

A review of the pharmacological effects of *Arctium Lappa* (burdock)

Yuk-Shing Chan^a, Long-Ni Cheng^a, Jian-Hong Wu^a, Enoch Chan^a, Yiu-Wa Kwan^b, Simon Ming-Yuen

Lee^c, George Pak-Heng Leung^d, Peter Hoi-Fu Yu^a, Shun-Wan Chan^a

^a State Key Laboratory of Chinese Medicine and Molecular Pharmacology, Department of Applied

Biology and Chemical Technology, The Hong Kong Polytechnic University, Hong Kong

^b Institute of Vascular Medicine, School of Biomedical Sciences, Faculty of Medicine, The Chinese

University of Hong Kong, Hong Kong

^c Institute of Chinese Medical Sciences, University of Macau, Av. Padre Tomas Pereira S.J., Taipa,
Macau, PR China

^d Department of Pharmacology and Pharmacy, The University of Hong Kong, Hong Kong

Correspondence

*Dr. George Pak-Heng Leung, Department of Pharmacology and Pharmacy, The University of Hong Kong, Hong Kong SAR, PR of China. E-mail address: gphleung@hkucc.hku.hk; Phone: +852-28192861; Fax: +852-28170859

*Dr. Shun-Wan Chan, State Key Laboratory of Chinese Medicine and Molecular Pharmacology, Department of Applied Biology and Chemical Technology, The Hong Kong Polytechnic University, Hong Kong SAR, PR of China. E-mail address: bcswchan@polyu.edu.hk; Phone: +852-34008718; Fax: +852-23649932

Abstract. *Arctium lappa*, commonly known as burdock, is being promoted/recommended as a healthy and nutritive food in Chinese societies. Burdock has been used therapeutically in Europe, North America and Asia for hundreds of years. The roots, seeds and leaves of burdock have been investigated in view of its popular uses in Traditional Chinese Medicine (TCM). In this review, the reported therapeutic effects of the active compounds present in the different botanical parts of burdock are summarized. In the root, the active ingredients have been found to “detoxify” blood in TCM term and promote blood circulation to the skin surface, improving the skin quality/texture and curing skin diseases like eczema. Antioxidants and anti-diabetic compounds have also been found in the root. In the seeds, some active compounds possess anti-inflammatory effects, and potent inhibitory effects on the growth of tumors such as the pancreatic carcinoma. In the leaf extract, the active compounds isolated can inhibit the growth of micro-organisms in the oral cavity. The medicinal uses of burdock in treating chronic diseases like cancers, diabetes and AIDS have been reported. However, it is also essential to be aware of the side-effects of burdock including contact dermatitis and other allergic/inflammatory responses that might be evoked by burdock.

Key Words: *Arctium lappa* (burdock); Traditional Chinese Medicine; Anti-inflammatory; Pharmacology

Introduction

Starting from the end of twentieth century, the majority of people in developed countries are becoming wealthier and more health conscious. They tend to spend extra money on different functional foods or nutraceuticals in order to pursue healthy aging. Natural products have been used in the treatment of various chronic human pathological conditions because they are rich in antioxidants (Guo *et al.*, 2008). In traditional Chinese medicine (TCM), it is believed that food and medicine stem from the same origin but with different uses and applications (Chan *et al.*, 2010). Therefore, it is common for Chinese people to incorporate different medicinal herbs into their diet to produce various “healthy” food recipes in order to achieve better taste, more attractive appearance and improved texture of the food and most importantly to improve health.

Burdock, a perennial herb in the family of *Compositae*, stores most of its nutrients during the first year. These nutrients are then used for the flower-blooming process afterwards. The plant, which can be found worldwide, has been cultivated as a vegetable for a period of long time in Asia (Morita *et al.*, 1993). Burdock, called “Niubang” in Chinese, has been used in China and some Western countries for over 3000 years and its therapeutic uses were documented in *The Compendium of Materia Medica* (*Bencao gangmu* in Chinese) written by Li Shizhen, the most famous/important figure in the history and development of TCM, during the Ming

dynasty (Yu *et al.*, 2003).

Burdock is traditionally used to treat diseases such as sore throat and infections such as rashes, boils and various skin problems. In TCM understanding, these pathological events are mainly due to the accumulation of toxin in the body. The dried root of one-year old burdock (Figure 1) is the major part used for different therapeutic purposes although burdock leaves and fruit/seeds are also used. It is suggested that the root of this herb is particularly effective and invaluable in eliminating heavy metals from our body. Therefore it appears to have the function of draining toxins in terms of TCM theory (Yu *et al.*, 2003).

In contrast to some famous and expensive medicinal herbs such as *Ganoderma lucidum* (Lingzhi) and *Panax ginseng* (Ginseng) that have been used for a long period of time with their rich and highly acclaimed nutritional values, burdock possesses various therapeutic values but is still sold at a low price. Moreover, it can be easily cultivated. In light of the aforementioned properties of this herb, the aim of this review is to summarize the currently available scientific information on burdock so as to provide a comprehensive overview of this herb.

Active ingredients found in burdock

With the advancement of different state-of-the-art analytical techniques, more active ingredients of burdock have been identified over the last decade (Park *et al.*,

2007). The major active ingredients isolated from this herb are: tannin, arctigenin, arctin, beta-eudesmol, caffeic acid, chlorogenic acid, inulin, trachelogenin 4, sitosterol-beta-D-glucopyranoside, lappaol and diartigenin (Table 1). Apart from these compounds, burdock also contains various common nutrients (Table 2).

Pharmacological effects

The extracts from different parts of burdock have long been considered good for health. They help enhance the body's immune system and improve metabolic functions (Lin *et al.*, 2002). Biological activities and pharmacological functions reported for the *Arctium* species include anti-inflammatory, anti-cancer, anti-diabetic, anti-microbial and antiviral activities.

Anti-inflammatory effects

Inhibition of inducible nitric oxide synthase (iNOS) expression and nitric oxide (NO) production, suppression of pro-inflammatory cytokine expression, inhibition of the nuclear factor-kappa B (NF- κ B) pathway, activation of antioxidant enzymes and scavenging of free radicals are the essential mechanisms of burdock's anti-inflammatory action.

The extract of burdock has been shown to exhibit anti-inflammatory response by inhibiting degranulation and release of cysteinyl leukotrienes (Cys-LTs) by peripheral

blood mononuclear cells (PBMCs). Cys-LTs are synthesized inflammatory mediators as histamine and prostaglandins. The blockade of Cys-LT is regarded as inhibition of inflammatory response. Also, the extract of burdock significantly inhibited acute mouse ear edema due to induced allergic response. Therefore, there has been evidence suggesting that burdock has significant anti-inflammatory effect (Knipping *et al.*, 2008).

Lappol F, diartignin and arctigenin, found in the seeds or leaves of burdock, are lignans that can inhibit NO production. The excessive production of NO by iNOS (EC1.14.13.39) is involved in various inflammatory diseases such as rheumatoid arthritis, autoimmune disease, chronic inflammation and atherosclerosis. Therefore, inhibition of NO production by iNOS in macrophages are potential treatments for certain inflammatory diseases (Wang *et al.*, 2007). Lappol F and diartignin strongly inhibited NO production in lipopolysaccharide (LPS)-stimulated murine macrophage RAW264.7 cells with IC₅₀ values of 9.5 and 9.6 μ M, respectively (Park *et al.*, 2007). Further study elucidated that diartignin could directly target NF- κ B-activating signaling cascade by direct inhibition of the DNA binding ability of NF- κ B and inhibition of NF- κ B-regulated iNOS expression (Kim *et al.*, 2008).

Arctigenin, a phenylpropanoid dibenzylbutyrolactone lignan, potently inhibited iNOS expression and NO production through suppression of NF- κ B activation and inhibition of I- κ B α phosphorylation and p65 nuclear translocation in LPS-activated

macrophages (Cho *et al.*, 2002). In addition, arctigenin strongly inhibited the expression of pro-inflammatory cytokines tumor necrosis factor- α (TNF- α) and IL-6, in LPS-stimulated RAW264.7 cells, THP-1 human monocyte-macrophage and differentiated human macrophage U937 (Cho *et al.*, 2002; Zhao *et al.*, 2009). Further study showed that arctigenin-induced inhibition of TNF- α production might be mediated by arctigenin's potent inactivation of mitogen-activated protein (MAP) kinases including ERK1/2, p38 kinase and JNK through the inhibition of MAP kinase kinase (MKK) activity, leading to inactivation of activator protein-1 (AP-1) (Cho *et al.*, 2004; Zhao *et al.*, 2009).

On the other hand, expression of inflammation-associated cyclooxygenase 2 (COX-2) and formation of prostaglandin E₂ (PGE₂) are the results of increased NO production. Inhibitor of COX-2 causes a potent inflammatory effect, since the prostaglandin family is associated with the onset of inflammation. The methanolic extract of burdock has been proven to be effective in inhibiting the expression level of COX-2 mRNA. Therefore the anti-inflammatory effect of Burdock is attributed to the lowered PGE₂ release (Wang *et al.*, 2007).

In view of the inflammatory processes, inflammation has usually been investigated together with the pathway of free radicals. There have been a lot of studies on the association between free radicals, oxidative stress and inflammation (Weber *et al.*, 2005; Abreu *et al.*, 2006; Pontiki *et al.*, 2006). Instead of just looking at

the action of drugs/herbs on pro-inflammatory cytokines or/and other inflammatory mediators, their free radical scavenging capacities should also be considered. There are increasing studies focusing on both the effects of pro-inflammatory signaling and free-radical scavenging capacity of individual drug/herb, which may contribute to the resultant anti-inflammatory effect of them (Lee *et al.*, 2007). Recent studies have demonstrated that burdock's anti-inflammatory characteristics on carrageenan-induced rat paw edema and carbon tetrachloride (CCl₄)-induced hepatotoxicity. The carrageenan-induced rat paw edema assay is a widely used model for acute inflammatory testing. Burdock has shown to have significant inhibition on the growth of rat paw edema in a dose-related manner, thus suggesting some significant anti-inflammatory activities of burdock (Lin *et al.*, 1996). Lin *et al.* (1996) demonstrated the antioxidant power of burdock extract by detecting the signal intensities of 5,5-dimethyl-1-pyrroline-N-oxide (DMPO)-OOH in relation to superoxide dismutase (SOD) concentration. For hepatoprotective effect, burdock was shown to suppress the CCl₄ or acetaminophen-intoxicated mice as well as the ethanol plus CCl₄-induced rat liver damage. The underlying hepatoprotective ability of burdock could be related to the decrease of oxidative stress on hepatocytes by increasing glutathione (GSH), cytochrome P-450 content and NADPH-cytochrome c reductase activity and by decreasing malondialdehyde (MDA) content, hence alleviating the severity of liver damage based on histopathological observations (Lin

et al., 2000; Lin *et al.*, 2002). In summary, the anti-inflammatory action of burdock is attributed to its high free radical scavenging capacities and antioxidant activity.

Anti-cancer activities

During the development of tumors, very large amounts of nutrients (oxygen and nutrients) are required in order to sustain the rapid proliferation of the tumor cells. However, tumor cells can still survive under extreme conditions like low oxygen and low carbohydrate availability due to their relatively high tolerance to hostile environment. Arctigenin, an active compound found in the seeds of burdock, has the ability to eradicate nutrient-deprived cancer cells (Awale *et al.*, 2006). In addition to its board spectrum of activities on different cancer cell lines, e.g. PANC-1 and AsPC-1, arctigenin seems to exhibit a highly preferential cytotoxicity to cancer cells that are bathed in glucose-deprived conditions (Awale *et al.*, 2006). This is because arctigenin has a potent inhibitory effect on the phosphorylation of Akt (Guo *et al.*, 2008), which is stimulated under glucose-deprived conditions. Hence, the rate of glucose formation in cancer cells is decreased, which in turn leads to cell death due to a lack of nutrients (Awale *et al.*, 2006).

Protection of cells from harmful substances can greatly reduce the chance of tumor formation and thus suppresses cancer cell proliferation. Flavonoid-type anti-oxidants and some other active polyphenol antioxidants found in the root of

burdock may account for the suppressive effects on cancer metastasis (Tamayo *et al.*, 2000). It has been shown that extracts of the root protect cells from toxic substances and lower the mutations of cells (Miyamoto *et al.*, 1993)

Tannin, a phenolic compound, is one of the most common active compounds found in the root of burdock. It induces macrophage responses, inhibits tumor growth, and possesses immuno-modulatory properties (Miyamoto *et al.*, 1993). However, tannin is potentially toxic in nature. It may cause stomach upset and at high concentrations it has some dangerous side effects such as nephrotoxicity and hepatic necrosis (Miyamoto *et al.*, 1993). Therefore, the use of tannin should be carefully monitored.

Anti-diabetic activity

Burdock has been used to treat diabetes by TCM practitioners. Several studies have suggested that the root or/and fruit are possible parts with hypoglycemic effect. Sitosterol-beta-D-glucopyranoside is considered to be the most potent and efficacious substance among the large profile of active compounds found in the root of burdock. It has demonstrated potent inhibitory effects on alpha glucosidase activities. Alpha glucosidases are involved in the processing of glycoprotein and glycogenolysis. Inhibitors of glycosidase are potential therapeutic agents in treating diabetes mellitus and obesity (Mitsuo *et al.*, 2005). In addition, gamma-glucoside-fructose ester, also

known as inulin, can help to regulate blood glucose levels. Inulin, a natural carbohydrate present in the root of burdock, can act on cell surface receptors to keep the blood glucose level constant, therefore improving the tolerance to high glucose level. Also, the production of short chain fatty acids is also increased (Silver and Krantz Jr, 1931). The anti-diabetic activity of total lignan from the fruit of burdock has been studied in a model of alloxan-induced diabetes in mice and rats. It has been proven that total lignan from burdock is a safe anti-diabetic agent and may help prevent diabetic complications (Xu *et al.*, 2008).

Anti-microbial and antiviral activity

It has been reported that the lyophilized extract of the leaves of burdock exhibits anti-microbial activity against oral micro-organisms and is the most effective against bacteria related to endodontic pathogens such as: *Bacillus subtilis*, *Candida albicans*, *Lactobacillus acidophilus* and *Pseudomonas aeruginosa* (Pereira *et al.*, 2005). Chlorogenic acid isolated from the leaves also have shown restraining effects on *Escherichia coli*, *Staphylococcus aureus*, and *Micrococcus luteus* (Lin *et al.*, 2004). Therefore, the leaves of Burdock may be useful in treating tooth/gum diseases that are related to micro-organisms in the oral cavity. It is also a potential topical remedy for skin problems such as eczema, acne, and psoriasis. In addition, the polyacetylene ingredients extracted from the root of burdock also possess potent anti-bacterial and

anti-fungal activities (Takasugi *et al.*, 1987).

Constituents of burdock have also demonstrated antiviral activity. Phenolic constituents like caffeic acid and chlorogenic acid possess strong inhibitory effect on herpesvirus (HSV-1, HSV-2) and adenovirus (ADV-3, ADV-11) (Chiang *et al.*, 2002). Arctigenin, one of the lignanoid ingredients, has demonstrated activities against human immunodeficiency virus type-1 (HIV-1) both *in vivo* and *in vitro* (Schroder *et al.*, 1990; Eich *et al.*, 1996). These suggest potential uses of these promising natural compounds isolated from burdock to treat infection by these viruses, especially HIV.

Other activities

Lignans isolated from burdock have been shown to be potent platelet-activating factor (PAF) receptor antagonists, calcium antagonists and hypotensive agent (Ichikawa *et al.*, 1986; Iwakami *et al.*, 1992). Arctiin, a lignin isolated from burdock seeds, has protective effect against 2-amino-1-methyl-6-phenylimidazo [4,5-b] pyridine (PhIP)-induced carcinogenesis (Hirose *et al.*, 2000). Besides arctiin, polyphenolics in burdock, especially caffeic acid and chlorogenic acid, also have significant anti-mutagenic activity, and the anti-mutagenic capacity of the extract of burdock has a positive correlation with polyphenolic content (Liu and Tang, 1997). The anti-decrepitude effect of burdock has also been noted. Li *et al.* (2004) have elucidated that the main mechanism of burdock's anti-decrepitude effect involves

improvement of SOD activity and reduction of MDA and lipofuscin content. Furthermore, burdock has been used as an adjunctive therapy or alternative medicine for the treatment of gout, hypertension, arteriosclerosis and other inflammatory disorders (Li *et al.*, 2004).

However, burdock has also been reported to have side-effects. The most commonly reported side-effect of burdock is the induction of contact dermatitis. Patients suffered from contact dermatitis after extended topical use of the root oil of burdock. Another reported case was a massage liniment containing burdock extracts had caused contact dermatitis (Paulsen, 2002). There was also case of development of anaphylaxis due to burdock consumption. A Japanese man had developed urticaria 10 times after consuming boiled burdock as food, with redness occurring over his entire body. In addition, he experienced difficulties in breathing an hour after consuming boiled burdock. It was found that this patient had a low blood pressure of 64/29 mmHg. He was diagnosed to be in anaphylactic shock (Sasaki *et al.*, 2003). Therefore, it seems to be a misconception that herbs that are of natural sources have less side-effect comparing to drugs. It was suggested that adverse clinical effects for herbal drugs range from allergic skin reactions, the Stevens-Johnson syndrome and photosensitization to toxic dermatosis. Since most herbs are readily accessible by the general public, increasing number of cases of herb-induced adverse effects is expected (Niggemann and Gruber, 2003). Therefore, public awareness about the possibility of

adverse effects of medicinal herbs shall be enhanced.

Conclusions

Burdock contains many active ingredients (isolated from different parts of the plant) that have been shown to possess many therapeutic effects for the treatment of various diseases. Multiple reports in the literature have demonstrated a wide range of possible clinical uses of this herb because of its anti-inflammatory, anti-tumor/cancer, anti-diabetic, anti-microbial and antiviral effects. In conclusion, the medicinal use of burdock in treating chronic diseases like cancers, diabetes and AIDS is promising. However, it is also essential to be aware of the side-effects of burdock including contact dermatitis and other allergic/inflammatory responses that might be evoked by burdock. It is expected that further investigations will lead to a better understanding of some other roles that *Arctium lappa* plays in preventing and treating of human diseases, as well as the potential adverse effects and toxicity of the herb. That could provide us with more information on the beneficial effect and the potential risk of consuming burdock as a functional food.

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References

- Abreu, P., Matthew, S., Gonzalez, T., *et al.* (2006). Anti-inflammatory and antioxidant activity of a medicinal tincture from *Pedilanthus tithymaloides*, *Life Sci.* **78**, 1578-1585.
- Awale, S., Lu, J., Kalauni, S. K., *et al.* (2006). Identification of arctigenin as an antitumor agent having the ability to eliminate the tolerance of cancer cells to nutrient starvation, *Cancer Res.* **66**, 1751-1757.
- Bhat, S. H., Azmi, A. S., and Hadi, S. M. (2007). Prooxidant DNA breakage induced by caffeic acid in human peripheral lymphocytes: Involvement of endogenous copper and a putative mechanism for anticancer properties, *Toxicol. Appl. Pharm.* **218**, 249-255.
- Bouayed, J., Rammal, H., Dicko, A., *et al.* (2007). Chlorogenic acid, a polyphenol from *Prunus domestica* (Mirabelle), with coupled anxiolytic and antioxidant effects, *J. Neurol. Sci.* **262**, 77-84.
- Bralley, E., Greenspan, P., Hargrove, J. L., *et al.* (2008). Inhibition of hyaluronidase activity by select sorghum brans, *J. Med. Food.* **11**, 307-312.
- Chan, E., Wong, C. Y. K., Wan, C. W., *et al.* (2010). Evaluation of Anti-Oxidant Capacity of Root of *Scutellaria baicalensis* Georgi, in Comparison with Roots of *Polygonum multiflorum* Thunb and *Panax ginseng* CA Meyer, *Am J Chinese Med.* **38**, 815-827.
- Chen, F. A., Wu, A. B., and Chen, C. Y. (2004). The influence of different treatments on the free radical scavenging activity of burdock and variations of its active components, *Food Chem.* **86**, 479-484.
- Chiang, L. C., Chiang, W., Chang, M. Y., *et al.* (2002). Antiviral activity of *Plantago* major extracts and related compounds in vitro, *Antivir. Res.* **55**, 53-62.
- Cho, M. K., Jang, Y. P., Kim, Y. C., *et al.* (2004). Arctigenin, a phenylpropanoid dibenzylbutyrolactone lignan, inhibits MAP kinases and AP-1 activation via potent MKK inhibition: the role in TNF-alpha inhibition, *Int. Immunopharmacol.* **4**, 1419-1429.
- Cho, M. K., Park, J. W., Jang, Y. P., *et al.* (2002). Potent inhibition of lipopolysaccharide-inducible nitric oxide synthase expression by dibenzylbutyrolactone lignans through inhibition of I-kappa B alpha phosphorylation and of p65 nuclear translocation in macrophages, *Int.*

- Immunopharmacol.* **2**, 105-116.
- Eich, E., Pertz, H., Kaloga, M., *et al.* (1996). (-)-Arctigenin as a lead structure for inhibitors of human immunodeficiency virus type-1 integrase, *J. Med. Chem.* **39**, 86-95.
- Gao, Y., Dong, X., Kang, T. G., *et al.* (2002). Activity of in vitro anti-influenza virus of arctigenin. **33**, 724-726.
- Guo, J. F., Zhou, J. M., Zhang, Y., *et al.* (2008). Rhabdastrellic acid-A inhibited PI3K/Akt pathway and induced apoptosis in human leukemia HL-60 cells, *Cell Biol. Int.* **32**, 48-54.
- Hirose, M., Yamaguchi, T., Lin, C., *et al.* (2000). Effects of arctiin on PhIP-induced mammary, colon and pancreatic carcinogenesis in female Sprague-Dawley rats and MeIQx-induced hepatocarcinogenesis in male F344 rats, *Cancer Lett.* **155**, 79-88.
- Ichikawa, K., Kinoshita, T., Nishibe, S., *et al.* (1986). The Ca-2+ Antagonist Activity of Lignans, *Chem. Pharm. Bull.* **34**, 3514-3517.
- Ishihara, K., Yamagishi, N., Saito, Y., *et al.* (2006). Arctigenin from Fructus Arctii is a novel suppressor of heat shock response in mammalian cells, *Cell Stress Chaperon.* **11**, 154-161.
- Iwakami, S., Wu, J. B., Ebizuka, Y., *et al.* (1992). Platelet Activating Factor(Paf) Antagonists Contained in Medicinal-Plants - Lignans and Sesquiterpenes, *Chem. Pharm. Bull.* **40**, 1196-1198.
- Kim, B. H., Hong, S. S., Kwon, S. W., *et al.* (2008). Diarctigenin, a Lignan Constituent from *Arctium lappa*, Down-Regulated Zymosan-Induced Transcription of Inflammatory Genes through Suppression of DNA Binding Ability of Nuclear Factor-kappa B in Macrophages, *J. Pharmacol. Exp. Ther.* **327**, 393-401.
- Knipping, K., van Esch, E., Wijering, S. C., *et al.* (2008). In Vitro and In Vivo Anti-Allergic Effects of *Arctium lappa* L, *Exp. Biol. Med. (Maywood)*. **233**, 1469.
- Lee, C. P., Shih, P. H., Hsu, C. L., *et al.* (2007). Hepatoprotection of tea seed oil (*Camellia oleifera* Abel.) against CCl₄-induced oxidative damage in rats, *Food Chem. Toxicol.* **45**, 888-895.
- Li, Y. J., Liu, S. M., Li, S. L., *et al.* (2004). The Experimental Study of the Effect of Anti-decrepitude of *Arctium lappa* L. **15**, 545-546.
- Li, Y. J., Shi, W., Li, Y. D., *et al.* (2008). Neuroprotective effects of chlorogenic acid against apoptosis of PC12 cells induced by methylmercury, *Environ. Toxicol. Phar.* **26**, 13-21.
- Lin, C. C., Lin, J. M., Yang, J. J., *et al.* (1996). Anti-inflammatory and radical scavenge effects of *Arctium lappa*, *Am. J. Chin. Med.* **24**, 127-137.

- Lin, S. C., Chung, T. C., Lin, C. C., *et al.* (2000). Hepatoprotective effects of *Arctium lappa* on carbon tetrachloride- and acetaminophen-induced liver damage, *Am J Chin Med.* **28**, 163-173.
- Lin, S. C., Lin, C. H., Lin, C. C., *et al.* (2002). Hepatoprotective effects of *Arctium lappa* Linne on liver injuries induced by chronic ethanol consumption and potentiated by carbon tetrachloride, *J. Biomed. Sci.* **9**, 401-409.
- Lin, X. C., Liu, C. Y., Chen, K. S., *et al.* (2004). Extraction and content comparison of chlorogenic acid in *Arctium lappa* L. leaves collected from different terrain and its restraining bacteria test, *Nat. Prod. Res. & Dev.* **16**, 328-330.
- Liu, L., and Tang, L. (1997). Studies on antimutagenicity of Burdock, *Acta Academiae Medicine Nanjing.* **4**, 343-345.
- Matsumoto, T., Hosono-Nishiyama, K., and Yamada, H. (2006). Antiproliferative and apoptotic effects of butyrolactone lignans from *Arctium lappa* on leukemic cells, *Planta Med.* **72**, 276-278.
- Mitsuo, M., Nobuo, Y., and Katsuya, T. (2005). Inhibitory compounds of alpha glucosidase activity from *Arctium lappa* L, *J. Oleo Sci.* **54**, 589-594.
- Miyamoto, K., Nomura, M., Sasakura, M., *et al.* (1993). Antitumor-Activity of Oenothetin-B, a Unique Macrocyclic Ellagitannin, *Jpn. J. Cancer Res.* **84**, 99-103.
- Mizushima, Y., Nakanishi, R., Kuriyama, I., *et al.* (2006). beta-sitosterol-3-O-beta-D-glucopyranoside: A eukaryotic DNA polymerase lambda inhibitor, *J. Steroid Biochem.* **99**, 100-107.
- Morita, T., Ebihara, K., and Kiriya, S. (1993). Dietary Fiber and Fat-Derivatives Prevent Mineral-Oil Toxicity in Rats by the Same Mechanism, *J. Nutr.* **123**, 1575-1585.
- Niggemann, B., and Gruber, C. (2003). Side-effects of complementary and alternative medicine, *Allergy.* **58**, 707-716.
- Pari, L., and Prasath, A. (2008). Efficacy of caffeic acid in preventing nickel induced oxidative damage in liver of rats, *Chem-Biol. Interact.* **173**, 77-83.
- Park, S. Y., Hong, S. S., Han, X. H., *et al.* (2007). Lignans from *Arctium lappa* and their inhibition of LPS-induced nitric oxide production, *Chem. Pharm. Bull.* **55**, 150-152.
- Paulsen, E. (2002). Contact sensitization from Compositae-containing herbal remedies and cosmetics, *Contact Dermatitis.* **47**, 189-198.
- Pereira, J. V., Bergamo, D. C., Pereira, J. O., *et al.* (2005). Antimicrobial activity of *Arctium lappa* constituents against microorganisms commonly found in endodontic infections, *Braz. Dent. J.* **16**, 192-196.
- Pontiki, E., Hadjipavlou-Litina, D., Chaviara, A. T., *et al.* (2006). Evaluation of anti-inflammatory and antioxidant activities of mixed-ligand Cu(II) complexes

- of then and its Schiff dibases with heterocyclic aldehydes and 2-amino-2-thiazoline, *Bioorg. Med. Chem. Lett.* **16**, 2234-2237.
- Rault-Nania, M. H., Demougeot, C., Gueux, E., *et al.* (2008). Inulin supplementation prevents high fructose diet-induced hypertension in rats, *Clin. Nutr.* **27**, 276-282.
- Sasaki, Y., Kimura, Y., Tsunoda, T., *et al.* (2003). Anaphylaxis due to burdock, *Int. J. Dermatol.* **42**, 472-473.
- Schroder, H. C., Merz, H., Steffen, R., *et al.* (1990). Differential in vitro anti-HIV activity of natural lignans, *Z Naturforsch C.* **45**, 1215-1221.
- Silver, A. A., and Krantz Jr, J. C. (1931). The effect of the ingestion of burdock root on normal and diabetic individuals a preliminary report, *Ann. Intern. Med.* **5**, 274.
- Takasaki, M., Konoshima, T., Komatsu, K., *et al.* (2000). Anti-tumor-promoting activity of lignans from the aerial part of *Saussurea medusa*, *Cancer Lett.* **158**, 53-59.
- Takasugi, M., Kawashima, S., Katsui, N., *et al.* (1987). Studies on Stress Metabolites .5. 2 Polyacetylenic Phytoalexins from *Arctium-Lappa*, *Phytochemistry.* **26**, 2957-2958.
- Tamayo, C., Richardson, M. A., Diamond, S., *et al.* (2000). The chemistry and biological activity of herbs used in Flor-Essence (TM) herbal tonic and Essiac (TM), *Phytother. Res.* **14**, 1-14.
- Tsuneki, H., Ma, E. L., Kobayashi, S., *et al.* (2005). Antiangiogenic activity of beta-eudesmol in vitro and in vivo, *Eur. J. Pharmacol.* **512**, 105-115.
- Wang, B. S., Yen, G. C., Chang, L. W., *et al.* (2007). Protective effects of burdock (*Arctium lappa* Linne) on oxidation of low-density lipoprotein and oxidative stress in RAW 264.7 macrophages, *Food Chem.* **101**, 729-738.
- Weber, V., Rubat, C., Duroux, E., *et al.* (2005). New 3-and 4-hydroxyfuranones as anti-oxidants and anti-inflammatory agents, *Bioorgan. Med. Chem.* **13**, 4552-4564.
- Xia, Z. Q., Costa, M. A., Pelissier, H. C., *et al.* (2001). Secoisolariciresinol dehydrogenase purification, cloning, and functional expression - Implications for human health protection, *J. Biol. Chem.* **276**, 12614-12623.
- Xu, Z. H., Wang, X. Y., Zhou, M. M., *et al.* (2008). The antidiabetic activity of total lignan from fructus arctii against alloxan-induced diabetes in mice and rats, *Phytother. Res.* **22**, 97-101.
- Yayli, N., Yasar, A., Gulec, C., *et al.* (2005). Composition and antimicrobial activity of essential oils from *Centaurea sessilis* and *Centaurea armena*, *Phytochemistry.* **66**, 1741-1745.
- Yu, B. S., Yan, X. P., Xiong, J. Y., *et al.* (2003). Simultaneous determination of

chlorogenic acid, forsythin and arctiin in Chinese traditional medicines preparation by reversed phase-HPLC, *Chem. Pharm. Bull.* **51**, 421-424.

Zhao, F., Wang, L., and Liu, K. (2009). In vitro anti-inflammatory effects of arctigenin, a lignan from *Arctium lappa* L., through inhibition on iNOS pathway, *J. Ethnopharmacol.* **122**, 457-462.

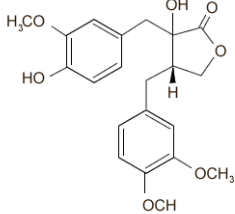
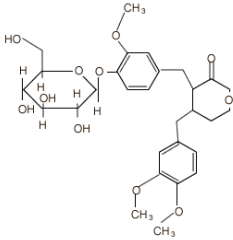
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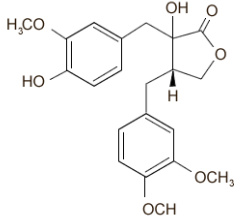
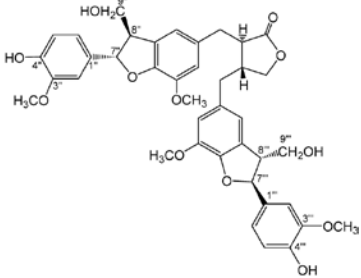
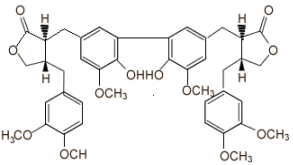
Table 1 General compounds and effects of burdock reported in the literature

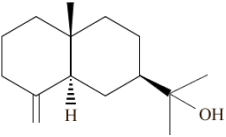
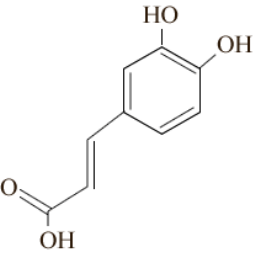
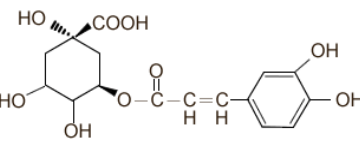
Table 2 Major nutritional ingredients contained in the burdock roots

Fig. 1 The root of burdock

Table 1 General compounds and effects of burdock reported in the literature

Classification	Compound	Molecular Formula	Parts of the plant	Effect	Reference
Lignans	 <p>Arctigenin</p>	$C_{12}H_{24}O_7$	Leaves, fruits, seeds, roots	Suppressor of heat shock; antitumor; Anti-influenza virus	(Ishihara <i>et al.</i> , 2006) (Awale <i>et al.</i> , 2006) (Gao <i>et al.</i> , 2002)
	 <p>Arctiin</p>	$C_{27}H_{34}O_{11}$	Leaves, fruits, roots	Anti-tumor-promoting activity; chemopreventive activity; antiproliferative activity against B cell hybridoma cell, MH60	(Takasaki <i>et al.</i> , 2000) (Hirose <i>et al.</i> , 2000) (Matsumoto <i>et al.</i> , 2006)

	 <p style="text-align: center;">Trachelogenin</p>	$C_{21}H_{24}O_7$	Fruits	Ca^{2+} antagonist activity ; Anti-HIV properties	(Ichikawa <i>et al.</i> , 1986) (Xia <i>et al.</i> , 2001)
	 <p style="text-align: center;">Lappaol F</p>	$C_{40}H_{42}O_{12}$	Fruits, seeds	Inhibiting NO production	(Park <i>et al.</i> , 2007)
	 <p style="text-align: center;">Diarctigenin</p>	$C_{42}H_{46}O_{12}$	Fruits, roots, seeds	Inhibiting NO production;	(Park <i>et al.</i> , 2007)

Terpenoids	 <p data-bbox="660 422 840 454">Beta-eudesmol</p>	C ₁₅ H ₂₆ O	Fruits	Antibacterial, .antiangiogenic	(Yayli <i>et al.</i> , 2005) (Tsuneki <i>et al.</i> , 2005)
Polyphenols	 <p data-bbox="683 805 817 837">Caffeic acid</p>	C ₉ H ₈ O ₄	Stems, leaves, the skin of roots	Antioxidative; free radical scavenging activity	(Pari and Prasath, 2008) (Bhat <i>et al.</i> , 2007)
	 <p data-bbox="660 1101 840 1133">Chlorogenic acid</p>	C ₁₆ H ₁₈ O ₉	Leaves; the skin of roots	Neuroprotective ; Antioxidative ; anti-anaphylaxis and anti-HIV ;	(Li <i>et al.</i> , 2008) (Bouayed <i>et al.</i> , 2007) (Chen <i>et al.</i> , 2004)

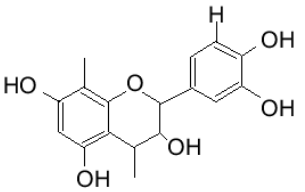
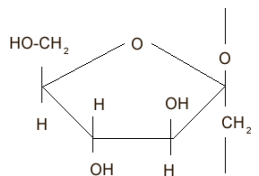
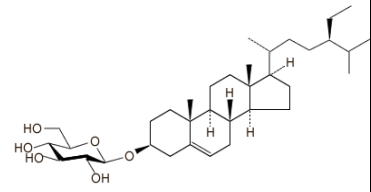
	 <p>Tannin</p>	$C_{76}H_{52}O_{46}$	Roots	Anti-tumor; immuno-modulator; haluronidase inhibition	(Miyamoto <i>et al.</i> , 1993) (Bralley <i>et al.</i> , 2008)
Fructose	 <p>Inulin</p>	$(C_6H_{10}O_5)_n$	Roots	Prebiotic effectiveness ; antihypertension; anti-diabetes	(Li <i>et al.</i> , 2008) (Rault-Nania <i>et al.</i> , 2008) (Silver and Krantz Jr, 1931)
Sterols	 <p>Sitosterol-beta-D-glucopyranoside</p>	$C_{35}H_{60}O_6$	Roots	mammalian DNA polymerase λ ; anti-diabetes and obesity	(Mizushina <i>et al.</i> , 2006) (Silver and Krantz Jr, 1931)

Table 2 Major nutritional ingredients contained in the burdock roots

Types	Nutrient ingredients							
Amino Acid	Essential amino acids		Aspartic acid (25-28%)			Arginine (18-20%)		
Metal elements	Potassium	Calcium	Iron	Magnesium	Manganese	Sodium	Zinc	Copper
Vitamins	B1	B2	C	A				
Others	Crude fiber	Phosphorus	Carotene					



Fig. 1. The root of burdock