg^{-1} , raw herb) caused a significant analgesic effect in an acetic acid-induced writhing mouse model with the inhibition of 39.28% and 53.41%, respectively; the positive control drug xiao-yan-tong pill displayed inhibition of 67.07% at a concentration of 0.1 g kg^{-1} . Moreover, 50% ethanol extract of the flowers of *H. plantaginea* also had a significant antinociceptive effect in the hot plate mouse model at a dose of 2.0 g kg^{-1} compared with the control group.

6.2 Anti-cancer activity

The crude extracts and steroids from *H. plantaginea* have significant activities against tumor cells such as the L615, MDA-MB-231, MCF-7, SMMC-7721, HL-60, Jurkat, K562, HepG2, MDCK, and YAC-A cell lines *in vitro*. However, no study has been conducted on the anti-cancer activity of *H. plantaginea* *in vivo*.

The ethanol extract from the flowers of *H. plantaginea* exhibited significant anti-cancer activity against the L615 tumor cells.¹⁴ The steroid compounds **2**, **15**, and **25** exhibited significant anti-cancer activities against MDA-MB-231, MCF-7 and SMMC-7721 tumor cells.³¹ Moreover, compounds **8-10** exhibited significant anti-cancer activities against the leukaemia HL-

60 cells in a dose-dependent manner with the IC_{50} values of 2.9, 1.0 and 1.7 $\mu\text{g mL}^{-1}$, respectively.¹¹ In addition, compounds **1**, **4**, **5**, and **7-11** exhibited significant or moderate anti-cancer activities against leukaemia HL-60, Jurkat, and K562 cells and others (Table 2).^{30,60} Thus, steroids are supposed to contribute to the anti-cancer effect of *H. plantaginea*.

6.3 Antioxidant activity

Recently, some studies have revealed the anti-oxidative effect of the compounds in the 2,2-diphenyl-1-picrylhydrazyl (DPPH) assay *in vitro*.^{37-39,45} The flavonoid compounds **31**, **33** and **48** exhibited strong antioxidant effects with the IC_{50} values of 36.3, 77.6 and 83.2 μM , respectively, compared to the positive control L-ascorbic acid (V_c) (33.9 μM). Moreover, three megastigmane glycosides **88-90** showed very strong antioxidant effects with the IC_{50} values of 1.34, 1.66 and 1.28 μM , respectively, compared to the positive control V_c (1.01 μM).

6.4 Anti-acetylcholinesterase and anti-viral activities

Three benzylphenethylamine alkaloids from the whole plants of *H. plantaginea*, namely, 8-demethoxy-10-O-methylhostasine (**58**), ungermine (**71**) and norsanguinine (**72**) demonstrated significant anti-acetylcholinesterase (AChE) activity, with the IC_{50} values of 2.32, 3.85 and 1.43 μM , respectively, compared to the positive control tacrine (0.20 μM).^{42,43} Moreover, 7-deoxy-*trans*-dihydronarciclasine (**69**) showed strong inhibitory activity against tobacco mosaic virus (TMV) than the positive control ribavirin, with the IC_{50} values of 1.80 and 2989.60 μM , respectively.⁴³

6.5 Other biological activities

In modern research, the *H. plantaginea* plant has been reported to have a variety of activities besides the above-mentioned fields. The roots of *H. plantaginea* are considered to be effective on recalcitrant contact dermatitis in patients.²² Additionally, the whole plants of *H. plantaginea* can improve bone hyperplasia in patients.⁶¹

Table 2 Anti-cancer activity of the isolated compounds

Compounds	HL-60 (μM)	Jurkat (μM)	K562 (μM)	HepG2 (μM)	MCF-7 (μM)	SGC-7901 (μM)	MDCK ($\mu\text{g mL}^{-1}$)	YAC-1 ($\mu\text{g mL}^{-1}$)	SMMC-7721 ($\mu\text{g mL}^{-1}$)
1	>40	>40	>40	9.95	>40	39.5	287.38	5.38	2.84
4	20.58	16.96	17.65	8.17	>40	9.92	— ^a	— ^a	— ^a
5	18.28	13.46	15.25	7.14	>40	7.32	— ^a	— ^a	— ^a
7	3.11	4.32	5.25	1.14	1.51	1.72	5.12	— ^a	— ^a
8	3.50	3.95	4.35	1.13	1.23	1.85	— ^a	— ^a	— ^a
9	2.45	2.84	2.76	0.16	0.56	0.44	51.16	24.23	— ^a
10	2.92	4.21	3.14	1.13	0.89	0.34	22.57	12.83	16.17
11	2.41	4.54	3.24	1.16	1.02	0.59	— ^a	— ^a	— ^a
10-Hydroxycamptothecin	0.04	0.03	1.60	11.90	16.81	19.80	— ^a	— ^a	— ^a
Cisplatin	1.92	3.66	17.63	4.58	57.25	5.79	— ^a	— ^a	— ^a

^a Not determined.



7. Conclusion and future perspectives

This review summarized information on the botany, traditional use, phytochemistry, quality control and pharmacology of *H. plantaginea*. The amount of data gathered from different studies revealed that this medicinal plant is rich in many secondary metabolites and vast biological active constituents. However, the bioactive constituent action mechanisms and their structure–function relationships need to be investigated for further development into therapeutics.

Systematic efficacy studies are necessary to examine the standardized extracts of *H. plantaginea* and to identify the bioactive constituents responsible for the pharmacological effects. Furthermore, the anti-inflammatory activity and its mechanisms are limited, which consequently limit the use of this medicinal plant to treat inflammation-related diseases. The secondary metabolites of steroids and flavonoids from *H. plantaginea* may be drug candidates for treating inflammation-related diseases because of their potent anti-inflammatory activities. To this end, no study has been performed to evaluate the toxic effects of *H. plantaginea*. However, further clinical evaluation must be performed to perceive the detailed effect of this plant on humans.

We believe that this review can be of particular value by providing fundamental insights into the medicinal value of this plant. Moreover, this review can provide a reference for clinical medication, sustainable development and utilization of this plant.

Conflicts of interest

The authors declare no conflict of interest.

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Notes and references

- 1 L. G. Fu and T. Q. Chen, *Higher plants of China*, 2002, vol. 13, pp. 91–92.
- 2 J. Tang and F. Z. Wang, *Flora of China*, 1980, vol. 14, pp. 49–52.
- 3 Q. J. Li, Y. F. He, C. Wang, Y. R. Chang, L. P. Wang, Q. K. Wang and Y. Q. Liu, *J. Anhui Agric. Sci.*, 2010, **38**, 11826–11829.
- 4 J. Z. Zhang, A. P. Si, G. F. Sun, L. Shi and Q. X. Zhang, *Acta Hortic. Sin.*, 2004, **31**, 549–554.
- 5 W. G. Sehmid, *The genus Hosta*, Timber Press, London, UK, 1991, pp. 428–428.
- 6 Q. Y. Li and Y. P. Xia, *Zhongguo Yuanlin*, 2004, **20**, 77–79.
- 7 J. X. Liu, C. H. Zhao, X. R. Liu, Y. Z. Xi and Y. L. Zhang, *Plant Syst. Evol.*, 2011, **294**, 99–107.
- 8 D. Q. Wang, *Guishaia*, 1989, **9**, 297–298.
- 9 X. Q. Chen and E. B. David, *Flora of China*, 2000, vol. 24, pp. 204–205.
- 10 State Administration of Traditional Chinese Medicine, *Zhonghua Bencao*, 1999, **8**, 107–109.
- 11 Y. Mimaki, A. Kameyama, M. Kuroda, Y. Sashida, T. Hirano, K. Oka, K. Koike and T. Nikaido, *Phytochemistry*, 1997, **44**, 305–310.
- 12 D. J. Park, H. J. Im, H. G. Kim, W. H. Yang, S. H. Yong and M. S. Choi, *J. Environ. Biol.*, 2018, **39**, 507–516.
- 13 M. R. Jia and Y. Zhang, *Dictionary of Chinese ethnic medicine*, 2016, pp. 468–468.
- 14 State Administration of Traditional Chinese Medicine, *Zhonghua Bencao*, 2004, **32**, 142.
- 15 Y. Xin, *Chin. Tradit. Pat. Med.*, 2015, **37**, 653–656.
- 16 J. H. Zhang, H. X. Xie, P. F. Xue, H. G. Zhang, X. Y. Liu and M. S. Baiji, *Chin. Pharm. J.*, 2010, **45**, 335–337.
- 17 J. W. He, L. Yang and G. Y. Zhong, *Chin. Tradit. Herb. Drugs*, 2016, **47**, 4295–4300.
- 18 L. Yang, Y. Q. Wang, J. W. He, X. M. Wang and J. X. Zhu, *Chin. Med. Mat.*, 2016, **39**, 216–222.
- 19 L. Yang, J. J. Zhao, Y. Q. Fang, J. X. Zhu, X. M. Wang, J. W. He and G. Y. Zhong, *Chin. J. Exp. Tradit. Med. Formulae*, 2016, **22**, 230–234.
- 20 G. Y. Zhong, L. S. Xu, G. J. Xu and T. Namba, *China J. Chin. Mater. Med.*, 2002, **27**, 89–94.
- 21 S. Z. Li, *Compendium of Materia Medica*, 1977, vol. 2, pp. 1208–1208.
- 22 S. J. Yun, J. Y. Lee, G. H. Kim, T. H. Kim, A. Y. Lee, S. H. Lee and J. S. Hong, *J. Eur. Acad. Dermatol. Venereol.*, 2018, **32**, e28–e29.
- 23 Y. Xin and W. L. J. Ao, *Zhong guo yao xue da hui ji di shi jie zhong guo yao shi zhou da hui*, 2010, p. 6.
- 24 M. Y. Wang, Y. Peng, C. S. Peng, J. Q. Qu and X. B. Li, *J. Asian Nat. Prod. Res.*, 2018, **20**, 501–509.
- 25 Q. Liu, G. F. Sun, J. Z. Zhang and X. D. Li, *Sci. Agric. Sin.*, 2015, **48**, 4323–4334.
- 26 M. Yu, Y. Yu, L. P. Xu, H. Z. Liu and S. Y. Liu, *North. Hortic.*, 2015, **27**, 198–201.
- 27 H. Dai, Y. M. Fang, L. B. Huang and M. Zhang, *Journal of Jiangsu Forestry Science & Technology*, 2014, **41**, 37–41.
- 28 M. X. Guan and R. Dong, *North. Hortic.*, 2013, **25**, 182–185.
- 29 J. H. Zhang, *Inner mongolia medical college*, 2009.
- 30 J. Q. Liu, C. F. Wang, M. H. Qiu and W. X. Hu, *Chin. Tradit. Herb. Drugs*, 2010, **41**, 520–526.
- 31 X. Liu, *Studies on the bioactive constituents from Zingiber officinale Rosc and Hosta plantaginea (Lam.) Aschers*, Third Military Medical University, 2017.
- 32 J. Y. Qu, M. Y. Wang, C. M. Wang, G. Y. Zhong and X. B. Li, *Chin. Tradit. Herb. Drugs*, 2011, **42**, 217–221.
- 33 W. Y. Li, *Preliminary study on the chemical compositions and bioactivities of Mongolian medicine Hosta plantaginea (Lam.) Ascherson*, Huazhong University of Science and Technology, 2009.



- 34 M. Y. Wang, Z. H. Xu, Y. Peng, G. Y. Zhong and X. B. Li, *Chem. Nat. Compd.*, 2016, **52**, 1047–1051.
- 35 J. He, *Study on quality standard and chemical constituents of Hosta ventricosa stearn*, Guangzhou University of Chinese Medicine, 2010.
- 36 H. X. Xie, J. H. Zhang, H. G. Zhang and P. F. Xue, *China J. Chin. Mater. Med.*, 2009, **44**, 733–735.
- 37 J. W. He, L. Yang, Z. Q. Mu, Y. Y. Zhu, G. Y. Zhong, Z. Y. Liu, Q. G. Zhou and F. Cheng, *RSC Adv.*, 2018, **8**, 18175–18179.
- 38 L. Yang, S. T. Jiang, Q. G. Zhou, G. Y. Zhong and J. W. He, *Molecules*, 2017, **22**, 1825.
- 39 J. W. He, X. Y. Huang, Y. Q. Wang, J. Liang, R. H. Liu, G. Y. Zhong and L. Yang, *Nat. Prod. Res.*, 2019, **33**, 1599–1604.
- 40 H. Yu, Q. H. Wang, J. J. Han, B. Y. Q. E. Bao and W. L. J. Ao, *Chin. Tradit. Pat. Med.*, 2017, **39**, 107–111.
- 41 Q. G. Zhou, L. Yang, J. W. He and G. Y. Zhong, *China J. Chin. Mater. Med.*, 2019, **44**, 3312–3315.
- 42 Y. H. Wang, S. Gao, F. M. Yang, Q. Y. Sun, J. S. Wang, H. Y. Liu, C. S. Li, Y. T. Di, S. L. Li, H. P. He and X. J. Hao, *Org. Lett.*, 2007, **9**, 5279–5281.
- 43 Y. H. Wang, Z. K. Zhang, F. M. Yang, Q. Y. Sun, H. P. He, Y. T. Di, S. Z. Mu, Y. Lu, Y. Chang, Q. H. Zheng, M. Ding, J. H. Dong and X. J. Hao, *J. Nat. Prod.*, 2007, **70**, 1458–1461.
- 44 Q. H. Wang, J. J. Han and B. Y. Q. E. Bao, *J. Food Biochem.*, 2017, **41**, e12320.
- 45 X. H. Bao, Q. F. Wang, B. Y. Q. E. Bao, J. J. Han and W. L. J. Ao, *Chem. Nat. Compd.*, 2017, **53**, 614–617.
- 46 H. X. Xie and P. F. Xue, *China Pharm.*, 2014, **23**, 12–13.
- 47 J. N. Ma, C. M. Mao, Z. J. Tian and W. D. Liu, *Chin. Tradit. Pat. Med.*, 2017, **39**, 639–641.
- 48 H. F. Li, M. J. Wang and H. N. Zhu, *China Med. Her.*, 2017, **14**, 34–37.
- 49 W. L. J. Ao, M. G. Bai, H. G. J. L. T. Wang and S. Weijimis, *Drug Stand. China*, 2012, **13**, 347–350.
- 50 X. J. Li, P. F. Xue, A. H. Ju, J. P. Gao and Y. L. Ren, *Neimenggu Yixueyuan Xuebao*, 2011, **33**, 307–310.
- 51 J. He, Y. Gao and W. M. Li, *Tradit. Chin. Drug Res. Clin. Pharmacol.*, 2010, **21**, 192–194.
- 52 H. X. Xie, P. F. Xue, H. G. Zhang, L. Q. Fan and H. M. Bian, *J. Beijing Univ. Tradit. Chin. Med.*, 2009, **32**, 624–625.
- 53 J. W. He, L. Yang, J. X. Zhu, X. M. Wang, Z. R. Zou, W. W. He and G. Y. Zhong, *J. Jiangxi Norm. Univ.*, 2016, **40**, 183–185.
- 54 Y. Xin and Dalahu, *Asia-Pacific Traditional Medicine*, 2015, **11**, 5–7.
- 55 C. Y. Li, P. F. Xue, M. N. Liu, P. P. Wang and L. Wang, *Lishizhen Med. Mater. Med. Res.*, 2015, **26**, 1559–1560.
- 56 A. L. Guan and M. D. L. Naren, *J. Med. Pharm. Chin. Minor.*, 2015, **21**, 21–22.
- 57 J. L. Bao, *Shijie Zuixin Yixue Xinxi Wenzhai*, 2016, **16**, 172.
- 58 Z. M. He, *Journal of Sichuan Traditional Chinese Medicine*, 1994, **13**, 53.
- 59 H. X. Xie, P. F. Xue, J. Zhou, L. Fan and F. H. Yuan, *Acta Acad. Med. Nei. Mongol.*, 2010, **32**, 36–38.
- 60 M. M. Wu, X. J. Li, P. F. Xue, S. L. Shi, C. Y. Li, H. X. Xie and J. H. Zhang, *Chin. Med. J. Res. Pract.*, 2018, **32**, 16–18.
- 61 X. Tang, *Zhongguo Xiangcun Yisheng Zazhi*, 1992, **4**, 32.

