

# Sarsaparilla (*Smilax officinalis*) Database file in the Tropical Plant Database of herbal remedies

**Family:** Smilacaceae

**Genus:** *Smilax*

**Species:** *officinalis*, *aristolochiaefolia*, *glabra*, *febrifuga*, *ornata*, *regelii*, *japicanga*

**Synonyms:** *Smilax medica*

**Common Names:** Sarsaparilla, salsaparrilha, khao yen, saparna, smilace, smilax, zarzaparilla, jupicanga

**Part Used:** Root

The following text has been reprinted from: [The Healing Power of Rainforest Herbs](#) © 2005 by [Leslie Taylor](#) .

## SARSAPARILLA

### Herbal Properties and Actions

MAIN ACTIONS	OTHER ACTIONS	STANDARD DOSAGE
<ul style="list-style-type: none"><li>• detoxifies organs</li><li>• cleanses blood</li><li>• aids absorption</li><li>• kills bacteria</li><li>• stimulates digestion</li><li>• increases urination</li><li>• protects liver</li><li>• promotes perspiration</li></ul>	<ul style="list-style-type: none"><li>• relieves pain</li><li>• kills fungi</li><li>• reduces inflammation</li><li>• kills germs</li><li>• reduces fever</li><li>• immunomodulator</li><li>• fights free radicals</li><li>• relieves rheumatism</li></ul>	Root <b>Decoction:</b> 1/2 to 1 cup 2-3 times daily <b>Capsules:</b> 1-2 g twice daily <b>Tincture:</b> 2-3 ml twice daily

---

Sarsaparilla is a brambled, woody vine that grows up to 50 m long, with paired tendrils for climbing (often high into the rainforest canopy). It produces small flowers and black, blue, or red berry-like fruits which are eaten greedily by birds. *Smilax*, a member of the lily family, is native to tropical and temperate parts of the world and comprises about 350 species worldwide. It is native to South America, Jamaica, the Caribbean, Mexico, Honduras, and the West Indies. The name *sarsaparilla* or *zarzaparilla* comes from the Spanish word *zarza* (bramble or bush), *parra* (vine), and *illa* (small)—a small, brambled vine.

The stems of many *Smilax* species are covered with prickles and, sometimes, these vines are cultivated to form impenetrable thickets

(which are called *catbriers* or *greenbriers*). The root, used for medicinal purposes, is long and tuberous—spreading 6-8 feet—and is odorless and fairly tasteless. Many species of *Smilax* around the world share the name *sarsaparilla*; these are very similar in appearance, uses, and even chemical structure. These include *S. officinalis*, *S. japicanga*, and *S. febrifuga* from South America (Brazil, Ecuador and Colombia); *S. regelii*, *S. aristolochiaefolia*, and *S. ornata* from Mexico and Latin America; and *S. glabra* from China. Sarsaparilla vine should not be confused with the large sasparilla and sassafras trees (the root and bark of which were once used to flavor root beer). Sarsaparilla has been used as an ingredient in root beer and other beverages for its foaming properties—not for its flavoring properties.

## **TRIBAL AND HERBAL MEDICINE USES**

Sarsaparilla root has been used for centuries by the indigenous peoples of Central and South America for sexual impotence, rheumatism, skin ailments, and as a general tonic for physical weakness. It has long been used by tribes in Peru and Honduras for headaches and joint pain, and against the common cold. Many shamans and medicine men in the Amazon use sarsaparilla root internally and externally for leprosy and other skin problems (such as psoriasis and dermatitis.) Leprosy can be common in areas where the disease is carried by armadillos (and in the Amazon, armadillos are "on the menu" in indigenous diets). Sarsaparilla root also was used as a general tonic by indigenous tribes in South America, where New World traders found it and introduced it into European medicine in the 1400s.

European physicians considered sarsaparilla root a tonic, blood purifier, diuretic, and sweat promoter. A *Smilax* root from Mexico was introduced into European medicine in 1536, where it developed a strong following as a cure for syphilis and rheumatism. Since this time, *Smilax* roots have had a long history of use for syphilis and other sexually-transmitted diseases throughout the world. With its reputation as a blood purifier, it was registered as an official herb in the *U.S. Pharmacopoeia* as a syphilis treatment from 1820 to 1910. From the 1500s to the present, sarsaparilla has been used as a blood purifier and general tonic and also has been used worldwide for gout, syphilis, gonorrhoea, rheumatism, wounds, arthritis, fever, cough, scrofula, hypertension, digestive disorders, psoriasis, skin diseases, and cancer.

## **PLANT CHEMICALS**

Sarsaparilla contains the plant steroids sarsasapogenin, smilagenin, sitosterol, stigmasterol, and pollinastanol; and the saponins sarsasaponin, smilasaponin, sarsaparilloside, and sitosterol glucoside, among others. The majority of sarsaparilla's pharmacological properties and actions have been attributed to these steroids and saponins. The saponins have been reported to facilitate the body's absorption of other drugs and phytochemicals, which accounts for its history of use in herbal formulas as an agent for bioavailability and to enhancement the power and effect of other herbs.

Saponins and plant steroids found in many species of plants (including sarsaparilla) can be synthesized into human steroids such as

estrogen and testosterone. This synthesis has never been documented to occur in the human body - only in the laboratory. Yet plant steroids and their actions in the human body have been a subject of much interest, sketchy research and, unfortunately, disinformation - mainly for marketing purposes. Sarsaparilla has been marketed (fraudulently) to contain testosterone and/or other anabolic steroids. While it is a rich source of natural plant steroids and saponins, it never has been proven to have any anabolic effects, nor has testosterone been found in sarsaparilla or any other plant source thus far.

Flavonoids in sarsaparilla have been documented to have immune modulation and liver protective activities. A U.S. patent was awarded in 2003 describing these flavonoids to be effective in treating autoimmune diseases and inflammatory reactions through their immunomodulating effects. Sarsasapogenin and smilagenin were subjects of a 2001 U.S. patent which reported that these Smilax steroids had the ability to treat senile dementia, cognitive dysfunction, and Alzheimer's disease. In the patent's animal studies references, smilagenin reversed the decline of brain receptors in aged mice and restored the receptor levels to those observed in young animals, reversed the decline in cognitive function, and enhanced memory and learning. These studies, however, have not been published in any peer-reviewed journals - only in the context of the patent, thus far.

Sarsaparilla's main plant chemicals include: acetyl-parigenin, astilbin, beta-sitosterol, caffeoyl-shikimic acids, dihydroquercetin, diosgenin, engeletin, essential oils, epsilon-sitosterol, eucryphin, eurryphin, ferulic acid, glucopyranosides, isoastilbin, isoengenitin, kaempferol, parigenin, parillin, pollinastanol, resveratrol, rhamnose, saponin, sarsapogenin, sarsaparilloside, sarsapogenin, sarsasapogenin, shikimic acid, sitosterol-d-glucoside, smilagenin, smilasaponin, smilax saponins A-C, smiglaside A-E, smitilbin, stigmasterol, taxifolin, and titogenin.

## **BIOLOGICAL ACTIVITIES AND CLINICAL RESEARCH**

Clinical research has validated the traditional use of sarsaparilla for skin conditions such as psoriasis, eczema, acne, and leprosy. In 1942, it was reported in the *New England Journal of Medicine* to improve the condition of psoriasis dramatically. There the results of a clinical study with 92 patients was detailed which reported that it improved psoriasis lesions in 62% of cases and completely cleared lesions in 18% of cases. One of the possible mechanisms of action in psoriasis is sarsaparilla's blood cleansing properties. Individuals with psoriasis have been found to have high levels of endotoxins circulating in the bloodstream (endotoxins are cell wall fragments of normal gut bacteria). Sarsapogenin, one of sarsaparilla's main steroids, was found to bind to these endotoxins and remove them, thus improving psoriasis.

This endotoxin-binding action is probably why the root has been used for centuries as a "blood purifier." Other health conditions associated with high endotoxin levels include eczema, arthritis, and ulcerative colitis. Sarsaparilla's effective use in the treatment of leprosy has been documented in a 1959 human trial. The effectiveness of sarsaparilla in the treatment of adolescent acne caused by excessive androgens has received some experimental support as well.

A 2001 U.S. patent was filed on sarsaparilla (*Smilax china*) for psoriasis and respiratory diseases. This patent cited clinical observations and studies with children and human adults with *Psoriasis vulgaris*, pustular psoriasis, erythroderma psoriaticum lesions, and associated itching-reporting marked clinical improvements with dosages of 3-6 g daily. It also reported that, upon

discontinuation of sarsaparilla after only two months of treatment, there was further gradual remission of lesions and no side effects. In addition, this patent indicated sarsaparilla was shown to be a preventative and therapeutic agent for respiratory and allergic diseases such as acute bronchitis, bronchial asthma, asthmatic bronchitis, and chronic bronchitis. Again, these studies and observations reported in the patent have yet to be published in any peer-reviewed journals.

Sarsaparilla has long been used in the treatment of syphilis. Clinical observations in China demonstrated that sarsaparilla was effective (according to blood tests) in about 90% of acute and 50% of chronic cases. In the 1950s the antibiotic properties of sarsaparilla were documented; other studies documented its antifungal and antimycobacterial activities. Its anti-inflammatory activity has been demonstrated in several in vitro and in vivo studies, using different laboratory-induced models of arthritis and inflammation. One of these studies attributes the beneficial effect for arthritis to sarsaparilla's immune modulatory action. Sarsaparilla also has demonstrated liver protective effects in rats, with researchers concluding that it is able to prevent immune-mediated liver injury. Improvement of appetite and digestion has been noted with sarsaparilla, as well as its diuretic actions in humans. The root has been reported to have stimulatory activity on the kidneys in humans and, in chronic nephritis, it was shown to increase the urinary excretion of uric acid.

## **CURRENT PRACTICAL USES**

Sarsaparilla is becoming more widely available in health food stores, with a variety of tablets, capsules, and tincture products sold today. Most of the sarsaparilla root in herbal commerce today comes from cultivation projects in Mexico and Latin America as well as China. In naturopathic and herbal medicine, it is used mostly in combination with other herbs for its tonic, detoxifying, blood purifying, and lymph-cleansing properties. In retail stores and products, it can be found as an ingredient in various herbal remedies made for skin disorders, libido enhancement, hormone balancing, and sports nutrition formulas. It's also commonly used in herbal preparations as a synergist or bioavailability aid - as it is thought that the saponins in sarsaparilla root increase the absorption of other chemicals in the gut. No known toxicity or side-effects have been documented for sarsaparilla; however, ingestion of large dosages of saponins may cause gastrointestinal irritation.

### **Sarsaparilla Plant Summary**

**Main Preparation Method:** capsules or decoction

**Main Actions (in order):**

blood cleanser, immunomodulator (selectively reduces overactive immune cells), antimutagenic (cellular protector), detoxifier, tonic (tones, balances, strengthens overall body functions)

**Main Uses:**

1. for psoriasis, dermatitis, leprosy, and other skin disorders
2. as a blood purifier and general detoxification aid

3. 3. as a general tonic (tones, balances, strengthens), stimulant, and hormonal regulator
4. 4. for arthritis, rheumatism and autoimmune disorders which cause inflammation
5. 5. for syphilis and other sexually transmitted diseases

**Properties/Actions Documented by Research:**

anti-inflammatory, antibacterial, antifungal, antimutagenic (cellular protector), blood cleanser, detoxifier, diuretic, hepatoprotective (liver protector), immunomodulator (selectively reduces overactive immune cells), neuroprotective (protects brain cells)

**Other Properties/Actions Documented by Traditional Use:**

absorption aid, analgesic (pain-reliever), anticancerous, antioxidant, antirheumatic, antiseptic, aphrodisiac, diaphoretic (promotes sweating), digestive stimulant, febrifuge (reduces fever), stimulant, tonic (tones, balances, strengthens), wound healer

**Cautions:** Excessive dosages can cause gastrointestinal irritation.

**Traditional Preparation:** One-half to 1 cup of a standard root decoction 2-3 times daily. Alternatively, 1-2 grams of root powder in tablets or capsules twice daily or 2-3 ml of a standard tincture or fluid extract may be taken twice daily.

**Contraindications:** Large doses may cause gastrointestinal upset.

**Drug Interactions:** Sarsaparilla may increase the absorption of some drugs and compounds. Some report that it can increase the absorption of Digitalis glycosides while accelerating the elimination of hypnotic drugs.

**Worldwide Ethnomedical Uses**

- Argentina** for rheumatism, and to increase perspiration and libido
- Brazil** for acne, anorexia, arthritis, blood purification, digestive disorders, eczema, fever, gallstones, gout, hives, kidney problems, kidney stones, impotence, leprosy, muscle weakness, psoriasis, rheumatism, skin disorders, sterility, syphilis, ulcers, urinary insufficiency, venereal diseases, and as an aphrodisiac, laxative, and to increase perspiration
- China** for abscesses, arthritis, boils, cystitis, diarrhea, digestive disorders, dysentery, enteritis, fever, malaria, mercury poisoning, rheumatism, rheumatoid arthritis, skin problems, sores, syphilis, urinary insufficiency, and as an aphrodisiac and tonic
- England** for abscesses, anorexia, antiseptic, blood cleansing, cancer, dysentery, eczema, fatigue, gout, immune enhancement, impotence, infections, inflammation, itching, leprosy, mercury poisoning, muscle weakness, PMS, psoriasis, rheumatism, rheumatoid arthritis, skin problems, syphilis, tonic, venereal diseases and as an aphrodisiac, antiseptic, diuretic, and to increase perspiration
- Europe** or arthritis, inflammation, kidney problems, psoriasis, rheumatism, skin problems, sweat promotion, urinary disorders, urinary insufficiency, venereal diseases
- Latin America** for aches, acne, arthritis, colds, digestive disorders, fever, gout, impotence, pain, psoriasis, rheumatism, skin problems, sweat promotion, syphilis, venereal disease, weakness, and as an aphrodisiac and tonic

<b>Mexico</b>	for arthritis, blood purification, burns, cancer, digestive disorders, dyspepsia, eczema, fever, gonorrhoea, inflammation, leprosy, nephritis, rash, rheumatism, scrofula, skin problems, syphilis, venereal disease, and to increase perspiration and urination
<b>United States</b>	for acne, arthritis, bladder problems, blood purification, burns, cancer, convalescence, coughs, diabetes, digestive disorders, eczema, eye infections, fatigue, fever, gonorrhoea, gout, herpes, hives, hypertension, impotence, infertility, inflammation, itching, kidney problems, laxative, liver protection, pleurisy, PMS, psoriasis, rheumatism, scrofula, shingles, skin problems, stomach disorders, stress, sweat promotion, syphilis, tuberculosis, ulcers, ulcerative colitis, urinary disorders, urinary insufficiency, vaginal discharge, venereal disease, warts, wounds, and as an expectorant
<b>Elsewhere</b>	for abscesses, arthritis, asthma, boils, burns, cancer, colds, conjunctivitis, cystitis, dyspepsia, dysentery, eczema, edema, epilepsy, gonorrhoea, gout, herpes, impotence, inflammation, intestinal gas, kidney problems, leprosy, lymph inflammation, malaria, menstrual disorders, psoriasis, rashes, stimulant, tonic, toothache, tumor, urogenital diseases, venereal disease, wounds, and as an aphrodisiac, stimulant, and tonic

---

**The above text has been reprinted from**

***[The Healing Power of Rainforest Herbs](#)* by Leslie Taylor, copyrighted 2005**

All rights reserved. No part of this document may be reproduced or transmitted in any form or by any means, electronic or mechanical, including photocopying, recording, or by any information storage or retrieval system, including websites, without written permission. Please read the [Conditions of Use](#), and [Copyright Statement](#) for this web page and web site.

## **Referenced Quotes on Sarsaparilla**

1. "Since ancient times Sarsaparilla has been considered an excellent blood purifier. More recently it has been used alone or with other herbs to combat psoriasis, eczema, warts and other skin infections. In homeopathy it is frequently used for skin eruptions that are accompanied by intense itching."
2. "Sarsaparilla is used all over the world for a wide range of symptoms. These include lung and stomach congestion, skin diseases, herpes, syphilis, psoriasis, arthritis, rheumatism, gout, nervous disorders, epilepsy, chronic liver disorders, colds fevers, and stomach and intestinal gas. It helps promote good circulation, clear toxins, balance the glandular system, and stimulate metabolism and male sexual potency. The natural steroidal glycosides found in smilax make it a favorite for body builders and anyone who would like to be stronger and energized. It promotes rejuvenation and can be used as an anti-inflammatory."
3. "ACTIONS: Stimulates metabolism, Enhances glandular balance, Boosts hormone, production, Cleanses blood, Aids muscle building. TRADITIONAL USE: Smilax is used for glandular balance, and is believed to aid building of muscle mass. Recognized for its properties believed to increase metabolic rate. Naturally stimulates production of hormones. Known to encourage muscle mass when used adjunct to specific exercise. High in vitamins and minerals. Has been used in the treatment of gout, rheumatism, kidney and bladder dysfunctions and skin conditions. MERIDIAN INDICATIONS: Warms male organs, Strengthens muscles, Nourishes blood, kidney, Stomach, Liver EVA POINTS: Triple Warmer, Bladder 65, Stomach, Kidney, Liver."

4. "Medical Action and Uses: Alterative, tonic. Used in chronic skin diseases, rheumatism, passive dropsy."

5. "A sweet herb used for impotency, liver problems, stress, rheumatism, gout, venereal disease (i.e., syphilis), leukorrhea, herpes, other disorders caused by blood impurities, epilepsy, and nervous system disorders. Reduces fever, clears skin disorders such as eczema and psoriasis, and controls diabetes. Also good for stomach and kidney disorders. Regulates hormones, increases energy, and protects against harmful radiation."

11. "Sarsaparilla to increase circulation, clear toxins and stimulate metabolism."

13. "Sarsaparilla's medicinal use has been as a tonic and a blood purifier. A blood purifier or depurative refers to an agent that cleanses and purifies the system. Sarsaparilla's reputation in this regard probably stems from its importation from the Caribbean and South America to Europe in the sixteenth century for the treatment of syphilis.

During military operations in Portugal in 1812, a British Inspector General of Hospitals noted that the Portuguese soldiers suffering from syphilis who used sarsaparilla recovered much faster and more completely than their British counterparts, who were treated with mercury.

Sarsaparilla was also used by the Chinese in the treatment of syphilis. Clinical observations in China demonstrated that sarsaparilla is effective, according to blood tests, in about 90 percent of acute cases and 50 percent of chronic cases.

An interesting note is that sarsaparilla species have been used all over the world in many different cultures for the same conditions, namely gout, arthritis, fevers, digestive disorders, skin disease, and cancer.

The mechanism of action of sarsaparilla is largely unknown, although the plant does contain several saponins and has been shown to be clinically effective in the treatment of psoriasis. This evidence points to a possible effect on binding of cholesterol and bacterial toxins in the intestines.

Evidence seems to support sarsaparilla as an endotoxin binder. Endotoxins are cell wall constituents of bacteria that are absorbed from the gut. Normally, the liver filters out these and other gut-derived compounds before they reach the general circulation. If the amount of endotoxin absorbed is excessive or if the liver is not functioning adequately, the liver can become overwhelmed, and endotoxins will spill into the blood. In a controlled study of ninety-two patients, an endotoxin-binding saponin (sarsaponin) from sarsaparilla greatly improved the psoriasis in 62 percent of the patients and resulted in complete clearance in 18 percent."

14. "Smilax is another herb with a reputation as a remarkable tonic and male rejuvenator. This herb has been receiving considerable attention lately, but not of a research nature. Rather it has become a favorite of the body-building crowd. Is there any justification for this interest?

As a tonic, smilax has been used primarily to increase vitality and virility, and is used throughout Central and South America and some parts of southern North America to treat the symptoms of sexually transmitted diseases. This practice has even spread to Europe. Chinese physicians verified the antisiphilic property of sarsaparilla. In clinical observations, its effectiveness on primary syphilis was rated at 90 percent. Allowing for some halo effect, the results are still staggering. This may help explain how the herb came to be used that way in Europe. It was simply effective.

Worldwide opinion also concurs on other uses for smilax, including an anti-inflammatory effect in arthritis and gout, and a detoxification effect in cancer and skin disorders.

The effectiveness of smilax in the treatment of skin disorders, such as the acne of adolescence caused by raging androgens, has received some experimental support.

The tonic effect of smilax may be the result of its ability to stimulate the removal of accumulated waste products from the cells, blood and lymph.

One final note: Smilax contains a wide variety of saponins, mainly sarsasapogenin, smilagenin, sitosterol and stigmasterol. These substances commonly occur in plants with immune enhancing action, in the adaptogens, in tonics, and in herbs used for their nutritive value. Smilax saponins have not been investigated thoroughly, but they may hold the key to the popularity of the plant for body-building purposes. Sarsaparilla saponins have, for example, been used in the synthesis of sex hormones."

**Hobbs, Christopher, "Sarsaparilla, A literature review" HerbalGram No. 17 - Summer 1988, Pg 10. :**

"Gerard, in his Great Herbal, mentions that the Honduran and Peruvian sarsaparilla "are a remedy against long continual pain of the joints and head, and against the cold." Spanish "sarza parilia", *S. aspera*, he takes to be similar, but weaker in action(11).

According to Monardes, the Spanish botanist, Mexican sarsaparilla was introduced into European medicine about 1536 at Seville (12). Other species soon followed from Guatamala and Honduras. They were highly regarded as a remedy for syphilis, which was also imported from the new world in the late 1400's, and for rheumatism. From Spain, the herb found its way into the pharmacists shops all over Europe and England.

Few plants have had the rise and fall in popularity that sarsaparilla has had. When it was introduced it was considered remarkably effective for diverse chronic diseases, and many doctors of the time wrote about its benefits. Generally considered an alterative tonic, blood purifier, diuretic and diaphoretic, it was given alone or in combination with other herbs, as well as with mercury for long-standing venereal disease.

Pereira, a leading physician in London in the mid-nineteenth century, felt that sarsaparilla works when "the malady is of long continuance, and the constitution is enfeebled and emaciated, either by the repeated attacks of the disease, or by the use of mercury," and that it is "the great restorer of a appetite, flesh, colour, strength and vigour."

11. Gerad, John (1633). *The Herbal or General History of Plants*, reprinted by Dover, NY (1975)

12. Lloyd, J. U. (1929) *Origin and History of all the Pharmacopeial Vegetable Drugs*, Caxton Press, Cincinnati"

---

## **Published Research on Sarsaparilla**

All available third-party research on sarsaparilla can be found at [PubMed](#). A partial listing of the published third party research on sarsaparilla updated through Feb 2019 is shown below:

### **Anti-inflammatory & Anti-Arthritic Actions:**

Dong, L., et al. "Corrigendum to "Astilbin from *Smilax glabra* Roxb. attenuates inflammatory responses in complete Freund's adjuvant-induced arthritis rats." *Evid. Based Complement. Alternat. Med.* 2018 Dec; 2018: 6279328.

Tetty, C., et al. "*Smilax china* leaf extracts suppress pro-inflammatory adhesion response in human umbilical vein endothelial cells and proliferation of HeLa cells." *Arch. Physiol. Biochem.* 2018 Oct:1-5.

Xie, Y., et al. "Anti-inflammatory furostanol saponins from the rhizomes of *Smilax china* L." *Steroids.* 2018 Dec; 140: 70-76.

Bao, Y., et al. "Therapeutic effects of *Smilax glabra* and *Bolbostemma paniculatum* on rheumatoid arthritis using a rat paw edema



model." *Biomed. Pharmacother.* 2018 Dec; 108: 309-315.

Shu, J., et al. "Three new flavonoid glycosides from *Smilax glabra* and their anti-inflammatory activity." *Nat. Prod. Res.* 2018 Aug; 32(15): 1760-1768.

Dong, L., et al. "Astilbin from *Smilax glabra* Roxb. attenuates inflammatory responses in complete freund's adjuvant-induced arthritis rats." *Evid. Based Complement. Alternat. Med.* 2017; 2017: 8246420.

Tian, L., et al. "Steroidal Saponins from the genus *Smilax* and their biological activities." *Nat. Prod. Bioprospect.* 2017 Aug; 7(4): 283-298.

Zhong, C., et al. "Phenolic compounds from the rhizomes of *Smilax china* L. and their anti-inflammatory activity." *Molecules.* 2017 Apr; 22(4).

Lu, C., et al. "Polysaccharides from *Smilax glabra* inhibit the pro-inflammatory mediators via ERK1/2 and JNK pathways in LPS-induced RAW264.7 cells." *Carbohydr. Polym.* 2015 May; 122: 428-36.

Lu, C., et al. "Optimization of astilbin extraction from the rhizome of *Smilax glabra*, and evaluation of its anti-inflammatory effect and probable underlying mechanism in lipopolysaccharide-induced RAW264.7 macrophages." *Molecules.* 2015 Jan; 20(1): 625-44.

Lu, C., et al. "Antioxidant and anti-inflammatory activities of phenolic-enriched extracts of *Smilax glabra*." *Evid. Based Complement. Alternat. Med.* 2014; 2014: 910438.

Luo, Y., et al. "[Effect of *Smilax china* bioactive fraction on tumor necrosis factor- $\alpha$  and interleukin-4 contents in uterine tissue of rats with chronic pelvic inflammatory disease]." *Nan. Fang Yi Ke Da Xue Xue Bao.* 2014 Feb; 34(2): 236-40.

Ma, Y., et al. "[Pharmaceutical screening of the effective fraction from *Smilax* for treatment of chronic pelvic inflammatory disease]." *Nan Fang Yi Ke Da Xue Xue Bao.* 2013 Jan; 33(1): 145-9.

He, X., et al. "[Comparison of anti-inflammatory effect and analysis of astilbin red and white transverse section *Smilax glabra* in 28 collection sites]." *Zhongguo Zhong Yao Za Zhi.* 2012 Dec; 37(23): 3595-8.

Liagre, B., et al. "Inhibition of human rheumatoid arthritis synovial cell survival by hecogenin and tigogenin is associated with increased apoptosis, p38 mitogen-activated protein kinase activity and upregulation of cyclooxygenase-2." *Int. J. Mol. Med.* 2007 Oct; 20(4): 451-60.

Shao, B., et al. "Steroidal saponins from *Smilax china* and their anti-inflammatory activities." *Phytochemistry.* 2007 Mar; 68(5): 623-30.

Shu, X., et al. "The anti-inflammation effects of *Smilax china* ethylacetate extract in rats and mice." *Zhongguo. Zhong. Yao. Za. Zhi.* 2006 Feb; 31(3): 239-43.

Shu, X., et al. "Anti-inflammatory and anti-nociceptive activities of *Smilax china* L. aqueous extract." *J. Ethnopharmacol.* 2006 Feb; 103(3): 327-32.

Ji, W., et al. "Effects of Rebixiao granules on blood uric acid in patients with repeatedly attacking acute gouty arthritis." *Chin. J. Integr. Med.* 2005 Mar; 11(1): 15-21.

Jiang, J., et al. "Immunomodulatory activity of the aqueous extract from rhizome of *Smilax glabra* in the later phase of adjuvant-induced arthritis in rats." *J. Ethnopharmacol.* 2003; 85(1): 53-9.

Ageel, A., et al. "Experimental studies on antirheumatic crude drugs used in Saudi traditional medicine." *Drugs Exp. Clin. Res.* 1989; 15(8): 369-72.

### **Kidney Protecting Actions:**

Huang, L., et al. "The anti-hyperuricemic effect of four astilbin stereoisomers in *Smilax glabra* on hyperuricemic mice." *J. Ethnopharmacol.* 2019 Mar 6. pii: S0378-8741

Wang, S., et al. "The flavonoid-rich fraction from rhizomes of *Smilax glabra* Roxb. ameliorates renal oxidative stress and inflammation in uric acid nephropathy rats through promoting uric acid excretion." *Biomed. Pharmacother.* 2019 Mar; 111: 162-168.

Luo, Q., et al. "Total flavonoids from *Smilax glabra* Roxb blocks epithelial-mesenchymal transition and inhibits renal interstitial fibrosis by targeting miR-21/PTEN signaling." *J. Cell. Biochem.* 2019 Mar; 120(3): 3861-3873.

Liu, Y., et al. "Protective effects of sarsasapogenin against early stage of diabetic nephropathy in rats." *Phytother. Res.* 2018 Aug; 32(8): 1574-1582.

Wang, M, Zhao., et al. "Astilbin improves potassium oxonate-induced hyperuricemia and kidney injury through regulating oxidative stress and inflammation response in mice." *Biomed. Pharmacother.* 2016 Oct; 83: 975-988.

Chen, L., et al. "Astilbin attenuates hyperuricemia and ameliorates nephropathy in fructose-induced hyperuricemic rats." *Planta Med.* 2011 Nov; 77(16): 1769-73.

Chen, L., et al. "Anti-hyperuricemic and nephroprotective effects of *Smilax china* L." *J. Ethnopharmacol.* 2011 May; 135(2): 399-405.

Li, G., et al. "Effect of astilbin on experimental diabetic nephropathy *in vivo* and *in vitro*." *Planta Med.* 2009 Nov; 75(14): 1470-5.

Humpert, F. "The effect of a sarsaparilla preparation (renotrat) in chronic nephritis, with particular reference to the uric acid content of the blood and urine." *Klin. Wochschr.* 1933; 12: 1696.

Rittmann, R., al. "A new agent in kidney therapy." *Klin. Wochschr.* 1930; 9: 401-8.

#### **Liver Protecting Actions:**

Murali, A., et al. "Effect of *Smilax zeylanica* roots and rhizomes in paracetamol induced hepatotoxicity." *J. Complement. Integr. Med.* 2012 Nov 9; 9(1).

Rafatullah, S., et al. "Hepatoprotective and safety evaluation studies on sarsaparilla." *Int. J. Pharmacognosy* 1991; 29: 296-301.

#### **Immune Stimulant & Immune Modulating Actions:**

Wang, Y., et al. "Smiglaside A ameliorates LPS-induced acute lung injury by modulating macrophage polarization via AMPK-PPAR $\gamma$  pathway." *Biochem. Pharmacol.* 2018 Oct; 156: 385-395.

Lim, S., et al "Timosaponin AIII and its metabolite sarsasapogenin ameliorate colitis in mice by inhibiting NF- $\kappa$ B and MAPK activation and restoring Th17/Treg cell balance." *Int. Immunopharmacol.* 2015 Apr; 25(2): 493-503.

Zheng, Z., et al. "Macrophage biospecific extraction and HPLC-ESI-MSn analysis for screening immunological active components in *Smilacis Glabrae Rhizoma*." *J. Pharm. Biomed. Anal.* 2013 Apr; 77: 44-8.

Wang, M., et al. "Structural characterization and macrophage immunomodulatory activity of a novel polysaccharide from *Smilax glabra* Roxb." *Carbohydr. Polym.* 2017 Jan; 156: 390-402.

Yi, H., et al. "Astilbin inhibits the adhesion of T lymphocytes via decreasing TNF-alpha and its associated MMP-9 activity and CD44 expression." *Int. Immunopharmacol.* 2008 Oct; 8(10): 1467-74.

Guo, J., et al. "Identification of a new metabolite of astilbin, 3'-O-methylastilbin, and its immunosuppressive activity against contact dermatitis." *Clin Chem.* 2007 Mar; 53(3): 465-71.

Spelman, K., et al. "Modulation of cytokine expression by traditional medicines: a review of herbal immunomodulators." *Altern. Med. Rev.* 2006 Jun; 11(2): 128-50.

Chu, K., et al. "Smilaxin, a novel protein with immunostimulatory, antiproliferative, and HIV-1-reverse transcriptase inhibitory activities from fresh *Smilax glabra* rhizomes." *Biochem. Biophys. Res. Commun.* 2006 Feb; 340(1): 118-24.

Wang, J., et al. "Astilbin prevents concanavalin A-induced liver injury by reducing TNF-alpha production and T lymphocytes adhesion." *J. Pharm. Pharmacol.* 2004; 56(4): 495-502.

Jiang, J., et al. "Immunomodulatory activity of the aqueous extract from rhizome of *Smilax glabra* in the later phase of adjuvant-induced arthritis in rats." *J. Ethnopharmacol.* 2003; 85(1): 53-9.

Chen, T., et al. "A new flavanone isolated from *Rhizoma smilacis glabrae* and the structural requirements for its derivatives for preventing immunological hepatocyte damage." *Planta Med.* 1999; 65(1): 56-9.

Santos, W., et al. "Haemolytic activities of plant saponins and adjuvants. Effect of *Periandra mediterranea* saponin on the humoral response to the FML antigen of *Leishmania donovani*." *Vaccine* 1997; 15(9): 1024-29.

#### **Anti-Psoriasis & Anti-Leprosy Actions:**

Di, T., et al. "Astilbin inhibits Th17 cell differentiation and ameliorates imiquimod-induced psoriasis-like skin lesions in BALB/c mice via Jak3/Stat3 signaling pathway." *Int. Immunopharmacol.* 2016 Mar; 32: 32-38.

Vijayalakshmi, A., et al. "Screening of flavonoid "quercetin" from the rhizome of *Smilax china* Linn. for anti-psoriatic activity." *Asian Pac J. Trop. Biomed.* 2012 Apr; 2(4): 269-75.

Juhlin, L., et al. "The influence of treatment and fibrin microclot generation in psoriasis." *Br. J. Dermatol.* 1983; 108: 33-7.

Rollier, R. "Treatment of lepromatous leprosy by a combination of DDS and sarsaparilla (*Smilax ornata*)." *Int. J. Leprosy* 1959; 27: 328-40.

Thurman, F. "The treatment of psoriasis with sarsaparilla compound." *New England Journal of Medicine* 1942; 337: 128-33.

#### **Antioxidant Actions:**

Zhu, Y., et al. "Neuroprotective effects of astilbin on MPTP-induced Parkinson's disease mice: Glial reaction,  $\alpha$ -synuclein expression and oxidative stress." *Int. Immunopharmacol.* 2019 Jan; 66: 19-27.

Lee, D., et al. "Using phytochemicals to investigate the activation of nicotine detoxification via upregulation of CYP2A6 in Animal models exposed tobacco smoke condensate by intratracheal instillation." *Evid. Based Complement. Alternat. Med.* 2018 Dec; 2018: 7635197.

Lee, H., et al. "Chemical constituents of *Smilax china* L. stems and their inhibitory activities against glycation, aldose reductase,  $\alpha$ -glucosidase, and lipase." *Molecules.* 2017 Mar; 22(3).

Wang, Y., et al. "Smiglaside A ameliorates LPS-induced acute lung injury by modulating macrophage polarization via AMPK-PPAR $\gamma$  pathway." *Biochem. Pharmacol.* 2018 Oct; 156: 385-395.

Lincha, V., et al. "Effects of constituent compounds of *Smilax china* on nicotine-induced endothelial dysfunction in human umbilical vein endothelial cells." *Biol. Pharm. Bull.* 2016; 39(6): 984-92.

Kong, G., et al. "Astilbin alleviates LPS-induced ARDS by suppressing MAPK signaling pathway and protecting pulmonary endothelial glycocalyx." *Int. Immunopharmacol.* 2016 Jul; 36: 51-58.

Cai, Y., et al. "Medicinal effect and its JP2/RyR2-based mechanism of *Smilax glabra* flavonoids on angiotensin II-induced hypertrophy model of cardiomyocytes." *J. Ethnopharmacol.* 2015 Jul; 169: 435-40.

Lu, C., et al. "Inhibitory effects of chemical compounds isolated from the rhizome of *Smilax glabra* on nitric oxide and tumor necrosis factor- $\alpha$  production in lipopolysaccharide-induced RAW264.7 cell." *Evid. Based Complement. Alternat. Med.* 2015; 2015: 602425.

Lu, C., et al. "Antioxidant and anti-inflammatory activities of phenolic-enriched extracts of *Smilax glabra*." *Evid. Based Complement. Alternat. Med.* 2014; 2014: 910438.

Kim, K., et al. "*Smilax china* root extract detoxifies nicotine by reducing reactive oxygen species and inducing CYP2A6." *J. Food Sci.* 2014 Oct; 79(10): H2132-9.

Park, G., et al. "Antioxidant effects of the sarsaparilla via scavenging of reactive oxygen species and induction of antioxidant enzymes in human dermal fibroblasts." *Environ. Toxicol. Pharmacol.* 2014 Jul; 38(1): 305-15.

Sang, H., et al. "The protective effect of *Smilax glabra* extract on advanced glycation end products-induced endothelial dysfunction in HUVECs via RAGE-ERK1/2-NF- $\kappa$ B pathway." *J. Ethnopharmacol.* 2014 Aug; 155(1): 785-95.

Yoon, S., et al. "Fermentation of *Smilax china* root by *Aspergillus usami* and *Saccharomyces cerevisiae* promoted concentration of resveratrol and oxyresveratrol and the free-radical scavenging activity." *J. Sci. Food Agric.* 2014 Jul; 94(9): 1822-6.

Diao, H., et al. "Astilbin protects diabetic rat heart against ischemia-reperfusion injury via blockade of HMGB1-dependent NF- $\kappa$ B signaling pathway." *Food Chem. Toxicol.* 2014 Jan; 63: 104-10.

Xia, D., et al. "Protective effect of *Smilax glabra* extract against lead-induced oxidative stress in rats." *J. Ethnopharmacol.* 2010 Jul; 130(2): 414-20.

Ranilla, L., et al. "Phenolic compounds, antioxidant activity and *in vitro* inhibitory potential against key enzymes relevant for hyperglycemia and hypertension of commonly used medicinal plants, herbs and spices in Latin America." *Bioresour. Technol.* 2010 Jun; 101(12): 4676-89.

Ma, D., et al. "Effect of sarsasapogenin and its derivatives on the stimulus coupled responses of human neutrophils." *Clin. Chim. Acta.* 2001 Dec; 314(1-2): 107-12.

#### **Antidepressant, Memory Enhancement, & Anti-Alzheimer's Actions:**

He, X., et al. "Smilagenin protects dopaminergic neurons in chronic MPTP/probenecid-lesioned Parkinson's disease models." *Front. Cell. Neurosci.* 2019 Feb 5;13:18.

Dong, D., et al. "Astrocytes mediated the nootropic and neurotrophic effects of Sarsasapogenin-AA13 via upregulating brain-derived neurotrophic factor." *Am. J. Transl. Res.* 2017 Sep; 9(9): 4015-4025.

Huang C., et al. "Sarsasapogenin-AA13 ameliorates A $\beta$ -induced cognitive deficits via improving neuroglial capacity on A $\beta$  clearance and antiinflammation." *CNS Neurosci. Ther.* 2017 Jun; 23(6): 498-509.

Feng, B., et al. "Sarsasapogenin reverses depressive-like behaviors and nicotinic acetylcholine receptors induced by olfactory bulbectomy." *Neurosci. Lett.* 2017 Feb; 639: 173-178.

Sy, L., et al. "Identification of "sarsasapogenin-aglyconed" timosaponins as novel A $\beta$ -lowering modulators of amyloid precursor protein processing." *Chem. Sci.* 2016 May; 7(5): 3206-3214.

Li, J., et al. "Reversal of dopamine neurons and locomotor ability degeneration in aged rats with smilagenin." *Neuroscience.* 2013 Aug; 245: 90-8.

Zhang, R., et al. "Smilagenin attenuates beta amyloid (25-35)-induced degeneration of neuronal cells via stimulating the gene expression of brain-derived neurotrophic factor." *Neuroscience.* 2012 May; 210: 275-85.

Hu, Y., et al. "Regulation of M1-receptor mRNA stability by smilagenin and its significance in improving memory of aged rats." *Neurobiol. Aging.* 2010 Jun; 31(6): 1010-9.

Zhang, Y., et al. "Role of glial cell derived neurotrophic factor in the protective effect of smilagenin on rat mesencephalic dopaminergic neurons damaged by MPP+." *FEBS Lett.* 2008 Mar; 582(6): 956-60.

Jeon, S., et al. "Beta-secretase (BACE1)-inhibiting stilbenoids from *Smilax* Rhizoma." *Phytomedicine.* 2007 Jun; 14(6): 403-8.

Ban, J., et al. "Catechin and epicatechin from *Smilacis chinae* rhizome protect cultured rat cortical neurons against amyloid beta protein (25-35)-induced neurotoxicity through inhibition of cytosolic calcium elevation." *Life Sci.* 2006 Nov; 79(24) :2251-9.

Ren, L., et al. "Antidepressant-like effects of sarsasapogenin from *Anemarrhena asphodeloides* BUNGE (Liliaceae)." *Biol. Pharm. Bull.* 2006 Nov; 29(11): 2304-6.

Ban, J., et al. "Protection of amyloid beta protein (25-35)-induced neurotoxicity by methanol extract of *Smilacis chinae* rhizome in cultured rat cortical neurons." *J. Ethnopharmacol.* 2006 Jun; 106(2): 230-7.

Barraclough, P., et al. "5-.beta.-sapogenin and pseudosapogenin derivatives and their use in the treatment of dementia." United States Patent 7,138,427: November 21, 2006.

Hu Y, et al. "A new approach to the pharmacological regulation of memory: Sarsasapogenin improves memory by elevating the low muscarinic acetylcholine receptor density in brains of memory-deficit rat models." *Brain Res.* 2005 Oct; 1060(1-2): 26-39.

Xia, Z. et al. Steroidal sapogenins and their derivatives for treating Alzheimer's disease." United States Patent 6,812,213; November 2, 2004.

### **Anti-obesity & Antidiabetic Actions:**

Yang, L., et al. "Dietary supplement of *Smilax china* L. ethanol extract alleviates the lipid accumulation by activating AMPK pathways in high-fat diet fed mice." *Nutr. Metab.* 2019 Jan 21; 16:6.

Pérez-Nájera, V., et al. "*Smilax aristolochiifolia* root extract and its compounds chlorogenic acid and astilbin inhibit the activity of  $\alpha$ -amylase and  $\alpha$ -glucosidase enzymes." *Evid. Based Complement. Alternat. Med.* 2018 Jun; 2018: 6247306.

Turdu, G., et al. "Plant dipeptidyl peptidase-IV inhibitors as antidiabetic agents: a brief review." *Future Med Chem.* 2018 May; 10(10): 1229-1239.

Liu, Y., et al. "Protective effects of sarsasapogenin against early stage of diabetic nephropathy in rats." *Phytother. Res.* 2018 Aug; 32(8): 1574-1582.

Lee H., et al. "Chemical constituents of *Smilax china* L. stems and their inhibitory activities against glycation, aldose reductase,  $\alpha$ -glucosidase, and lipase." *Molecules.* 2017 Mar 11; 22(3).

Kang, Y., et al. "Antiobesity effects of the water-soluble fraction of the ethanol extract of *Smilax china* L. leaf in 3T3-L1 adipocytes." *Nutr. Res. Pract.* 2015 Dec; 9(6): 606-12.

Pereira, F., et al. "Antihyperlipidemic and antihyperglycemic effects of the Brazilian salsaparilhas *Smilax brasiliensis* Spreng. (Smilacaceae) and *Herreria salsaparrilha* Mart. (Agavaceae) in mice treated with a high-refined-carbohydrate containing diet." *Food Res. Int.* 2015 Oct; 76(Pt 3): 366-372.

Sang, H., et al. "The protective effect of *Smilax glabra* extract on advanced glycation end products-induced endothelial dysfunction in HUVECs via RAGE-ERK1/2-NF- $\kappa$ B pathway." *J. Ethnopharmacol.* 2014 Aug; 155(1): 785-95.

### **Anti-Allergy Actions:**

Ki, N., et al. "The hot-water extract of *Smilacis Chinae* Rhizome suppresses 2,4-dinitrochlorobenzene and house dust mite-induced atopic dermatitis-like skin lesions in mice." *Phytother. Res.* 2016 Apr; 30(4): 636-45.

Itharat, A., et al. "Anti-allergic activities of *Smilax glabra* rhizome extracts and its isolated compounds." *J. Med. Assoc Thai.* 2015 Apr; 98 Suppl 3: S66-74.

### **Cytotoxic & Anticancerous Actions:**

Tetty, C., et al. "*Smilax china* leaf extracts suppress pro-inflammatory adhesion response in human umbilical vein endothelial cells and proliferation of HeLa cells." *Arch. Physiol. Biochem.* 2018 Oct 30: 1-5.

She, T., et al. "Sarsaparilla (*Smilax glabra* Rhizome) extract activates redox-dependent ATM/ATR pathway to inhibit cancer cell growth by S phase arrest, apoptosis, and Autophagy." *Nutr. Cancer.* 2017 Nov-Dec; 69(8): 1281-1289.

Fu, S., et al. "Flavonoids and tannins from *Smilax china* L. rhizome induce apoptosis via mitochondrial pathway and MDM2-p53

signaling in human lung adenocarcinoma cells." *Am. J. Chin. Med.* 2017; 45(2): 369-384.

Zhang, C., et al. "Astilbin decreases proliferation and improves differentiation in HaCaT keratinocytes." *Biomed. Pharmacother.* 2017 Sep; 93: 713-720.

Hao, G., et al. "*Smilax glabra* Roxb targets Akt(p-Thr308) and inhibits Akt-mediated signaling pathways in SGC7901 cells." *J. Drug Target.* 2016; 24(6): 557-65.

She, T., et al. "Sarsaparilla (*Smilax glabra* rhizome) extract inhibits migration and invasion of cancer cells by suppressing TGF- $\beta$ 1 pathway." *PLoS One.* 2015 Mar; 10(3): e0118287.

She, T., et al. "Sarsaparilla (*Smilax glabra* rhizome) extract inhibits cancer cell growth by S phase arrest, apoptosis, and autophagy via redox-dependent ERK1/2 pathway." *Cancer Prev. Res.* 2015 May; 8(5): 464-74.

Hu, L., et al. "*Smilax china* L. rhizome extract inhibits nuclear factor- $\kappa$ B and induces apoptosis in ovarian cancer cells." *Chin. J. Integr. Med.* 2015 Dec; 21(12): 907-15.

Nho, K., et al. "Anti-metastatic effect of *Smilax china* L. extract on MDA-MB-231 cells." *Mol. Med. Rep.* 2015 Jan; 11(1): 499-502.

Yu, H., et al. "Extracellular signal regulated kinase inhibition is required for methanol extract of *Smilax china* L. induced apoptosis through death receptor 5 in human oral mucocarcinoma cells." *Mol. Med. Rep.* 2014 Feb; 9(2): 663-8.

Xia, D., et al. "Protective effects of the flavonoid-rich fraction from rhizomes of *Smilax glabra* Roxb. on carbon tetrachloride-induced hepatotoxicity in rats." *J. Membr. Biol.* 2013 Jun; 246(6): 479-85.

Shen, S., et al. "Sarsasapogenin induces apoptosis via the reactive oxygen species-mediated mitochondrial pathway and ER stress pathway in HeLa cells." *Biochem. Biophys. Res. Commun.* 2013 Nov; 441(2): 519-24.

Challinor, V., et al. "Steroidal saponins from the roots of *Smilax* sp.: structure and bioactivity." *Steroids.* 2012 Apr; 77(5):504-11.

Gao, Y., et al. "Mitochondrial apoptosis contributes to the anti-cancer effect of *Smilax glabra* Roxb." *Toxicol Lett.* 2011 Nov; 207(2): 112-20.

Ni, Y., et al. "Mitochondrial ROS burst as an early sign in sarsasapogenin-induced apoptosis in HepG2 cells." *Cell. Biol. Int.* 2008 Mar; 32(3): 337-43.

Trouillas, P., et al. "Structure-function relationship for saponin effects on cell cycle arrest and apoptosis in the human 1547 osteosarcoma cells: a molecular modelling approach of natural molecules structurally close to diosgenin." *Bioorg. Med. Chem.* 2005 Feb; 13(4): 1141-9.

Thabrew, M., et al. "Cytotoxic effects of a decoction of *Nigella sativa*, *Hemidesmus indicus* and *Smilax glabra* on human hepatoma HepG2 cells." *Life Sci.* 2005 Aug; 77(12): 1319-30.

### **Antimicrobial Actions:**

Tian, L., et al. "Steroidal saponins from the genus *Smilax* and their biological activities." *Nat. Prod. Bioprospect.* 2017 Aug; 7(4): 283-298.

Wang, W., et al. "Anti-HIV-1 activities of extracts and phenolics from *Smilax china* L." *Pak. J. Pharm. Sci.* 2014 Jan; 27(1): 147-51.

Xu, S., et al. "Chemical constituents from the rhizomes of *Smilax glabra* and their antimicrobial activity." *Molecules.* 2013 May; 18(5): 5265-87.

Ooi, L., et al. "Antiviral and anti-proliferative glycoproteins from the rhizome of *Smilax glabra* Roxb (Liliaceae)." *Am. J. Chin. Med.* 2008; 36(1): 185-95.

Sautour, M., et al. "Bioactive steroidal saponins from *Smilax medica*." *Planta Med.* 2006 Jun; 72(7): 667-70.

Tewtrakul S, et al. "Anti-HIV-1 protease- and HIV-1 integrase activities of Thai medicinal plants known as Hua-Khao-Yen." *J. Ethnopharmacol.* 2006 Apr; 105(1-2): 312-5.

Sautour, M., et al. "Steroidal saponins from *Smilax medica* and their antifungal activity." *J. Nat. Prod.* 2005 Oct; 68(10): 1489-93.

Ooi, L. S., et al. "New mannose-binding lectin isolated from the rhizome of Sarsaparilla *Smilax glabra* Roxb. (Liliaceae)." *J. Agric. Food Chem.* 2004 Oct; 52(20): 6091-5.

Caceres, A., et al. "Plants used in Guatemala for the treatment of dermatophytic infections. 1. Screening for antimycotic activity of 44 plant extracts." *J. Ethnopharmacol.* 1991; 31(3): 263-76.

Tschesche, R. "Advances in the chemistry of antibiotic substances from higher plants." H. Wagner and L. Horhammer. *Pharmacognosy and Phytochemistry*. New York: Springer Verlag, 1971. 274-76.

Fitzpatrick, F. "Plant substances active against mycobacterium tuberculosis." *Antibiotics and Chemotherapy* 1954; 4(5): 528-36.

#### **Herb-Drug Interactions:**

Li, Y., et al. "*Smilax glabra* rhizoma affects the pharmacokinetics and tissue distribution of methotrexate by increasing the P glycoprotein mRNA expression in rats after oral administration. *Mol. Med. Rep.* 2017 Nov; 16(5): 7633-7640.

**This is an informational site only and no products are sold. The statements contained herein have not been evaluated by the Food and Drug Administration. The information contained in this plant database file is intended for education, entertainment and information purposes only. This information is not intended to be used to diagnose, prescribe or replace proper medical care. The plant described herein is not intended to treat, cure, diagnose, mitigate or prevent any disease. Please refer to our [Conditions of Use](#) for using this plant database file and web site.**