

Wound Healing Property Review of Siam Weed, *Chromolaena odorata*

Anushika Sirinthipaporn, Wannee Jiraungkoorskul¹

Mahidol University International College, Mahidol University, Salaya Campus, Nakhon Pathom, ¹Department of Pathobiology, Faculty of Science, Mahidol University, Bangkok, Thailand

ABSTRACT

Chromolaena odorata (Family: *Asteraceae*) synonyms as *Eupatorium odoratum* is a traditional medicinal plant that is widely used for its wound healing property. In particular, the several parts of this herb have been used to treat wounds, burns, and skin infections. Furthermore, it has also been shown to possess anticancer, antidiabetic, anti-hepatotoxic, anti-inflammatory, antimicrobial, and antioxidant properties. Its phytochemical components are alkaloids, flavonoids, flavanone, essential oils, phenolics, saponins, tannins, and terpenoids. The other important constituents of this plant are Eupolin, chromomoric acid, quercetagenin, and quercetin, all of which contribute to its remedial properties. Published information on the wound healing property of *C. odorata* was gathered by the use of different scientific websites such as Google Scholar, Science Direct, PubMed, and Web of Knowledge to provide an up-to-date review showing its importance.

Key words: Antioxidant, *Chromolaena odorata*, healing property, plant, traditional medicine, wound

WOUND HEALING PLANTS

The ancient history of wound healing treatments in several countries was reported; for example, Tirunelveli Hills in Southern India;^[1] Northern Himalaya Range, Abbottabad district, Pakistan;^[2] several districts in Bangladesh;^[3] Kpando area of Volta Region in Ghana;^[4] and Kuruma tribes, Wayanad districts of Kerala, India.^[5] The natural or biological products are studied for wound and burn healing agents in many countries such as India, China, and Thailand.^[6] Because of poor hygienic status, wound infection is still one of the most common diseases in developing countries.^[7] Some examples of wound healing plants include korphad, *Aloe vera*;^[8] Madeira vines, *Anredera diffusa*;^[9] jungle geranium, *Ixora coccinea*;^[10] Indian mulberry, *Morinda pubescens*;^[11] simple-leaf chaste tree, *Vitex trifolia*; and peacock chaste tree, *Vitex altissima*.^[12] The present review provided an up-to-date information about the properties of *Chromolaena odorata*, one of the wound healing plants that is being investigated for its diverse health benefits.

PLANT DESCRIPTION OF CHROMOLAENA ODORATA

C. odorata or Siam weed has a minimum 10-year life span. *C. odorata* is a scrambling perennial shrub which grows 2–3 m in height with straight, pithy, brittle stems that branch readily. The arrowhead-shaped

leaves are 6–12 cm in length and 3–7 cm in width, with three veins in a pitchfork appearance. The leaves grow in opposite pairs along the stems and branches. There are 15–25 tubular florets per head, each 10 mm long and several colors such as white, purple, pink, or blue. The color of seeds is brown-gray to black and is 4–5 mm long with a pale brown pappus that is 5 or 6 mm long. The roots are narrow and fibrous and generally reach 0.3 km in depth.^[13–15] *C. odorata* shows morphological in terms of flower color, leaf shape, odor of the crushed leaves, and plant architecture variable in its native environment.^[16]

TAXONOMICAL CLASSIFICATION

The taxonomy of *C. odorata* is in the Kingdom: *Plantae*; Subkingdom: *Viridiplantae*; Infrakingdom: *Streptophyta*; Superdivision: *Embryophyta*; Division: *Tracheophyta*; Subdivision: *Spermatophytina*; Class: *Magnoliopsida*; Superorder: *Asteranae*; Order: *Asterales*; Family: *Asteraceae*; Genus: *Chromolaena*; Species: *C. odorata*.^[17] The plant genus *Chromolaena* is a genus of the family *Asteraceae* which comprises over 165 species that are distributed across tropical and subtropical regions. The name is derived from the Greek word meaning “color.” Due to its species name “*odorata*,” the leaves exhibit a strong odor when they are crushed.

NOMENCLATURE

C. odorata aka *Eupatorium odoratum* is a weedy herb native of Central and South America, which has spread throughout the tropical and subtropical areas.^[18,19] It was first introduced to Southeast Asia in the 1920s and Africa in around 1940 as a plantation cover crop and has ever since spread worldwide.^[20,21] The vernacular names of *C. odorata*

Correspondence:

Dr. Wannee Jiraungkoorskul,
Department of Pathobiology, Faculty of Science, Mahidol University,
Bangkok 10400, Thailand.
E-mail: pathobiologymu@gmail.com

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include sunflower family, Christmas bush, Jack in the bush, communist weed, Siam weed, devil weed (English); sekou toure, acheampong, jabinde, matapa, mighbe (African); herbe du Laos (French); Siam kraut (German); kesengesil (Guam); bagh dhoka, tivra gandha (Hindi); rumput belalang, rumput putih, rumput golkar (Indonesian); pokok kapal terbang, rumput jepun, rumput Siam (Malayalam); ropani, seekhrasarp (Sanskrit); cariaquillo Santa Maria (Spanish); agono, hagonoy, huluhagonoi (Tagalog); sab suea (Thai); and co hoi (Vietnamese).^[21,22]

PHYTOCHEMICAL SUBSTANCES

The dried leaf of *C. odorata* contained ash (11%), crude fat (11%), fiber (15%), moisture (15%), crude protein (18%), and carbohydrate (31%).^[23] Its active phytochemical substances are as follows: (1) flavonoid aglycones (flavanones, flavonols, flavones) including acacetin, chalcones, eupatilin, luteolin, naringenin, kaempferol, quercetin, quercetagenin, and sinensetin;^[24-31] (2) terpenes and terpenoids;^[32] (3) essential oils;^[33-38] (4) alkaloids including pyrrolizidine;^[39-41] (5) saponins and tannins;^[23] (6) phenolic acids including ferulic acid, protocatechuic acid;^[42] (7) phytoprostane compound including chromomoric acid.^[43]

TRADITIONAL USES

From review literature regarding the traditional uses, phytochemical properties of *C. odorata* are anti-bacterial,^[27,44-47] anticancer,^[21,48] anticonvulsant,^[49] antidiabetic,^[50-52] anti-diarrheal,^[53,54] anti-fungal,^[55,56] anti-inflammatory,^[57-59] antioxidant,^[60-65] and antiparasitic,^[30,40] hemostatic and wound healing,^[15,22,23,66-69] and hepatoprotective activities.^[70,71]

WOUND HEALING PROPERTY

The efficiency of healing wounds come from the antioxidant property of the drug or plant which enhances conserving the fibroblast and keratinocyte proliferation on those wounds.^[22] *C. odorata* is popularly used for traditional wound healing in Vietnam; moreover, the leave aqueous extract has been used for the treatment of soft-tissue burns or skin infections.^[42,60,72,73]

IN VITRO STUDY

Phan *et al.*^[72] reported that Eupolin extract increased the proliferation of fibroblasts, endothelial cells, and keratinocytes in wound assay. Stimulation of keratinocyte migration, upregulation of production by keratinocytes of extracellular matrix proteins and basement membrane components, and protection of collagen lattice contraction by fibroblasts were reported. Moreover, Phan *et al.*^[74] also reported that Eupolin extract enhanced the expression of many adhesion complexes, for example, laminin-5, laminin-1, collagen IV, and fibronectin by human keratinocytes. Pandith *et al.*^[69] reported that *C. odorata* stimulated hemostatic process and wound healing activity by inducing the expression of genes, including heme oxygenase-1, thromboxane synthase, and anti-platelet aggregator matrix metalloproteinase 9 (MMP9). This plant can promote fibroblast cell migration and proliferation. Moreover, they found that heme oxygenase-1, the accelerating wound healing enzyme, was increased at the transcriptional and translational levels by *C. odorata* treatment. Thromboxane synthase, a vasoconstrictor, was increased and MMP-9, an anti-platelet aggregator, was decreased when treated with *C. odorata*.

IN VIVO STUDY

According to the study of Mahmood *et al.*,^[67] adult male Sprague-Dawley rats with wounds in the posterior neck were divided into four groups

for the twice daily application of normal saline, pure unboiled honey, 90% honey in combined with 10% *C. odorata* aqueous leave extract, and solcoseryl jelly. They reported the advantage of honey combined with this extract for the stimulation of wound healing process, decrease scar formation and period of epithelialization, and the rates of wounds sterility. Pandurangan *et al.*^[15] investigated the wound healing activity of 2.5%, 7.5%, and 10% w/w of leaves of *C. odorata* extract ointments for 14 days in rats. Their results revealed that varying concentrations of this herb extract in the ointment base was capable of producing significant cutaneous wound dressing activity by inducing wound contraction and wound closure time.

BLEEDING TIME STUDY

Anyasor *et al.*^[23] reported the aqueous extract of *C. odorata* (coagulation: 15.18 ± 0.023 min; clotting time 0.26 ± 0.014 min) showed significantly higher hemostatic activity than the ethanolic extract (21 min in coagulation time and clotting 2 min in clotting time). Akomas and Ijioma^[75] studied the effect of the oral administration of *C. odorata* in rats for 14 days. This herb significantly lowered bleeding times from 4.5 min in control group to 3.0 and 2.7 min, in low and high doses, respectively. The extract also lowered clotting time from 2.6 min in control group to 1.8 and 1.5 min, respectively. The bleeding and clotting times decreased in animals treated with *C. odorata* extract, suggesting that it remains the good hemostatic property and reduces the bleeding and clotting times by inducing the formation and activation platelets.^[75] The results obtained therefore indicates that *C. odorata* promotes wound healing, by stopping of bleeding which may be the first step in the wound healing mechanism.^[75]

CONCLUSION

C. odorata exhibits its wound healing property using multiple mechanisms. From the literature reviews, these mechanisms can be summarized as follows: (1) *C. odorata* extract contains many antioxidant compounds that enhance wound healing property.^[74] (2) *C. odorata* reduces the bleeding and clotting time may be the first line of action in the physiology of wound healing.^[75] (3) *C. odorata* can protect the cells from destruction by inhibiting the inflammatory mediators.^[73] (4) *C. odorata* has the antibacterial activities against both Gram-positive and Gram-negative bacteria, suggesting that it may reduce the wound infections.^[44] This review article has attempted to compile the new medicinal plant *C. odorata*, to be one of choices in the wound healing treatment.

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Conflicts of interest

There are no conflicts of interest.

REFERENCES

1. Ayyanar M, Ignacimuthu S. Herbal medicines for wound healing among tribal people in Southern India: Ethnobotanical and scientific evidences. *Int J Appl Res Nat Prod* 2009;2:29-42.

2. Abbasi AM, Khan MA, Ahmad M, Qureshi R, Arshad M, Jahan S, *et al.* Ethnobotanical study of wound healing herbs among the tribal communities in Northern Himalaya Ranges district Abbottabad, Pakistan. *Pak J Bot* 2010;42:3747-53.
3. Rani CA, Mohammed R. Ethnobotanical study of wound healing plants among the folk medicinal practitioners of several districts in Bangladesh. *Am Eur J Sustain Agric* 2012;6:371-7.
4. Barku VY, Boahen YO, Dali GA. Ethnobotanical study of wound healing plants in Kpando traditional area, Ghana. *Int J Phytomed* 2014;6:564-72.
5. Thomas B, Arumugam R, Veerasamy A, Ramamoorthy S. Ethnomedicinal plants used for the treatment of cuts and wounds by Kuruma tribes, Wayanadu districts of Kerala, India. *Asian Pac J Trop Biomed* 2014;4 Suppl 1:S488-91.
6. Kumar B, Vijayakumar M, Govindarajan R, Pushpangadan P. Ethnopharmacological approaches to wound healing – Exploring medicinal plants of India. *J Ethnopharmacol* 2007;114:103-13.
7. Hussein J, Mavalankar DV, Sharma S, D'Ambruoso L. A review of health system infection control measures in developing countries: What can be learned to reduce maternal mortality. *Global Health* 2011;7:14.
8. Maenthaisong R, Chaiyakunapruk N, Niruntraporn S, Kongkaew C. The efficacy of *Aloe vera* used for burn wound healing: A systematic review. *Burns* 2007;33:713-8.
9. Moura-Letts G, Villegas LF, Marçalo A, Vaisberg AJ, Hammond GB. *In vivo* wound-healing activity of oleanolic acid derived from the acid hydrolysis of *Anredera diffusa*. *J Nat Prod* 2006;69:978-9.
10. Upadhyay A, Chattopadhyay P, Goyary D, Mitra Mazumder P, Veer V. *Ixora coccinea* enhances cutaneous wound healing by upregulating the expression of collagen and basic fibroblast growth factor. *ISRN Pharmacol* 2014;2014:751824.
11. Mathivanan N, Surendiran G, Srinivasan K, Malarvizhi K. *Morinda pubescens* JE Smith (*Morinda tinctoria* Roxb) fruit extract accelerates wound healing in rats. *J Med Food* 2006;9:591-3.
12. Manjunatha BK, Vidya SM, Krishna V, Mankani KL, Singh SD, Manohara YN. Comparative evaluation of wound healing potency of *Vitex trifolia* L. and *Vitex altissima* L. *Phytother Res* 2007;21:457-61.
13. Henderson L. Alien Weeds and Invasive Plants. Vol. XII. Pretoria: ARC-PPRI; 2001.
14. Chakraborty AK, Rambhade S, Patil UK. *Chromolaena odorata* (L.): An overview. *J Pharm Res* 2011;4:573-6.
15. Pandurangan A, Rana K, Singh A. Evaluation wound healing activity of leaves of *Chromolaena odorata* Linn. *Int J Pharm Sci Lett* 2015;5:555-7.
16. Zachariades C, Day M, Muniappan R, Reddy GV. *Chromolaena odorata* (L.) King and Robinson (*Asteraceae*). In: Muniappan R, Reddy GV, editors. Biological Control of Tropical Weeds Using Arthropods. UK: Cambridge University Press; 2009. p. 130-62.
17. Integrated Taxonomic Information System (ITIS). *Chromolaena odorata* (L.) R.M. King & H. Rob. Taxonomic Serial No.:37034, Geological Survey, VA, USA; 2016.
18. Gautier L. Taxonomy and distribution of a tropical weed, *Chromolaena odorata* (L.) R. King and H. Robinson. *Candollea* 1992;47:645-62.
19. Omokhua AG, McGaw LJ, Finnie JF, Van Staden J. *Chromolaena odorata* (L.) R.M. King & H. Rob. (*Asteraceae*) in sub-Saharan Africa: A synthesis and review of its medicinal potential. *J Ethnopharmacol* 2016;183:112-22.
20. Rouw A. The invasion of *Chromolaena odorata* and competition with the native flora in a rain forest zone, South-west Côte d'Ivoire. *J Biogeogr* 1991;18:13-23.
21. Kouamé PB, Jacques C, Bedi G, Silvestre V, Loquet D, Barillé-Nion S, *et al.* Phytochemicals isolated from leaves of *Chromolaena odorata*: Impact on viability and clonogenicity of cancer cell lines. *Phytother Res* 2013;27:835-40.
22. Vaisakh MN, Pandey A. The invasive weed with healing properties: A review on *Chromolaena odorata*. *Int J Pharm Sci Res* 2012;3:80-3.
23. Anyasor GN, Aina DA, Olushola M, Aniyikawe AF. Phytochemical constituents, proximate analysis, antioxidants, anti-bacterial and wound healing properties of leaf extracts of *Chromolaena odorata*. *Ann Biol Res* 2011;2:441-51.
24. Barua RN, Sharma RP, Thyagarajan G, Hertz W. Flavonoids of *Chromolaena odorata*. *Phytochemistry* 1978;17:1807-8.
25. Wollenweber E, Dorr M, Muniappan R. Exudate flavonoids in a tropical weed, *Chromolaena odorata* (L.) R.M. King & H. Robinson. *Biochem Syst Ecol* 1995;23:873-4.
26. Wollenweber E, Roitman JN. A novel methy ether of quercetagenin from *Chromolaena odorata* leaf exudate. *Biochem Syst Ecol* 1996;24:479-80.
27. Suksamrarn A, Chotipong A, Suavansri T, Boongird S, Timsuksai P, Vimuttipong S, *et al.* Antimycobacterial activity and cytotoxicity of flavonoids from the flowers of *Chromolaena odorata*. *Arch Pharm Res* 2004;27:507-11.
28. Pisutthanan N, Liawruangrath B, Liawruangrath S, Bremner JB. A new flavonoid from *Chromolaena odorata*. *Nat Prod Res* 2006;20:1192-8.
29. Hung TM, Cuong TD, Dang NH, Zhu S, Long PQ, Komatsu K, *et al.* Flavonoid glycosides from *Chromolaena odorata* leaves and their *in vitro* cytotoxic activity. *Chem Pharm Bull (Tokyo)* 2011;59:129-31.
30. Ezenyi IC, Salawu OA, Kulkarni R, Emeje M. Antiplasmodial activity-aided isolation and identification of quercetin-4'-methyl ether in *Chromolaena odorata* leaf fraction with high activity against chloroquine-resistant *Plasmodium falciparum*. *Parasitol Res* 2014;113:4415-22.
31. Emami L, Ravada S, Meka B, Garaga M, Golakoti T. A new flavanone from the leaves of *Chromolaena odorata*. *Nat Prod Commun* 2015;10:1555-9.
32. Wafo P, Kamdem RS, Ali Z, Anjum S, Begum A, Olujemisi OO, *et al.* Kaurane-type diterpenoids from *Chromolaena odorata*, their X-ray diffraction studies and potent α -glucosidase inhibition of 16-kauren-19-oic acid. *Fitoterapia* 2011;82:642-6.
33. Bamba D, Bessière JM, Marion C, Pélissier Y, Fourasté I. Essential oil of *Eupatorium odoratum*. *Planta Med* 1993;59:184-5.
34. Bouda H, Taponjdou LA, Fontem DA, Gumedzo MY. Effect of essential oils from leaves of *Ageratum conyzoides*, *Lantana camara* and *Chromolaena odorata* on the mortality of *Sitophilus zeamais* (Coleoptera, Curculionidae). *J Stored Prod Res* 2001;37:103-9.
35. Pisutthanan N, Liawruangrath B, Liawruangrath S, Baramee A, Apisariyakul A, Korh J, *et al.* Constituents of the essential oil from aerial parts of *Chromolaena odorata* from Thailand. *Nat Prod Res* 2006;20:636-40.
36. Owolabi MS, Ogundajo A, Yusuf KO, Lajide L, Villanueva HE, Tuten JA, *et al.* Chemical composition and bioactivity of the essential oil of *Chromolaena odorata* from Nigeria. *Rec Nat Prod* 2010;4:72-8.
37. Joshi RK. Chemical composition of the essential oils of aerial parts and flowers of *Chromolaena odorata* (L.) R.M. King & H. Rob. From Western Ghats region of North West Karnataka, India. *J Essent Oil Bearing Plants* 2013;16:71-5.
38. Joshi RK. Chemical composition of the essential oil of *Chromolaena odorata* (L.) R. M. King & H. Rob. roots from India. *J Chem* 2013:1-4.
39. Biller A, Boppre M, Witte L, Hartmann T. Pyrrolizidine alkaloids in *Chromolaena odorata*. chemical and chemoeological aspects. *Phytochemistry* 1994;35:615-9.
40. Thoden TC, Boppre M, Hallmann J. Pyrrolizidine alkaloids of *Chromolaena odorata* act as nematocidal agents and reduce infection of lettuce roots by *Meloidogyne incognita*. *Nematology* 2007;9:343-9.
41. Yakubu MT. Effect of a 60-day oral gavage of a crude alkaloid extract from *Chromolaena odorata* leaves on hormonal and spermatogenic indices of male rats. *J Androl* 2012;33:1199-207.
42. Phan TT, Wang L, See P, Grayer RJ, Chan SY, Lee ST. Phenolic compounds of *Chromolaena odorata* protect cultured skin cells from oxidative damage: Implication for cutaneous wound healing. *Biol Pharm Bull* 2001;24:1373-9.
43. Heiss EH, Tran TV, Zimmermann K, Schwaiger S, Vouk C, Mayerhofer B, *et al.* Identification of chromomeric acid C-I as an Nrf2 activator in *Chromolaena odorata*. *J Nat Prod* 2014;77:503-8.
44. Irobi ON. Activities of *Chromolaena odorata* (Compositae) leaf extract against *Pseudomonas aeruginosa* and *Streptococcus faecalis*. *J Ethnopharmacol* 1992;37:81-3.
45. Johari SA, Kiong LS, Mohtar M, Isa MM, Man S, Mustafa S, *et al.* Efflux inhibitory activity of flavonoids from *Chromolaena odorata* against selected methicillin-resistant *Staphylococcus aureus* (MRSA) isolates. *Afr J Microbiol Res* 2012;6:5631-5.
46. Kigigha LT, Zige DV. Activity of *Chromolaena odorata* on enteric and superficial etiologic bacterial agents. *Am J Res Commun* 2013;1:266-76.
47. Stanley MC, Ifeanyi OE, Nwakaego CC, Esther IO. Antimicrobial effects of *Chromolaena odorata* on some human pathogens. *Int J Curr Microbiol Appl Sci* 2014;3:1006-12.
48. Adedapo AA, Oyagbemi AA, Fagbohun OA, Omobowale TO, Yakubu MA. Evaluation of the anticancer properties of the methanol leaf extract of *Chromolaena odorata* on HT29 lung cancer cell line. *The FASEB Journal* 2016;30:Supplement1193.6
49. Amazu LU, Omoregie P, Ajugwo AO, Ifezulike CC, Azikiwe CC. Anticonvulsant potency of the leaf extract of *Chromolaena odorata* in rats. *Unique Res J Med Med Sci* 2013;1:64-9.
50. Onkaramurthy M, Veerapur VP, Thippeswamy BS, Reddy TN, Rayappa H, Badami S. Anti-diabetic and anti-cataract effects of *Chromolaena odorata* Linn. in streptozotocin-induced diabetic rats. *J Ethnopharmacol* 2013;145:363-72.
51. Ijioma SN, Okafor AI, Ndokuba PI, Nwankwo AA, Akomas SC. Hypoglycemic, hematologic and lipid profile effects of *Chromolaena odorata* ethanol leaf extract in alloxan induced diabetic rats. *Ann Biol Sci* 2014;2:27-32.
52. Uhegbu FO, Imo C, Onwuegbuchulam CH. Lipid lowering, hypoglycemic and antioxidant

- activities of *Chromolaena odorata* (L) and *Ageratum conyzoides* (L) ethanolic leaf extracts in albino rats. *J Med Plants Stud* 2016;4:155-9.
53. Atindehou M, Lagnika L, Guérold B, Strub JM, Zhao M, Dorsselaer AV, et al. Isolation and identification of two anti-bacterial agents from *Chromolaena odorata* L. active against four diarrheal strains. *Adv Microbiol* 2013;3:115-21.
54. Aba PE, Joshua PE, Ezeonugu FC, Ezeja MI, Omoja VU, Umeakuana PU. Possible anti-diarrhoeal potential of ethanol leaf extract of *Chromolaena odorata* in castor oil-induced rats. *J Complement Integr Med* 2015;12:301-6.
55. Ngono NA, Ebelle Etame R, Ndifor F, Biyiti L, Amvam ZP, Bouchet P. Antifungal activity of *Chromolaena odorata* (L.) King and Robinson (*Asteraceae*) of Cameroon. *Chemotherapy* 2006;52:103-6.
56. Naidoo KK, Coopposamy RM, Naidoo G. Screening of *Chromolaena odorata* (L.) King and Robinson for anti-bacterial and anti-fungal properties. *J Med Plant Res* 2011;5:4859-62.
57. Owoyele VB, Adediji JO, Soladoye AO. Anti-inflammatory activity of aqueous leaf extract of *Chromolaena odorata*. *Inflammopharmacology* 2005;13:479-84.
58. Hanh TT, Hang DT, Minh CV, Dat NT. Anti-inflammatory effects of fatty acids isolated from *Chromolaena odorata*. *Asian Pac J Trop Med* 2011;4:760-3.
59. Pandith H, Zhang X, Thongpraditchoe S, Wongkrajang Y, Gritsanapan W, Baek SJ. Effect of Siam weed extract and its bioactive component scutellarein tetramethyl ether on anti-inflammatory activity through NF- κ B pathway. *J Ethnopharmacol* 2013;147:434-41.
60. Phan TT, See P, Lee ST, Chan SY. Anti-oxidant effects of the extracts from the leaves of *Chromolaena odorata* on human dermal fibroblasts and epidermal keratinocytes against hydrogen peroxide and hypoxanthine-xanthine oxidase induced damage. *Burns* 2001;27:319-27.
61. Akinmoladun AC, Ibukun EO, Dan-Ologe IA. Phytochemical constituents and antioxidant properties of extracts from the leaves of *Chromolaena odorata*. *Sci Res Essays* 2007;2:191-4.
62. Melinda KP, Rathinam X, Marimuthu K, Diwakar A, Ramanathan S, Kathiresan S, et al. A comparative study on the antioxidant activity of methanolic leaf extracts of *Ficus religiosa* L., *Chromolaena odorata* (L.) King & Robinson, *Cynodon dactylon* (L.) Pers. and *Tridax procumbens* L. *Asian Pac J Trop Med* 2010;3:348-50.
63. Rao KS, Chaudhury PK, Pradhan A. Evaluation of anti-oxidant activities and total phenolic content of *Chromolaena odorata*. *Food Chem Toxicol* 2010;48:729-32.
64. Vijayaraghavan K, Mohamed Ali S, Maruthi R. Studies on phytochemical screening and antioxidant activity of *Chromolaena odorata* and *Annona squamosa*. *Int J Innov Res Sci Eng Technol* 2013;2:7315-21.
65. Boudjeko T, Megnekou R, Woguia AL, Kegne FM, Ngomoyogoli JE, Tchapoum CD, et al. Antioxidant and immunomodulatory properties of polysaccharides from *Allanblackia floribunda* Oliv stem bark and *Chromolaena odorata* (L.) King and H.E. Robins leaves. *BMC Res Notes* 2015;8:759.
66. Akah PA. Mechanism of hemostatic activity of *Eupatorium odoratum*. *Int J Crude Drug Res* 1990;28:253-6.
67. Mahmood AA, Indran M, Salmah I, Sidix K, Suzainur KA. Evaluation of *in vivo* wound healing activity of *Chromolaena odorata* leaf extract on excision wounds model in rats. *J Food Technol* 2005;3:126-9.
68. Pandith H, Thongpraditchoe S, Wongkrajang Y, Gritsanapan W. *In vivo* and *in vitro* hemostatic activity of *Chromolaena odorata* leaf extract. *Pharm Biol* 2012;50:1073-7.
69. Pandith H, Zhang X, Liggett J, Min KW, Gritsanapan W, Baek SJ. Hemostatic and wound healing properties of *Chromolaena odorata* leaf extract. *ISRN Dermatol* 2013;(2013):168269, 8 pages.
70. Alisi CS, Onyeze GO, Ojako OA, Osuagwu CG. Evaluation of the protective potential of *Chromolaena odorata* Linn. extract on carbon tetrachloride-induced oxidative liver damage. *Int J Biochem Res Rev* 2011;1:69-81.
71. Asomugha RN, Okafor PN, Ijeh II, Orisakwe OE, Asomugha AL. Hepatic effects of aqueous extract of *Chromolaena odorata* in male Wistar albino rats. *Pharmacol Online* 2014;1:127-36.
72. Phan TT, Hughes MA, Cherry GW. Enhanced proliferation of fibroblasts and endothelial cells treated with an extract of the leaves of *Chromolaena odorata* (Eupolin), an herbal remedy for treating wounds. *Plast Reconstr Surg* 1998;101:756-65.
73. Thang PT, Patrick S, Teik LS, Yung CS. Anti-oxidant effects of the extracts from the leaves of *Chromolaena odorata* on human dermal fibroblasts and epidermal keratinocytes against hydrogen peroxide and hypoxanthine-xanthine oxidase induced damage. *Burns* 2001;27:319-27.
74. Phan TT, Allen J, Hughes MA, Cherry G, Wojnarowska F. Upregulation of adhesion complex proteins and fibronectin by human keratinocytes treated with an aqueous extract from the leaves of *Chromolaena odorata* (Eupolin). *Eur J Dermatol* 2000;10:522-7.
75. Akomas SC, Ijioma SN. Bleeding and clotting time effect of ethanolic extracts of *Chromolaena odorata* versus *Ocimum gratissimum* treated albino rats. *Compr J Med Sci* 2014;2:9-13.