



## Pharmacological Potential of *Matricaria recutita*-A Review

Vikas Gupta<sup>1\*</sup>, Payal Mittal<sup>2</sup>, Parveen Bansal<sup>1</sup>, Sukhbir L Khokra<sup>3</sup>, Dhirender Kaushik<sup>3</sup>

<sup>1</sup>National Institute of Ayurvedic Pharmaceutical Research, CCRAS, Patiala, Punjab, India

<sup>2</sup>Akal College of Pharmacy and Technical Education, Mastuana Sahib, Sangrur, India

<sup>3</sup>Institute of Pharmaceutical Sciences, Kurukshetra University, Kurukshetra, India

### ABSTRACT

*Matricaria recutita*, once it has been called as *Marticaria chamomilla*, *Chamomilla recutita*, and *Chamomilum nobile* family *Asteraceae*. The main constituents of this plant include the terpenoids  $\alpha$ -bisabolol and its oxides and azulenes, including chamazulene. *Matricaria recutita* shows different pharmacological activities like anti-inflammatory, anti-cancer, treatment of stress and depression, anti-allergic etc. This review focuses on the detailed chemical constituents, pharmacological activities of different parts of this plant.

**Keywords:** Marticaria chamomilla, terpenoids, chamazulene.

### INTRODUCTION

*Matricaria recutita*, once it has been called as Marticaria chamomilla, Chamomilla recutita, and Chamomilum nobile family *Asteraceae* and commonly it is known as German chamomile, Roman chamomile, English chamomile, Camomilla, and Flos Chamomile. [1-2]

It mainly grows indigenously in Europe, NW. Asia, N. Africa, and cultivated in N. America and in many parts of the world. [1, 3] This herb has been used as herbal remedies for thousands of years. This herb has been believed by Anglo-Saxons as one of nine sacred herbs given to humans by the lord. [4] One of the most commonly consumed single ingredient herbal tea is chamomile, prepared with dried flowers from *Matricaria recutita* L. The composite flower is white in color with a yellowish orange center. [1] Infusions and essential oils from fresh or dried flower heads have aromatic, flavoring and coloring properties. Both are used in a number of commercial products including soaps, detergents, perfumes, lotions, ointments, hair products, baked goods, confections, alcoholic beverages and herbal teas. Chamomile flowers contain 0.24- to 2.0 percent volatile oil that is blue in color. [1, 3] European Pharmacopoeia recommends chamomile contains no less than 4 mL/kg of blue essential oil. [5]

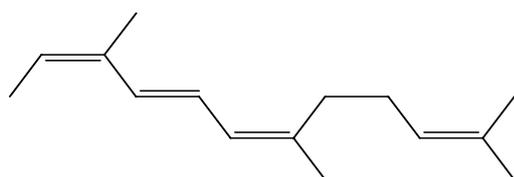
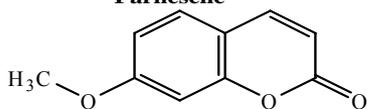
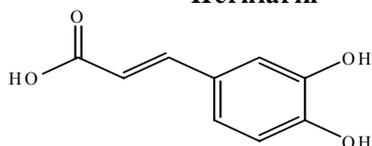
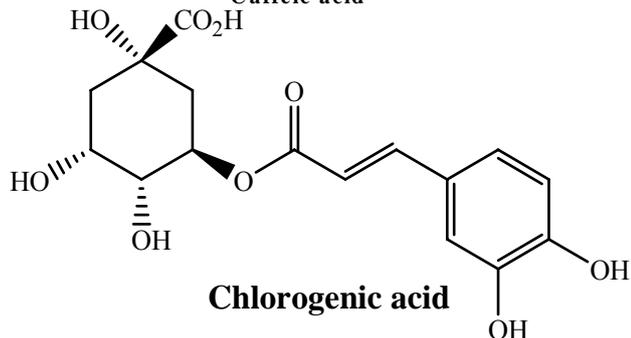
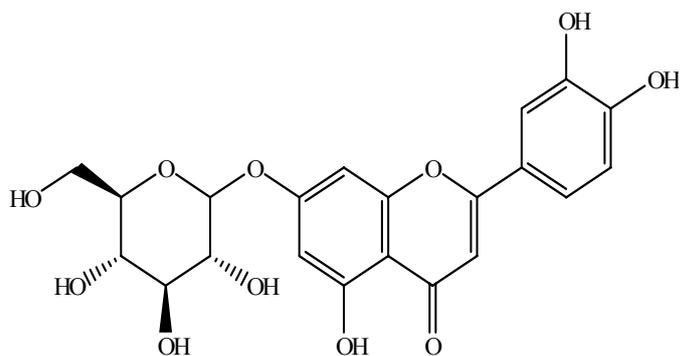
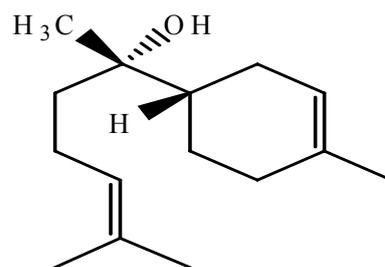
### CHEMICAL CONSTITUENTS

**\*Corresponding author: Mr. Vikas Gupta,**  
National Institute of Ayurvedic Pharmaceutical Research,  
CCRAS, Patiala, Punjab, India; **Tel:** 09914933022  
**E-mail:** vikas\_4308@rediffmail.com

Over 120 constituents have been identified in chamomile flowers. [6] The main constituents of the oil include the terpenoids  $\alpha$ -bisabolol and its oxides ( $\leq 78\%$ ) and azulenes, including chamazulene (1-15%). [7-11]

Chamazulene carboxylic acid and proazulenes occur in chamomile. [12] Precocenes were isolated from the essential oils of *Matricaria recutita*. [13] The essential oil of German chamomile showed specific inhibition toward aflatoxin G (1) (AFG (1)) production, and (E) - and (Z)-spiroethers were isolated as the active compounds from the oil. [14] Farnesene (12-28%), spathulenol and spiroethers, including the *cis/trans*-en-yn-dicycloethers (8-20%), are also present in the volatile oil. [15-17] Eleven bioactive phenolic compounds (coumarins: herniarin, umbelliferone; phenylpropanoids: chlorogenic acid, caffeic acid; flavones: apigenin, apigenin-7-O-glucoside, luteolin, luteolin-7-O-glucoside; flavonols: quercetin, rutin and flavanone: naringenin) are found in chamomile extract. The main constituents of the flowers include several phenolic compounds, primarily the flavonoids apigenin, quercetin, patuletin, luteolin and their glucosides Coumarins and Dicycloethers also occur in the flowers. The principal components of the essential oil extracted from the flowers are the terpenoids  $\alpha$ -bisabolol and its oxides and azulenes, including chamazulene. Chamomile (*Matricaria chamomilla*) in the above-ground organs synthesizes and accumulates (Z) - and (E)-2-beta-D: -glucopyranosyloxy-4-methoxy cinnamic acids (GMCA), the precursors of phytoanticipin herniarin (7-methoxycoumarin). [18-26] The largest group of medically important compounds forming the essential oils are primarily chamazulene, (-)- $\alpha$ -bisabolol, bisabololoxides, bisabolonoxide A, trans-

beta-farnesene, alpha-farnesene, spathulenol and the cis/trans-en-in-dicycloethers. Flavonoids, coumarins, mucilages, mono- and oligosaccharides also have pharmacological effects. [27-28] Fractionation of the aqueous extract of this plant led to the detection of several fractions with significant affinity for the central benzodiazepine receptor and to the isolation and identification of 5, 7, 4'-trihydroxyflavone (apigenin) in one of them. [29-30] The essential oil of German chamomile showed specific inhibition toward aflatoxin G (1) (AFG (1)) production, and (E) - and (Z)-spiroethers were isolated as the active compounds from the oil. [14] The major flavonoids in the white florets of chamomile (*Chamomilla recutita* [L.] Rauschert) were rapidly purified using a combination of polyamide solid-phase extraction and preparative HPLC. From the combined LC/MS, LC/MS/MS, and NMR data the apigenin glucosides were identified as apigenin 7-O-glucoside (Ap-7-Glc), Ap-7-(6"-malonyl-Glc), Ap-7-(6"-acetyl-Glc), Ap-7-(6"-caffeoyl-Glc), Ap-7-(4"-acetyl-Glc), Ap-7-(4"-acetyl,6"-malonyl-Glc), and a partially characterized apigenin-7-(mono-acetyl/mono-malonylglucoside) isomer. [31] Precocenes were isolated from the essential oils of *Matricaria recutita*. [13] We screened various isoprenoids to search for inducers of apoptosis-like cell death ("apoptosis") in the shoot primordia of *Matricaria chamomilla* and found that geraniol has the most potent apoptosis-inducing activity among terpenoids. [32] Chamazulene carboxylic acid is a natural profen with anti-inflammatory activity and a degradation product of proazulenic sesquiterpene lactones, e.g., matricin. Both 1 and proazulenes occur in chamomile (*Matricaria recutita*). [33] A Flavone 7-O-Glucoside-Specific Glucosidase from Ligulate Florets of *Chamomilla recutita* is also found. [34]

**Farnesene****Herniarin****Caffeic acid****Chlorogenic acid****Luteolin-7-O-glucoside****alpha-Bisabolol**

## GENERAL PHARMACOLOGICAL ACTIVITIES

### Anti-inflammatory

The freeze-dried extracts of chamomile (*Matricaria chamomilla* L.) and was found to suppress both the inflammatory effect and the leukocyte infiltration. *Matricaria chamomilla* was assessed for its anti-inflammatory activity on intact rats by measuring the suppression of carrageenan-induced paw edema produced by 1/10 of the intraperitoneal LD<sub>50</sub> dose for the 80 % ethanol extract. Results showed that the plant possessed good anti-inflammatory activity. [35-36]

### Immunomodulatory activity

Intragastric and parenteral administration of heteropolysaccharides of *Matricaria chamomilla* L. is found to normalize developing of the immune response upon air cooling and enhance (but do not normalize) this process upon immersion cooling. The immunomodulating effect of the heteropolysaccharides upon cooling is attributed to initiation of immunostimulating properties of heavy erythrocytes (macrocytes), activation of immunoregulation cells of peripheral blood, and increased sensitivity of effector cells to helper signals. [37]

### Acaricidal property

Acaricidal properties of decoctions, infusions and macerates of dried flower heads of chamomile, *Matricaria chamomilla* L. were tested in vitro against the mite *Psoroptes cuniculi* Delafond (Parasitiformes: Psoroptidae). This mite species is responsible for otoacariasis in domestic animals. Mites were exposed to the extracts for 24, 48 or 72 h. All the extracts tested showed highly significant acaricidal activity when compared with controls. Among them, a decoction of 10% was the only formulation which gave 100% activity at all the three observations times. [38]

### Antihyperglycemic

*Matricaria chamomilla* L. ethanolic extract treatment protected the majority of the pancreatic islet cells, with respect to the control group. As a result, *Matricaria*

*chamomilla* L. ethanolic extract exhibited significant antihyperglycemic effect and protected beta-cells in STZ-diabetic rats, in a dose-dependent manner, and diminished the hyperglycemia-related oxidative stress. [39]

#### Anti-cancer activity

The aqueous and methanolic extracts of chamomile showed differential apoptosis in cancer cells but not in normal cells at similar doses. [40]

#### Antipruritic effect

The single per oral administration of the ethyl acetate extract or essential oil of German chamomile (*Matricaria recutita* L.) showed remarkable antipruritic effects in the compound 48/80-induced itch-scratching test in ddY mice. [41]

#### Wound healing property

The aqueous extract of *M. recutita* (120 mg/kg/day) showed increased rate of wound contraction, together with the increased wound-breaking strength, hydroxyproline content. The chamomile extract in the form of rubbing oil had a good potential for acceleration of burn wound healing in rats. The extract of *M. chamomilla* administered topically has wound healing potential in linear incisional wound model in rats. Animals treated with chamomile presented significantly faster wound healing in comparison to those treated with corticosteroids. [42-45]

#### Treatment of oral mucositis

Methotrexate-induced oral mucositis in a patient with rheumatoid arthritis was successfully treated with Wild chamomile mouthwashes. [46]

#### Intracanal irrigant

Chamomile or tea tree oil was effective in removal of the smear layer. [47]

#### Treatment of infant botulism

Chamomile (principally, unwrapped chamomile) is a potential vehicle of *C. botulinum* spores, and ingestion of chamomile tea could represent a risk for infant botulism. [48]

#### Lousicidal, ovicidal and repellent property

*Matricaria chamomilla* essential oil has lousicidal, ovicidal and repellent efficacy against lice and flies infesting water buffaloes. [49]

#### Virucidal agent

Camomile oil exhibited a high selectivity index and seems to be a promising candidate for topical therapeutic application as virucidal agents for treatment of herpes genitalis. [50]

#### Treatment of gastrointestinal disorders

Methanol extracts of *Matricaria recutita* (flowers) and *Ginkgo biloba* (leaves) had a MIC > 100 microg/mL against the gram-negative bacterium *Helicobacter pylori* (HP). [51]

#### Antimicrobial activity

The essential oil from and *M. chamomila* were active against 3 strains of *S. aureus* and the *Candida* strains and can be used in the treatment of acute otitis externa. [52]

#### Antiulcer activity

Extracts from the plants *Iberis amara*, *Melissa officinalis*, *Matricaria recutita*, *Carum carvi*, *Mentha x piperita*, *Glycyrrhiza glabra*, *Angelica archangelica*, *Silybum marianum* and *Chelidonium majus*, singly and combined in the form of a commercial preparation, STW 5 (Iberogast). All extracts produced a dose dependent anti-ulcerogenic activity associated with a reduced acid output and an increased mucin secretion, an increase in prostaglandin E2 release and a decrease in leukotrienes. [53]

#### Treatment of stress and depression

Chamomilla 6cH is related to the recovery of basal behavioral conditions in mice subjected to stressful conditions. [54]

#### Uterotonic

Water extracts (infusions) from a group of medicinal plants were studied in terms of their activity enhancing the uterine tonus in a series of experiments with a preparation of an isolated rabbit and guinea pig uterine horn. [55]

#### Anti-allergic activity

The inhibitory effects of the dietary intake of the German chamomile extracts on compound 48/80-induced itch-scratch response were comparable to oxatamide (10 mg/kg, p.o.), an anti-allergic agent. [56]

#### Antisolar agent

Liquid and dry extracts of *Hamamelis virginiana*, *Matricaria recutita*, *Aesculus hippocastanum*, *Rhamnus purshiana* and *Cinnamomum zeylanicum* were prepared by reprecipitation, maceration and microwave oven extraction. The solar protection factors (SPF) of these preparations were determined by a spectrophotometric method. The results showed that after incorporation to a 2 % solution of the synthetic sunscreen octylmethoxycinnamate, the extracts showed intensification in SPF values, suggesting that this can be an interesting method to intensify SPF. [57]

#### Inhibition of poliovirus replication

Hydroalcoholic extract of *Matricaria chamomilla* added during the early stage of Poliovirus development inhibits cellular and viral RNA synthesis. [58]

#### Anxiolytic agent

A significant reduction in mean total Hamilton Anxiety Rating (HAM-A) scores was observed during chamomile versus placebo therapy (P = 0.047). [59]

#### Prevent osteoporosis

The aqueous extracts derived from *Matricaria chamomilla* may form the basis to design "functional foods" for the prevention of osteoporosis. [60]

*M. chamomila* is a popular medicinal plant useful in various ailments. A number of old ayurvedic texts have mentioned tremendous and a variety of uses of *Matricaria recutita*. Today evidence based studies are needed to establish these facts so that these wonder drugs with multifarious therapeutic activities can be put to human use.

## REFERENCES

1. WHO Monograph on selected Medicinal plants: <http://www.who.int/medicines/library/trm/medicinalplants/monographs.shtml>.
2. Micromedex Healthcare Series: MICROMEDEX, Inc., Englewood, Colorado (Vol. 115 expires 3/2003)
3. Wald G., Brendler T. PDR for Herbal Medicines. 1<sup>st</sup> ed. Montville, (NJ) Medical Economics Company publishers; 1998. 07645-1742.
4. Crevin JK, philpott J. Herbal Medicine Past and Present. Vol. II Duke University Press; 1990.
5. *European Pharmacopoeia*. 5th ed. Strasbourg, France: European Directorate for the Quality of Medicines of the Council of Europe; 1996:1976-1977.
6. Pino JA, Bayat F, Marbot R, Aguero J. Essential oil of chamomile *Chamomilla recutita* (L.) Rausch. from Iran. *J Essent Oil Res* 2002; 14: 407-408.
7. Pino JA, Marbot R, Aguiro J, Fuentes V. Essential oil of chamomile *Chamomilla recutita* (L.) Rausch. from Cuba. *J Essent Oil-Bear Plants* 2000; 3: 1-3.
8. Mann C, Staba EJ. The chemistry, pharmacology, and commercial formulations of Chamomile. In *Herbs, Spices, and Medicinal Plants: Recent Advances in Botany, Horticulture, and*

- Pharmacology Volume 1, Craker LE, Simon JE (eds). Oryx Press: Phoenix, 1986; 235–280.
9. Matos FJA, Machado MIL, Alencar JW, Craveiro AA. Constituents of Brazilian chamomile oil. *J Essent Oil Res* 1993; 5: 337–339.
  10. Mimica-Dukic N, Lukic V, Pavkov R, Gasic O. Study of chemical composition and microbiological contamination of chamomile tea. *Acta Hort* 1993; 333: 137–141.
  11. Stanev S, Zheljazkov V, Janculoff Y. Variation of chemical compounds in the essential oil from some native forms of chamomile (*Chamomilla recutita* L.). *Beitr Zuchtungsforshung* 1996; 2: 214–217.
  12. Ramadan M, Goeters S, Watzler B, Krause E, Lohmann K, Bauer R, Hempel B, Imming P. Chamazulene carboxylic acid and matricin: a natural profen and its natural prodrug, identified through similarity to synthetic drug substances. *J Nat Prod*. 2006; 69(7): 1041-5.
  13. Yaguchi A, Yoshinari T, Tsuyuki R, Takahashi H, Nakajima T, Sugita-Konishi Y, Nagasawa H, Sakuda S. Isolation and identification of precocenes and piperitone from essential oils as specific inhibitors of trichothecene production by *Fusarium graminearum*. *J Agric Food Chem*. 2009; 57(3):846-51.
  14. Yoshinari T, Yaguchi A, Takahashi-Ando N, Kimura M, Takahashi H, Nakajima T, Sugita-Konishi Y, Nagasawa H, Sakuda S. Spiroethers of German chamomile inhibit production of aflatoxin G and trichothecene mycotoxin by inhibiting cytochrome P450 monooxygenases involved in their biosynthesis. *FEMS Microbiol Lett*. 2008; 284(2):184-90.
  15. Yaguchi A, Yoshinari T, Tsuyuki R, Takahashi H, Nakajima T, Sugita-Konishi Y, Nagasawa H, Sakuda S. Isolation and identification of precocenes and piperitone from essential oils as specific inhibitors of trichothecene production by *Fusarium graminearum*. *J Agric Food Chem*. 2009; 57(3): 846-51.
  16. Yoshinari T, Yaguchi A, Takahashi-Ando N, Kimura M, Takahashi H, Nakajima T, Sugita-Konishi Y, Nagasawa H, Sakuda S. Spiroethers of German chamomile inhibit production of aflatoxin G and trichothecene mycotoxin by inhibiting cytochrome P450 monooxygenases involved in their biosynthesis. *FEMS Microbio Lett*. 2008; 284(2):184-90.
  17. Lis-Balchin M, Deans SG, Eaglesham E. Relationship between bioactivity and chemical composition of commercial essential oils. *Flavour Fragrance J* 1998; 13: 98–104.
  18. McKay DL, Blumberg JB. A review of the bioactivity and potential health benefits of chamomile tea (*Matricaria recutita* L.). *Phytother Res*. 2006; 20(7):519-30.
  19. Evdokimoff V, Tacci Bucci B, Cavazzutti G. [Analytical study of chemazulene from essential oil of *Matricaria chamomilla* L]. *Farmacoprat* 1972; 27(3):163-73.
  20. Schenck G, KH. [Determination of oil and azulene content in *Matricaria chamomilla* in different stages of its blossom.]. *Arch Pharm Ber Dtsch Pharm Ges* 1956; 289(4): 200-03.
  21. Redaelli C, Formentini L, Santaniello E. Evaluation of Coumarins and Dicycloethers in *Matricaria chamomilla* Flowers and Chamomile Extracts by Reverse-Phase HPLC. *Planta Med*. 1981; 42(6):130.
  22. Padula LZ, Rondina RV, Coussio JD. Quantitative determination of essential oil, total azulenes and chamazulene in German chamomile (*Matricaria chamomilla*) cultivated in Argentina. *Planta Med*. 1976; 30(3):273-80.
  23. Pietta P, Manera E, Ceva P. Simultaneous isocratic high-performance liquid chromatographic determination of flavones and coumarins in *Matricaria chamomilla* extracts. *J Chromatogr*. 1987; 404(1):279-81.
  24. Fonseca FN, Tavares MF, Horvath C. Capillary electrochromatography of selected phenolic compounds of *Chamomilla recutita*. *J Chromatogr A* 2007; 1154(1-2):390-9.
  25. Repčák M, Smajda B, Kovacik J, Eliasova A. Circadian rhythm of Z- and E-2-beta-D: -glucopyranosyloxy-4-methoxy cinnamic acids and herniarin in leaves of *Matricaria chamomilla*. *Plant Cell Rep*. 2009; 28(7):1137-43.
  26. Redaelli C, Formentini L, Santaniello E. HPLC Determination of Coumarins in *Matricaria chamomilla*. *Planta Med*. 1981; 43(12):412-3.
  27. Máday E, Szöke E, Muskáth Z, Lemberkovics E. A study of the production of essential oils in chamomile hairy root cultures. *Eur J Drug Metab Pharmacokinet*. 1999; 24(4):303-8.
  28. Schilcher H, Novotny L, Ubik K, Motl O, Herout V. Structure of a third bisabololoxide from *Matricaria chamomilla* L (AUTHOR'S TRANSL) *Arch Pharm (Weinheim)*. 1976; 309(3):189-96.
  29. Viola H, Wasowski C, Levi de Stein M, Wolfman C, Silveira R, Dajas F, Medina JH, Paladini AC. Apigenin, a component of *Matricaria recutita* flowers, is a central benzodiazepine receptors-ligand with anxiolytic effects. *Planta Med*. 1995; 61(3):213-6.
  30. Zanolli P, Avallone R, Baraldi M. Behavioral characterisation of the flavonoids apigenin and chrysin. *Fitoterapia* 2000; (71 Suppl 1): S117-23
  31. Svehlikova V, Bennett RN, Mellon FA, Needs PW, Piacente S, Kroon PA, Bao Y. Isolation, identification and stability of acylated derivatives of apigenin 7-O-glucoside from chamomile (*Chamomilla recutita* [L.] Rauschert). *Phytochemistry* 2004; 65(16):2323-32.
  32. Izumi S, Takashima O, Hirata T. Geraniol is a potent inducer of apoptosis-like cell death in the cultured shoot primordia of *Matricaria chamomilla* *Biochem Biophys Res Commun*. 1999; 259(3):519-22.
  33. Ramadan M, Goeters S, Watzler B, Krause E, Lohmann K, Bauer R, Hempel B, Imming P. Chamazulene carboxylic acid and matricin: a natural profen and its natural prodrug, identified through similarity to synthetic drug substances. *J Nat Prod*. 2006; 69(7):1041-5.
  34. Maier R, Carle R, Kreis W, Reinhard E. Purification and Characterization of a Flavone 7-O-Glucoside-Specific Glucosidase from Ligulate Florets of *Chamomilla recutita*. *Planta Med*. 1993; 59(5):436-41.
  35. Shipochliev T, Dimitrov A, Aleksandrova E. Anti-inflammatory action of a group of plant extracts] *Vet Med Nauki*. 1981;18(6):87-94.
  36. Al-Hindawi MK, Al-Deen IH, Nabi MH, Ismail MA. Anti-inflammatory activity of some Iraqi plants using intact rats. *J Ethnopharmacol* 1989; 26(2):163-8.
  37. Uteshev BS, Laskova IL, Afanasev VA. [The immunomodulating activity of the heteropolysaccharides from German chamomile (*Matricaria chamomilla*) during air and immersion cooling Eksp Klin Farmakol 1999; 62(6):52-5.
  38. Macchioni F, Perrucci S, Cecchi F, Cioni PL, Morelli I, Pampiglione S. Acaricidal activity of aqueous extracts of chamomile flowers, *Matricaria chamomilla*, against the mite *Psoroptes cuniculi* *Med Vet Entomol* 2004; 18(2):205-7.
  39. Cemek M, Kağa S, Simşek N, Büyükkuroğlu ME, Konuk M. Antihyperglycemic and antioxidant potential of *Matricaria chamomilla* L. in streptozotocin-induced diabetic rats. *Nat Med (Tokyo)*. 2008; 62(3):284-93.
  40. Srivastava JK, Gupta S. Antiproliferative and apoptotic effects of chamomile extract in various human cancer cells. *J Agric Food Chem*. 2007; 55(23):9470-8.
  41. Kobayashi Y, Takahashi R, Ogino F. Antipruritic effect of the single oral administration of German chamomile flower extract and its combined effect with antiallergic agents in ddY mice. *J Ethnopharmacol* 2005; 101(1-3):308-12.
  42. Nayak BS, Raju SS, Rao AV. Wound healing activity of *Matricaria recutita* L. extract. *J Wound Care*. 2007; 16(7):298-302.
  43. Jarrahi M. An experimental study of the effects of *Matricaria chamomilla* extract on cutaneous burn wound healing in albino rats. *Nat Prod Res*. 2008; 22(5):422-7.
  44. Jarrahi M, Vafaei AA, Taherian AA, Miladi H, Rashidi Pour A. Evaluation of topical *Matricaria chamomilla* extracts activity on linear incisional wound healing in albino rats. *Nat Prod Res*. 2008; 22(14):1197-202.
  45. Martins MD, Marques MM, Bussadori SK, Martins MA, Pavesi VC, Mesquita-Ferrari RA, Fernandes KP. Comparative analysis between *Chamomilla recutita* and corticosteroids on wound healing. An in vitro and in vivo study. *Phytother Res*. 2009; 23(2):274-8.
  46. Mazokopakis EE, Vrentzos GE, Papadakis JA, Babalis DE, Ganotakis ES. Wild chamomile (*Matricaria recutita* L.) mouthwashes in methotrexate-induced oral mucositis. *Phytomedicine* 2005; 12(1-2):25-7.
  47. Sadr Lahijani MS, Raof Kateb HR, Heady R, Yazdani D. The effect of German chamomile (*Matricaria recutita* L.) extract and tea tree (*Melaleuca alternifolia* L.) oil used as irrigants on removal of smear layer: a scanning electron microscopy study. *Int Endod J* 2006; 39(3):190-5.
  48. Bianco MI, Lúquez C, de Jong LI, Fernández RA. Presence of *Clostridium botulinum* spores in *Matricaria chamomilla* (chamomile) and its relationship with infant botulism. *Int J Food Microbio* 2008; 121(3):357-60.
  49. Khater HF, Ramadan MY, El-Madawy RS. Lousicidal, ovicidal and repellent efficacy of some essential oils against lice and flies infesting water buffaloes in Egypt. *Vet Parasitol* 2009.
  50. Koch C, Reichling J, Schneele J, Schnitzler P. Inhibitory effect of essential oils against herpes simplex virus type 2. *Phytomedicine* 2008; 15(1-2):71-8.

51. Mahady GB, Pendland SL, Stoa A, Hamill FA, Fabricant D, Dietz BM, Chadwick LR. In vitro susceptibility of *Helicobacter pylori* to botanical extracts used traditionally for the treatment of gastrointestinal disorders. *Phytother Res.* 2005; 19(11):988-91.
52. Nogueira JC, Diniz Mde F, Lima EO. In vitro antimicrobial activity of plants in Acute Otitis Externa. *Braz J Otorhinolaryngol* 2008; 74(1):118-24.
53. Khayyal MT, el-Ghazaly MA, Kenawy SA, Seif-el-Nasr M, Mahran LG, Kafafi YA, Okpanyi SN. Antiulcerogenic effect of some gastrointestinally acting plant extracts and their combination. *Arzneimittelforschung* 2001; 51(7):545-53.
54. Pinto SA, Bohland E, Coelho Cde P, Morgulis MS, Bonamin LV. An animal model for the study of Chamomilla in stress and depression: pilot study. *Homeopathy.* 2008; 97(3): 141-4
55. Shipochliev T. Uterotonic action of extracts from a group of medicinal plants]. *Vet Med Nauki* 1981; 18(4): 94-8.
56. Kobayashi Y, Nakano Y, Inayama K, Sakai A, Kamiya T. Dietary intake of the flower extracts of German chamomile (*Matricaria recutita* L.) inhibited compound 48/80-induced itch-scratch responses in mice. *Phytomedicine* 2003; 10(8):657-64.
57. Ramos MF, Santos EP, Bizarri CH, Mattos HA, Padilha MR, Duarte HM. Preliminary studies towards utilization of various plant extracts as antisolar agents. *Int J Cosmet Sci* 1996; 18(3):87-101.
58. Vilaginès P, Delaveau P, Vilagines R. Inhibition of poliovirus replication by an extract of *Matricaria chamomilla* (L) *C R Acad Sci III.* 1985; 301(6):289-94.
59. Amsterdam JD, Li Y, Soeller I, Rockwell K, Mao JJ, Shults J. A randomized, double-blind, placebo-controlled trial of oral *Matricaria recutita* (chamomile) extract therapy for generalized anxiety disorder. *J Clin Psychopharmacol* 2009; 29(4):378-82.
60. Kassi E, Papoutsi Z, Fokialakis N, Messari I, Mitakou S, Moutsatsou P. Greek plant extracts exhibit selective estrogen receptor modulator (SERM)-like properties. *J Agric Food Chem.* 2004; 52(23):6956-61.