Agrimony

Species (Family)

Agrimonia eupatoria L. (Rosaceae)

Synonym(s)

Agrimonia

Part(s) Used

Herb

Pharmacopoeial and Other Monographs

BHP 1996 (G9)
Complete German Commission E(G3)
PDR for Herbal Medicines 2nd edition(G36)

Legal Category (Licensed Products)

GSL (G37)

Constituents (1,2,G2,G22,G31,G40,G64)

Acids Palmitic acid, salicylic acid, silicic acid and stearic acid.

Flavonoids Apigenin, luteolin, luteolin-7-glucoside, quercetin, quercitrin, kaempferol and glycosides. (3)

Tannins 3-21%. Condensed tannins in herb; hydrolysable tannins (e.g. ellagitannin).

Vitamins Ascorbic acid (vitamin C), nicotinamide complex (about 100–300 μg/g leaf), thiamine (about 2 μg/g leaf) and vitamin K.

Other constituents Bitter principle, triterpenes (e.g. α -amyrin, ursolic acid, euscapic acid), phytosterols and volatile oil 0.2%.

Food Use

Agrimony is listed by the Council of Europe as a natural source of food flavouring (category N2). This category indicates that agrimony can be added to foodstuffs in small quantities, with a possible limitation of an active principle (as yet unspecified) in the final product. (G16)

Herbal Use

Agrimony is stated to possess mild astringent and diuretic properties. (1) It has been used for diarrhoea in children, mucous colitis, grumbling appendicitis, urinary incontinence, cystitis, and as a gargle for acute sore throat and chronic nasopharyngeal catarrh. (G2,G7)

Dosage

Dried herb 2-4 g by infusion three times daily. (G7)

Liquid extract 1-3 mL (1:1 in 25% alcohol) three times daily. (G7)

Tincture 1-4 mL (1:5 in 45% alcohol) three times daily. (G7)

Pharmacological Actions

In vitro and animal studies

Significant uricolytic activity has been documented for agrimony infusions and decoctions (15% w/v), following their oral administration to male rats at a dose of 20 mL/kg body weight (equivalent to 3 g dry drug). (4) Diuretic activity was stated to be minimal and elimination of urea unchanged. A hypotensive effect in anaesthetised cats has been documented for an agrimony extract given by intravenous injection; blood pressure was lowered by more than 40%. (5)

Marked antibacterial activity against *Staphylococcus aureus* and α-haemolytic streptococci has been reported for agrimony. (6)

An aqueous ethanol extract of the herb was tested for immunomodulative activity in the peritoneal cavities of mice. (7) Immunostimulant activity resulted in an increase in phagocytic activity and increases in the activities of lysozyme and peroxidase. Agrimonia eupatoria given in the diet of mice for 12 days prior to intraperitoneal administration of streptozotocin resulted in a reduction in hyperglycaemia. (8) Further investigation revealed stimulation of 2-deoxyglucose transport, glucose oxidation and incorporation of glucose into glycogen in mouse abdominal muscle. An aqueous extract (0.25–1 mg/ mL) stimulated insulin secretion from a BRIN-BD11 pancreatic B cell line. (9) These findings demonstrate that A. eupatoria aqueous extract given orally to mice

has antihyperglycaemic, insulin-releasing and insulin-like activity. (9)

A related species, A. pilosa, has also been investigated. In vivo antitumour activity in mice has been attributed to the tannin agrimoniin (10) which has not been reported as a constituent of A. eupatoria. Agrimoniin was administered intraperitoneally into ascites-type and solid tumours in rodents. (11) At doses of greater than 10 mg/kg, given before or after intraperitoneal inoculation with MM2 cells, it completely rejected tumour growth in mice. (11) Solid tumours of MH134 and Meth-A were inhibited by agrimoniin, and the number of peripheral blood cells was increased, indicating that agrimoniin has antitumour activity and that it exerts its effect by enhancing the immune response. In vitro studies have reported that agrimoniin induces the cytotoxicity of murine peritoneal exudate cells, (12) and that it induces interleukin 1 in human peripheral blood mononuclear cells and in mouse adherent peritoneal exudate cells in vivo. (13) Several phloroglucinols isolated from A. pilosa have demonstrated activity against Staphylococcus aureus, (14) and a methanol extract of the herb inhibited HIV-1 protease activity. (15) An aqueous suspension of A. pilosa herb (1 mg/kg and 5 mg/kg) given orally or intraperitoneally significantly reduced blood glucose concentrations in streptozotocin-induced diabetic rats. (16)

Clinical studies

The successful treatment of cutaneous porphyria in a group of 20 patients receiving agrimony infusions has been described. (17) An improvement in skin eruptions together with a decrease in serum iron concentrations and in urinary porphyrins was noted.

A compound herb preparation containing agrimony has been used to treat 35 patients suffering from chronic gastroduodenitis. After 25 days of therapy, 75% of patients claimed to be free from pain, 95% from dyspeptic symptoms and 76% from palpitation pains. Gastroscopy was said to indicate that previous erosion and haemorrhagic mucous changes had healed. No side-effects or signs of toxicity were documented.

Side-effects, Toxicity

None documented for A. eupatoria. A polar fraction containing flavonoids and triterpenes, but not tannins, produced a negative result in the Ames test. (1)

In mice, agrimoniin has been documented to cause stretching and writhing reactions when administered by intraperitoneal injection, and cyanosis and necrosis at the site of intravenous injection. These reactions were considered to be inflammatory reac-

tions. The LD₅₀ of agrimoniin in mice has been estimated as 33 mg/kg (by intravenous injection), 101 mg/kg (by intraperitoneal injection), and greater than 1 g/kg (by mouth).⁽¹¹⁾ Cytotoxic activity has been reported for A. pilosa⁽¹⁰⁾ (see In vitro and animal studies).

Contra-indications, Warnings

Excessive doses may interfere with existing drug treatment for high or low blood pressure, and anticoagulant therapy. In view of the tannin constituents, excessive use should be avoided.

Pregnancy and lactation Agrimony is reputed to affect the menstrual cycle. (G22) In view of the lack of toxicity data, excessive use of agrimony should be avoided during pregnancy and lactation.

Pharmaceutical Comment

Relatively limited information is available on the chemistry of agrimony, although more is known about the tannin constituents of the related species A. pilosa. Human studies have indicated that agrimony may be useful in the treatment of certain cutaneous and gastrointestinal disorders, although further studies are needed to confirm these reports. The tannin constituents may justify the astringent activity attributed to the herb. In view of the lack of toxicity data, excessive use of agrimony should be avoided.

References

See also General References G2, G3, G9, G16, G22, G31, G36, G37, G40, G42 and G64.

- 1 Bilia AR et al. Constituents and biological assay of Agrimonia eupatoria. Fitoterapia 1993; 64: 549-550.
- 2 Carnat A et al. L'aigremoine: étude comparée d'Agrimonia eupatoria L. et Agrimonia procera Wallr. Plantes médicinales et phytothérapie 1991; 25: 202-211.
- 3 Sendra J, Zieba J. Flavonoids from Agrimonia eupatoria L. Diss Pharm Pharmacol 1971; 24: 79-83.
- 4 Giachetti D et al. Ricerche sull'attivita diuretica ed uricosurica di Agrimonia eupatoria. Boll Soc Ital Biol Sper 1986; 62: 705-711.
- 5 Petkov V. Plants with hypotensive, antiatheromatous and coronarodilatating action. Am J Chin Med 1979; 7: 197-236.
- 6 Petkov V. Bulgarian traditional medicine: A source of ideas for phytopharmacological investigations. *J Ethnopharmacol* 1986; 15: 121-132.
- 7 Bukovsky M, Blanárik P. Immunomodulative

- 8 Swanston-Flatt SK et al. Traditional plant treatments for diabetes in normal and streptozotocin diabetic rats. Diabetologia 1990; 33: 462-464.
- 9 Gray AM, Flatt PR. Actions of the traditional antidiabetic plant, Agrimonia eupatoria (agrimony): effects on hyperglycaemia, cellular glucose metabolism and insulin secretion. Br J Nutr 1998; 80: 109-114.
- 10 Miyamoto K et al. Isolation of agrimoniin, an antitumour constituent, from the roots of Agrimonia pilosa Ledeb. Chem Pharm Bull (Tokyo) 1985; 33: 3977-3981.
- 11 Miyamoto K et al. Antitumour effect of agrimoniin, a tannin of Agrimonia pilosa Ledeb., on transplantable rodent tumors. Jpn J Pharmacol 1987; 43: 187–195.
- 12 Miyamoto K et al. Induction of cytotoxicity of peritoneal exudate cells by agrimoniin, a novel immunomodulatory tannin of Agrimonia pilosa Ledeb. Cancer Immunol Immunother 1988; 27:

- 59-62
- 13 Murayama T et al. Agrimoniin, an antitumour tannin of Agrimonia pilosa Ledeb., induces interleukin-1. Anticancer Res 1992; 12: 1471–1474.
- 14 Yamaki M et al. Antimicrobial activity of naturally occurring and synthetic phloroglucinols against Staphylococcus aureus. Phytother Res 1994; 8: 112-114.
- 15 Min BS et al. Screening of Korean plants against human immunodeficiency virus type 1 protease. Phytother Res 1999; 13: 680-682.
- 16 Hsu F-L, Cheng J-T. Investigation in rats of the antihyperglycaemic effect of plant extracts used in Taiwan for the treatment of diabetes mellitus. Phytother Res 1992; 6: 108-111.
- 17 Patrascu V et al. Rezultate terapeutice favorabile in porfiria cutanata cu Agrimonia eupatoria. Dermato-venerologia 1984; 29: 153-157.
- Chakarski I et al. Clinical study of a herb combination consisting of Agrimonia eupatoria, Hipericum perforatum, Plantago major, Mentha piperita, Matricaria chamomila for the treatment of patients with chronic gastroduodenitis. Probl Vatr Med 1982; 10: 78-84.