Research and Reviews: Journal of Chemistry Aloe Vera - A Mini Review

Gayathri P*

MNR College of Pharmacy, Sangareddy, Medak District, Telangana, India

Review Article

Received: 28/08/2016 Accepted: 02/09/2016 Published: 30/09/2016

ABSTRACT

*For Correspondence

Gayathri P, MNR College of Pharmacy, Sangareddy, Medak District, Telangana, India, Tel: +919701764943

E-mail: gayathri28@gmail.com

Keywords: Aloe Vera, herbal medicine, skin

Aloe Vera is a characteristic item that is currently a day much of the time utilized as a part of the field of cosmetology. In spite of the fact that there are different signs for its utilization, controlled trials are expected to decide its genuine adequacy. The aloe Vera plant, its properties, instrument of activity and clinical uses are quickly explored in this article.

INTRODUCTION

Aloe vera (syn. Aloe barbadensis Mill., Fam. Liliaceae), otherwise called Barbados or Curaçao Aloe, has been utilized as a part of customary and society solutions for a great many years to treat and cure an assortment of maladies. Inspite of the fact that the plant is local to northern parts of Africa, it has quickly spread over the world since its development is simple. An imperative qualification must be made between the firmly diuretic and laxative latex got from the pack sheath cells and the reasonable adhesive gel. The plant has been utilized by Egyptians, Assyrians, and Mediterranean human advancements, and additionally in Biblical times. An assortment of aloe animal types are still utilized as a part of people prescriptions of Africa and Asia. Seekers in the Congo apparently rub their bodies free adhesive gel to diminish sweat; some African tribes apply the gel for constant conjunctivitis; the gel is utilized as a part of India for the treatment of asthma. ^[1-10]

Description

Aloe vera is a succulent plant with thick, plump, serrated, lanceolate-formed leaves of green-greyish shading. Aloe Vera inward gel is gotten from the lower leaves of the plant by cutting the leaf open. The gel is clear, unscented, and bland and ought to be free of leaf skin or yellow parts. No reliable institutionalization has been built up, yet the International Aloe Science Council (IASC), an exchange relationship of globally based aloe makers and advertisers, obliges adherence to specific for the item to be certified.10 Other arrangements incorporate a hydrophilic cream containing 0.5% aloe vera gel and an emulsion comprising of 30% aloe Vera gel (Figure 1).



Figure 1. Aloe vera leaf.

CHEMICAL COMPOSITION

The aloe gel predominantly comprises of water (99.1%) and mesophyll cells (0.9% dry matter), which can be isolated into 3 particular portions: cell divider, micro particles, and fluid gel [accounting for 16.2%, 0.7%, and 83.1% of dry mash (w/w), respectively]. The transcendent sugar part is mannose as mannose-6-phosphate36 in each of the 3 divisions [20.4% in cell divider, 32.2% in micro particles, and 62.9% in the fluid gel (% of aggregate sugars)], trailed by different sugars in changing focuses relying upon the portion. By and large, the 5 impartial sugars (i. e. arabinose, xylose, mannose, galactose, glucose) represent 69.2% of the aggregate sugars in the gel.37 Muco polysaccharides are mostly present as acemannan [a exceedingly acetylated, β -1-4-connected polysaccharide (> 1kDa) made for the most part of mannose] with different side chain glycosylation patterns.38 The anthra guinone substance ought to be beneath 50 ppm and is viewed as a debasement from the leaf concentrate of aloe vera.7 Other fixings incorporate different amino acids, proteins, and vitamins, which have not been evaluated. The IASC keeps up a confirmation program, in which "entire aloe vera leaf gel" needs to stick to the accompanying details: solids (0.46% - 1.31%); pH (3.5-4.7); calcium(98.2-448 mg/L); magnesium (23.4-118 mg/L; malic corrosive (817.8-3.427.8 mg/L); acemannan in crude materials (\geq 5% by dry weight); isocitrate (\leq 5% for inward leaf by dry weight); crude materials powder content (\leq 40%); aloin (\leq 10 ppm in 0.5% aloe vera solids answer for oral utilization). Quality items ought to contain high sums (95%) of unadulterated aloe vera gel.39 One method for evaluating aloe polysaccharides is a colorimetric test, which has been proposed for use in quality control of business products.40 Ouality control and recognizable proof of business aloe vera items has additionally been proficient by atomic attractive reverberation spectrometry. [11-20]

Mechanism of action

Incitement of macrophage and fibroblast movement, expanded collagen and proteoglycan synthesis. Mannose-6-phosphate ties to development element receptor on fibroblasts and improves their activity Macrophage actuation through expanded nitric oxide synthase action by acemannan, prompting arrival of fibrogenic cytokines. Upregulation of phagocytosis and fungicidal movement of macrophages by acemannan95 Acemannan and other cell divider biomaterial may advance steadiness of development components and drag out incitement of granulation tissue48,80 Restraint of Thromboxan A2. May advance hypoglycemic impact by normalizing film bound protein exercises of phosphatases and hydrolases and expanded glucose metabolism; potential dynamic mixes incorporate the phytosterols lophenol, cycloartenol and their alkylated derivatives Mitigating impact of plant sterols like lupeol, campesterol, and β -sitosterol through bradikinase activation, prostaglandin F2 and E2, and in addition thromboxane A2 inhibitionand hindrance of IL-10 secretion Inhibitory impact on arrival of receptive oxygen species from human neutrophils by decreasing intracellular free calcium levels Increment in mRNA articulation of metalloproteinases and plasminogen activator may prompt angiogenic action in endothelial cells91^[21-30]



Figure 2. Aloe vera plant.

CLINICAL USES

Seborrheic dermatitis, psoriasis vulgaris, genital herpes, skin burns, diabetes (type 2), HIV infection, cancer prevention, ulcerative colitis, wound healing (results of aloe on wound healing are mixed with some studies reporting positive results and others showing no benefit or potential worsening), pressure ulcers, mucositis, radiation dermatitis, acne vulgaris lichen planus, frostbite, aphthous stomatitis, and constipation.

Pregnancy and breastfeeding: Oral aloe is not recommended during pregnancy due to theoretical stimulation of uterine contractions, and in breastfeeding mothers, it may sometime cause gastrointestinal distress in the nursing infant.^[31-40]

Used for treating animal diseases

Speeding up of twisted recuperating in mice and rats; lessening of radiation-affected skin responses in illuminated mice and rats; counteractive action of dynamic dermal ischaemia brought about by blazes and frostbite in rats and guinea pigs; antidiabetic in sort 2 diabetic and insulin-safe mice; chemo preventive in skin papillo magenesis in mice; calming in mice; improvement of safe responsiveness in chicks and mice; enhancement of UV-impelled resistant concealment in mice; advancement of gastric ulcer mending in rats; assurance of liquor dehydrogenase and decrease of blood ethanol fixations in rats; diminishment of salmonella-intervened irritation in mice; cancer prevention agent and cholesterol-bringing down impacts in matured rats ^[41-60].

Adverse effects

Topical utilization of aloe vera arrangements has been viewed as sheltered as surveyed by the Cosmetic Ingredient Review Expert Panel. However, a few case reports of the improvement of extreme touchiness responses and contact dermatitis in light of topically connected aloe gel arrangements have been published. This unfavourably susceptible response has been ascribed much of the time to anthraquinone defilements in the gel. Macrophage penetration and emesis has been seen in canines treated intravenously with acemannan. Oral use of aloe vera gel may bring down blood glucose levels and upgrade the movement of antidiabetic treatments. No genotoxic impacts have been watched taking after oral organization of an aloe vera inward leaf gel (Qmatrix® by Aloecorp, Inc., which is an institutionalized internal gel extricate that has not been warmed after extraction from the leaf) to rats after 90 days. An essential element for antagonistic impacts is the virtue of the aloe vera gel utilized, since anthraquinones like aloin may be identified with the advancement of excessive touchiness reactions. [61-80].

Cosmetic uses of aloe Vera

- Prevents Signs of Aging
- Moisturizes Skin
- Reduces Acne and Helps Lighten Blemishes

- Helps with Sunburns And Reduces Tan
- Heals External Wounds and Insect Bites
- Reduces Stretch Marks
- Promotes Hair Growth
- Reduces Dandruff
- Maintains pH Balance of the Scalp
- Conditions Hair
- Reduces Inflammation
- Eases Heartburn and Acid Reflux
- Reduces Cholesterol and Regulates Blood Sugar
- Maintains Oral Health
- Builds Immunity
- Lowers Risk of Cancer
- Helps in Treating Haemorrhoid

CONCLUSION

Taking everything into account, the utilization of aloe Vera gel or its segments for the treatment of an assortment of conditions and infections needs promote clinical confirmation through very much composed studies with characterized aloe extricates and coordinating fake treatment controls. Presently (June 2012), 5 national and universal clinical studies are recorded by the United States National Institutes of Health clinical trial database with a noteworthy accentuation on the utilization of aloe Vera in the treatment of wounds.122 This shows the logical hugeness of aloe Vera gel and the need to build up it as a substantial treatment choice for wounds. Be that as it may, the utilization of aloe Vera gel in topical applications has broadly been affirmed in the clinical studies as protected.

REFERENCES

- 1. Amanda M. Growing up with pesticides. Sci 2013; 341:40-41.
- Douiri LF and Boughdad A. Chemical composition and biological activity of Allium sativum essential oils against Callosobruchus maculatus, Mohieddine Moumni. IOSR-JESTFT 2013; 3: 30-36.
- 3. Dubey NK and Srivastavab KA. Current Status of Plant Products as Botanical Pesticides in storage pest management. J Bio Pest 2008; 1:182-186.
- 4. Brock T and Arts GH. Aquatic risks of pesticides, ecological protection goals, and common aims in European Union legislation. Integrated Environmental Assessment and Management. 2006;2: e20-e46.
- 5. Roger C and Vincent C. Essential oils in insect control: low-risk products in a high-stakes world. Annu Rev Entomol 2012;57: 405-424.

- Bakkali F and Averbeck S. Biological effects of essential oils. A review. Food Chem Toxicol. 2008; 46:446-475.
- 7. Mannan A and Ahmed I. Survey of artemisinin production by diverse Artemisia species in northern Pakistan. Malar J 2010;9: 310.
- 8. Noedl H, et al. Artemisinin-resistant malaria in Asia. N Engl J Med 2009;361: 40-541.
- Kordali S, et al. Toxicity of essential oils isolated from three Artemisia species and some of their major components to granary weevil, Sitophilus granarius (L.)(Coleoptera: Curculionidae). Industrial crops and products. 2006; 3:162-170.
- 10. Negahban M, et al. Fumigant toxicity of essential oil from Artemisia sieberi Besser against three storedproduct insects. Journal of Stored Products Research 2007; 43:123-128.
- 11. You C, et al. Identification of repellent and insecticidal constituents from Artemisia mongolica essential oil against Lasioderma serricorne. Journal of Chemistry 2015; 549057: 1-7.
- 12. Zhang WJ, et al. Bioactivity of essential oil of Artemisia argyi Lévl. et Van. and its main compounds against Lasioderma serricorne. J Oleo Sci 2014; 63: 829-837.
- 13. Zhang WJ, et al. Bioactivity of essential oil from Artemisia stolonifera (Maxim.) Komar. and its main compounds against two stored-product insects. J Oleo Sci 2015; 64:299-307.
- 14. Tomus Delectis Florae Reipublicae Popularis Sinicae Acendae Academiae Sinicae. Flora of China, 1999; Science Press, Beijing, China.
- 15. Chu SS, et al. Chemical composition and insecticidal activity against Sitophilus zeamais of the essential oils derived from Artemisia giraldii and Artemisia subdigitata. Molecules 2012;7: 7255-7265.
- 16. Papadopoulou SC and Athanassiou CG. Lariophagus distinguendus (F.)(Hyme., Chalcidoidea, Pteromalidae), an ectoparasitoid of Lasioderma serricorne (F.)(Col., Anobiidae), found for the first time in tobacco stores in Greece. Journal of Pest Science 2004;77: 183-184.
- 17. Abdelghany AY, et al Stored-product insects in botanical warehouses. Journal of stored products research 2010;46: 93-97.
- 18. Adams RP, Identification of Essential Oil Components by Gas Chromatography/Quadrupole Mass Spectroscopy. Journal of the American Society for Mass Spectrometry 2001; 16:1902-1903.
- 19. Cannon RJ, et al. Identification, synthesis, and characterization of novel sulfur-containing volatile compounds from the in-depth analysis of Lisbon lemon peels (Citrus limon L. Burm. f. cv. Lisbon). Journal of agricultural and food chemistry 2015;63: 1915-1931.
- 20. Lin J, et al Chemical composition, antimicrobial and antitumor activities of the essential oils and crude extracts of Euphorbia macrorrhiza. Molecules 2012;17: 5030-5039.
- 21. Maggio A, et al. Chemical composition of the essential oils of three endemic species of Anthemis Sect. Hiorthia (DC.) R. Fern. growing wild in Sicily and chemotaxonomic volatile markers of the genus Anthemis L.: an update. Chemistry & biodiversity 2014;11: 652-72.
- 22. Yu CW, et al. Essential oil alloaromadendrene from mixed-type Cinnamomum osmophloeum leaves prolongs the lifespan in Caenorhabditis elegans. J Agric Food Chem 2014; 62: 6159-6165.
- 23. Liu ZL and Ho SH. Bioactivity of the essential oil extracted from Evodia rutaecarpa Hook f. et Thomas against the grain storage insects, Sitophilus zeamais Motsch. and Tribolium castaneum (Herbst). Journal of Stored Products Research 1999; 35:317-328.
- 24. Sakuma M, Probit analysis of preference data. Applied entomology and zoology 1998;33: 339-347.

- 25. XiaoYang D and XinRong D. Chemical constituents of fresh Artemisia ruhripes Nakai leaves. Acta Botanica Boreali-Occidentalia Sinica 2010;30: 259-1263.
- 26. Wang Y, et al. Chemical constituents and insecticidal activities of the essential oil from Amomum tsaoko against two stored-product insects. J Oleo Sci 2014;63: 019-1026.
- 27. Wang Y, et al. Bioactivity of Essential Oil of Zingiber purpureum Rhizomes and Its Main Compounds against Two Stored Product Insects. J Econ Entomol 2015;108: 925-932.
- 28. Wang Y, et al. Chemical constituents and insecticidal activities of the essential oil from Alpinia blepharocalyx rhizomes against Lasioderma serricorne. J Serb Chem Soc 2015;80: 171-178.
- 29. Rattan RS, Mechanism of action of insecticidal secondary metabolites of plant origin. Crop protection 2010;29: 913-920.
- 30. Mills C, Et al. Inhibition of acetylcholinesterase by Tea Tree oil. J Pharm Pharmacol 2004; 56: 375-379.
- 31. Shaaya E and Rafaeli A. Essential oils as biorational insecticides potency and mode of action. InInsecticides design using advanced technologies. Springer, Berlin Heidelberg, 2007; pp: 249-261.
- 32. Felipe CF, et al. Alterations in behavior and memory induced by the essential oil of Zingiber officinale Roscoe (ginger) in mice are cholinergic-dependent. Journal of Medicinal Plants Research 2008; 2:163-170.
- 33. Bloomquist JR, Chloride channels as tools for developing selective insecticides. Arch Insect Biochem Physiol 2003; 54:145-156.
- 34. Priestley CM, et al. Thymol, a constituent of thyme essential oil, is a positive allosteric modulator of human GABAA receptors and a homo-oligomeric GABA receptor from Drosophila melanogaster. British journal of pharmacology 2003; 140:1363-1372.
- 35. Enan EE, et al. Molecular response of Drosophila melanogaster tyramine receptor cascade to plant essential oils. Insect Biochem Molec 2005; 35:309-321.
- 36. López MD and Pascual-Villalobos MJ Mode of inhibition of acetylcholinesterase by monoterpenoids and implications for pest control. Industrial Crops and Products 2010;31: 284-288.
- 37. Tak JH and Isman MB. Enhanced cuticular penetration as the mechanism for synergy of insecticidal constituents of rosemary essential oil in Trichoplusiani. Sci Rep 2015; 5:12690.
- 38. Zhukovskaya MI, Aminergic regulation of pheromone sensillae in the cockroach Periplaneta americana. Journal of Evolutionary Biochemistry and Physiology 2007; 43:318-326.
- 39. Tong F and Coats JR Quantitative structure–activity relationships of monoterpenoid binding activities to the housefly GABA receptor. Pest management science 2012;68: 122-1129.
- 40. Capinera JL. Insecticides and Wildlife. Entomol and Nematology 2014; pp: 1-18.
- 41. Verdi LG et al Gênero Baccharis (Asteraceae): Aspectos químicos, econômicos e biológicos. Quim Nova 2005;28: 85-94.
- 42. Karam TK, et al Carqueja (Baccharis trimera): utilização terapêutica e biossíntese. Rev Bras Plantas Med 2013; 15:280-286.
- 43. Heiden G and Pirani JR. Two new species of Baccharis subgen. Baccharis (Asteraceae, Astereae) with singleflowered female capitula from the Serra do Cipó, Minas Gerais, Brazil. Phytotaxa 2014;164: 141-148.
- 44. Gamberini MT, et al Inhibition of gastric secretion by a water extract from Baccharis triptera, Mart. Mem Inst Oswaldo Cruz 1991;86: 37-139.
- 45. Abad MJ and Bermejo P. Baccharis (Compositae): a review update. Arkivoc 2007; 7:76-96.

- 46. Trojan-Rodrigues M, et al. Plants used as antidiabetics in popular medicine in Rio Grande do Sul, southern Brazil. J Ethnopharmacol 2012; 139:155-163.
- 47. Florão A, et al. Essential oils from Baccharis species (Asteraceae) have anti-inflammatory effects for human cells. J Essent Oil Res 2012; 24:561-570.
- 48. Rezende TP, et al. Protective Effects of Baccharis dracunculifolia Leaves Extract against Carbon Tetrachloride- and Acetaminophen-Induced Hepatotoxicity in Experimental Animals. Molecules 2014; 19: 9257-9272.
- Toyama DO, et al. Effect of Chlorogenic Acid (5-Caffeoylquinic Acid) Isolated fromBaccharis oxyodonta on the Structure and Pharmacological Activities of Secretory Phospholipase A2 from Crotalus durissus terrificus. Biomed Res Int 2014;2014: 1-10.
- 50. Campos FR, et al. Baccharis (Asteraceae): Chemical Constituents and Biological Activities. Chem Biodivers 2016; 13:1-17.
- 51. Godard A, et al. Etude de la métallation des carbamates d'hydroxy-5, -6, -7 et -8 quinoéine. J Organometallic Chem.1987;336(1-2):1.
- 52. Katritzky AR and Lagowski JM. Prototropictautomerism of heteroaromafic compounds. IV. Five-membered rings with two or more hetero atoms. Adv Heterocyclic Chem. 1963; 2:27.
- 53. Shoji E, et al. Immiscible polymers in double spin-coated electroluminescent devices containing phenylsubstituted tris(8-hydroxyquinoline) aluminum derivatives soluble in a host polymer. J Polym Sci Part A. Polym Chem. 2003; 4:3006.
- 54. Manske RHF and Fulka M. The Skraup Synthesis of Quinolines.Organic Reactions, New York. 1953; 7:59-98.
- 55. Nakano Y and Imai D. Synthesis of 5-substituted quinolin-8-ols. Synthesis.1997;12:1425.
- 56. SulimanFO and Al-BusafiSN. Synthesis, characterization and DFT investigation of aluminum complexes of aryl- substituted-8-hydroxyquinoline. Dyes and Pigments. 2012; 92:1153-1159.
- 57. Heiskanen JP and Hormi OEO. 4-Aryl-8-hydroxyquinolines from 4-chloro-8-tosyloxyquinoline using a Suzuki– Miyaura cross-coupling approach. Tetrahedron. 2009;65(2):518-524.
- 58. Tomlin CDS, et al The Pesticide Manual, 11th Ed. British Crop Protection Council, UK. 1997:1606.
- 59. Gershon H and Clarke DD. Evidence of steric factors in the fungitoxic mechanisms of 8-quinolinol and its 2-, 3-, 4-, 5-, 6- and 7-chloro and bromo analogs. Monatsh Chem. 1994; 125:51-59.
- 60. Lee C-H and Jeon J-H. Insecticidal properties of Euphorbiaceae: Sebastiania corniculata-derived 8hydroxyquinoline and its derivatives against three planthopper species (Hemiptera: Delphacidae). J Korean Soc Appl Biol Chem. 2010; 53:464-469.
- 61. Gershon H and Gershon M. Antifungal activity of substituted 8-quinolinol-5- and 7-sulfonic acids: a mechanism of action is suggested based on intramolecular synergism. Mycopathologia. 2002; 155:213-217.
- 62. Darby CM and Nathan CF. Killing of non-replicating Mycobacterium tuberculosis by 8-hydroxyquinoline. J Antimicrob Chemother. 2010; 65:1424-1427.
- 63. Benjamin RD and Short MA. In vitro activity of a novel compound, the metal ion chelating agent AQ+, against clinical isolates of Staphylococcus aureus. J Antimicrob Chemother. 2006; 57:104-109.
- 64. Patel KB and Nimavat KS. Synthesis, characterization and comparative microbial screening of some 5alkoxymethyl-8-quinolinol. Res J Pharm Biol Chem Sci.2012;3;838-844.

- 65. Shen AY and Chenb CP Chelating agent possessing cytotoxicity and antimicrobial activity: 7morpholinomethyl-8-hydroxyquinoline. Life Sci.1999; 64:813-825.
- 66. Enquist PA et al. Derivatives of 8-hydroxyquinoline antibacterial agents that target intra- and extracellular Gram-negative pathogens. Bioorg Med Chem Lett. 2012; 22:3550-3553.
- 67. Collery P, et al. Inhibitory effects of gallium chloride and tris(8-quinolinolato) gallium(III) on A549 human malignant cell line. Anticancer Res. 2000; 20:955-958.
- 68. Corce V et al, Polyaminoquinoline Iron Chelators for Vectorization of Antiproliferative Agents: Design, Synthesis, and Validation. Bioconjugate Chem. 2012; 23:1952-1968.
- 69. Oliveri V, et al. Gluconjugates of 8-hydroxyquinolines as potential anti-cancer prodrugs. Dalton Trans. 2012; 41(15):4530-4535.
- 70. Zouhiri F, et al. J Structure-Activity Relationships and Binding Mode of Styrylquinolines as Potent Inhibitors of HIV-1 Integrase and Replication of HIV-1 in Cell Culture. J Med Chem. 2000; 43:1533-1540.
- 71. CrouchPJ, et al., Mechanisms of Aß mediated neurodegeneration in Alzheimer's disease. Cell Biol. 2008; 40:81-198.
- 72. Bush Al. Et al. The metallobiology of Alzheimer's disease.Trends Neurosci.2003;26:207-214.
- 73. OrlyW, et al. Multifunctional neuroprotective derivatives of rasagiline as anti-Alzheimer's disease drugs. Neurotherapeutics.2009;6:163-174.
- 74. Cherny RA, et al. Aqueous dissolution of Alzheimer's disease Aß amyloid deposits by biometal depletion.J. Biol. Chem.1999;274:23223-23228.
- 75. Adlard PA, et al. Rapid restoration of cognition in Alzheimer's transgenic mice with 8-hydroxyquinoline analogs is associated with decreased interstitial AB. Neuron.2008;59: 43-55.
- 76. Kaur D, et al. Genetic or pharmacological iron chelation prevents MPTP-induced neurotoxicity in vivo: A novel therapy for Parkinson's disease. Neuron. 2003; 37:899-909.
- 77. Tang CW and Van Slyke SA. Organic electroluminescent diodes. Appl Phys Lett. 1987;51:913-915.
- Hopkins TA, et al. Substituted aluminum and zinc quinolates with blue-shifted absorbance/luminescence bands: synthesis and spectroscopic, photoluminescence, and electroluminescence characterization. Chem. Mater.1996;8:344-351.
- 79. Ballardini R, et al. Phosphorescent 8-quinolinol metal chelates Excited-state properties and redox behavior. Inorg Chem. 1986; 25:3858-3865.
- 80. Wang S.et al. Luminescence and electroluminescence of Al(III), B(III), Be(II) and Zn(II) complexes with nitrogen donors. Coord Chem Rev. 2001; 215:79-98.