

**SRI LANKAN MEDICINAL PLANT
MONOGRAPHS AND ANALYSIS**

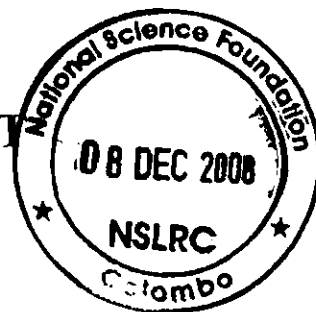
VOL - 8

ALOE VERA



**Lakshmi Arambewela
Sachintha Alagiyawanna**

SRI LANKAN MEDICINAL PLANTS
MONOGRAPHS AND ANALYSIS
VOL - 8



ALOE VERA

Lakshmi Arambewela and Sachintha Alagiyawanna

Edited by

Dilmani Warnasuriya

Industrial Technology Institute
(Ceylon Institute of Scientific and Industrial Research)
363, Bauddhaloka Mawatha
Colombo 07
Sri Lanka

Published by
National Science Foundation

2006

©Industrial Technology Institute & National Science Foundation, 2006.
First published in 2006.

ISBN 955-590-044-2

Published by National Science Foundation.

Arambewela, Lakshmi

Sri Lankan Medicinal plant Monograph and Analysis:

Aloe vera / Lakshmi Arambewela and Sachintha Alagiyawanna.-

Colombo: National Science Foundation, 2006.

Vol 8, iii, 38p. ill.; 30 cm. -(Sri Lankan Medicinal Plants, Vol. 8)

ISBN 955-590-044-2

Price: Rs. 200.00

i. 615.321 DDC 21

ii. Title

iii. Series

iv. Arambewela, Lakshmi jt.au.

v. Alagiyawanna, Sachintha jt.au.

1. Medicinal plants

2. Botany, Chemistry, Traditional medicine

The information found in this monograph is taken from available scientific literature. The authors accept no liability for any damages arising from any claims contained in this text except the work carried out by the authors.

Preface

Studies on medicinal plants of Sri Lanka have been carried out in the Herbal Technology Division of Industrial Technology Institute (former Ceylon Institute of Scientific and Industrial Research) for almost two decades. This monograph which is the eighth in this series incorporates information collected from literature surveys, researches and also experiences of the Herbal Technology Division staff. This monograph is intended for a varied reading public, herbal drug manufacturers who need to identify their herbal raw materials, Ayurvedic physicians who need some scientific information on medicinal plants, research workers requiring some quick background information on a plant, industrialists or entrepreneurs pondering on commercial ventures and the inquiring lay readers. We hope this monograph fulfils some requirements of each of them.

The authors wish to thank the members of the Herbal Technology Division for their contribution, the Information Service Center for providing information, Department of Plant Sciences and Department of Zoology of the University of Colombo for assisting in anatomical studies, Food Technology Division of the Industrial Technology Institute for helping in the analysis of powdered plant materials and the Microbiology Laboratory for photographing the slides. They also gratefully acknowledge the sponsor National Science Foundation for the research grant (RG / 2004 / TM / 01).

Herbal Technology Division
Industrial Technology Institute
P.O.Box 787
Colombo 07
Sri Lanka.

Aloe vera Linn.

Family

Liliaceae

Synonyms

Aloe vera (L) Burm. F., *Aloe vera* L. var. *littoralis* Konig ex Bak., *Aloe vera* L. var. *chinensis* Berger, *Aloe barbadensis* Mill., *Aloe perfoliata* L. Var. *Vara* L., *Aloe chinensis* Bak., *Aloe elongata* Murray., *Aloe indica* Royle., *Aloe vulgaris* Lam., *Aloe officinalis* Forsk., *Aloe perfoliata* L., *Aloe rubescens* DC^{1,2,3}.

Selected Vernacular Names

Sinhala – Komarika^{1,4}

English – Aloe^{1,4,5}

Tamil – Angai, Angini, Kodyyan, Kattali, Sirukattalai, Sottukkattalai, Veligam^{1,3}

Hindi – Ghiguvara, Ghikumari, Gvarapatha, kumari, kuvarapatha, Ghikunwar^{1,3}

Sanskrit – Adala, Ajara, Amara, Ambudhisvara, Aphala, Bahupatri, Bhringeshta, Brahmagni, Dirghapatrika, Grihakanya, Gritakumari, kantakani, Kanya, Kapila, Kumari, Mandala, Mata, Rama, Rasayani, Saha, Sthaleruha, Sthuladala, Sukantaka, Surasa, suvaha, tarani, Vipulasrava, Vira^{1,4}

Bengal- Ghritakumari, Kanya³

Malay- Kattuvala, Kattarvala³

Arabic – Musabar⁵

Chinese – Lu Hui⁵

French – Aloes⁵

German – Aloe⁵

Pharmacopoeia

Ayurveda Pharmacopoeia⁴

British Pharmacopoeia⁶

Pharmacopoeia of United States of America⁷

European Pharmacopoeia⁸

Indian Pharmacopoeia⁹

Distribution^{1,2}

The plant is native to North Africa and the desert regions of Arabia. It is naturalized in most of the tropics and warmer areas of the world including India, West Indies, Mediterranean countries, China and Sri Lanka. It is commercially cultivated in many countries including United States of America and Venezuela.

Morphology¹

A perennial herb with a very short, thick, cylindrical, woody stem sending out at the base numerous stolons; roots fibrous and fleshy; leaves not very numerous, sessile, densely crowded on the short stem with wide dilated bases, spreading below and then ascending, 30-45 cm long tapering to a blunt point, very thick and fleshy, flat or rather concave above, convex beneath, the margins set with hard, distinct, somewhat hooked prickles, surface quite smooth and shining, dark glaucous green, sometimes mottled; flowering stem (scape) 90 cm high, oblique at the base, erect, stout, smooth, cylindrical from the center of the leaf-tuft; flowers red, stalked, numerous, erect in bud, afterwards pendulous, arranged in a rather close, narrow, erect, spicate, raceme, terminating the scape, bracts, membranous, triangular, acute, reddish, veined and persistent; perianth gamophyllous, tubular, about 3.7 cm long, rather fleshy, deeply cut into 6, oblong, bluntish segments; stamens 6, hypogynous, a little longer than the perianth; anthers small, oblong, dorsifixed near the base, bright orange turning brown; ovary superior, oblong-ovoid, bluntly triangular, 3-locular, with a double row of axile ovules in each chamber.

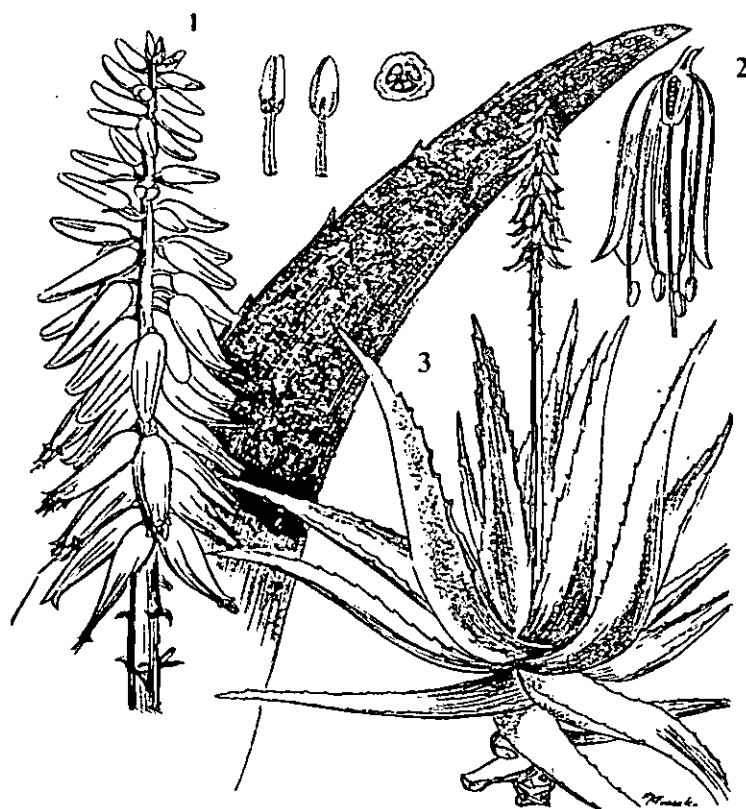


Fig - 1. *Aloe vera* plant

1. Inflorescence 2. Flower 3. Leaf

(Source- Jayaweera D. M. A., (1981). Medicinal plants used in Ceylon, Part III)

Plant Material of Interest

Leaves.

Official Drug

Parenchyma gel of leaves (Aloe gel, Aloe Vera), juice of leaves (exudate). The commercial drug is a hard dark mass (musabber) which is derived from yellow juice (dried juice of leaves)^{2,3}.

Pharmacognostic Features*

Anatomy

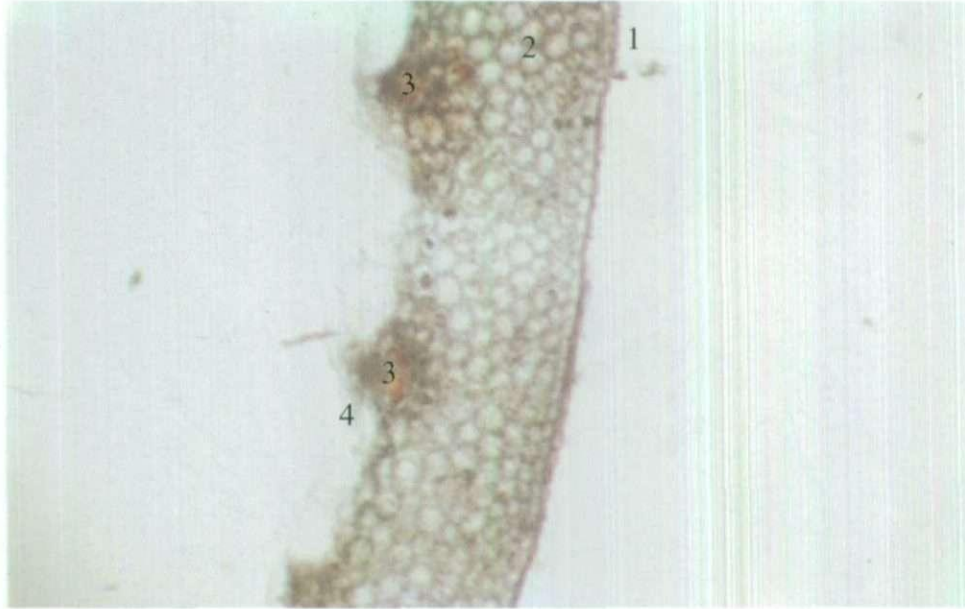


Fig - 2. Cross section of *Aloe vera* leaf (stained with safranin (10 x 10))

1. Epidermis 2. Parenchyma cells containing chloroplasts 3. Vascular bundle
4. Parenchyma (water storage tissue)

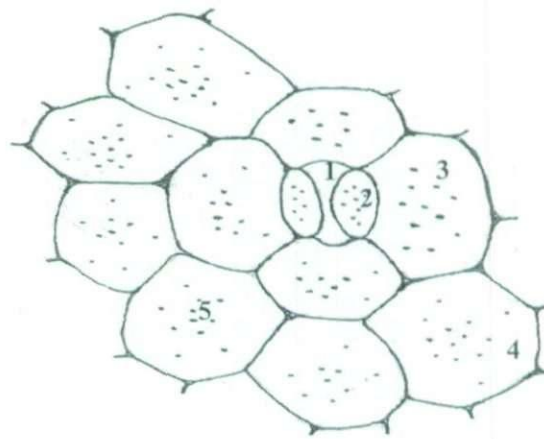


Fig - 3. Schematic diagram of epidermis of leaf

1. Stomatal pore 2. Guard cells 3. Subsidiary cells 4. Epidermal cells 5. Chloroplasts

Powder analysis

Part analyzed – Dried leaf

Organoleptic properties

Colour – Yellowish brown

Odour – Odourless

Taste – Slightly bitter taste

Microscopic characters

Parenchyma cells can be mainly seen. Stomata, Xylem vessels and fibers too can be seen.

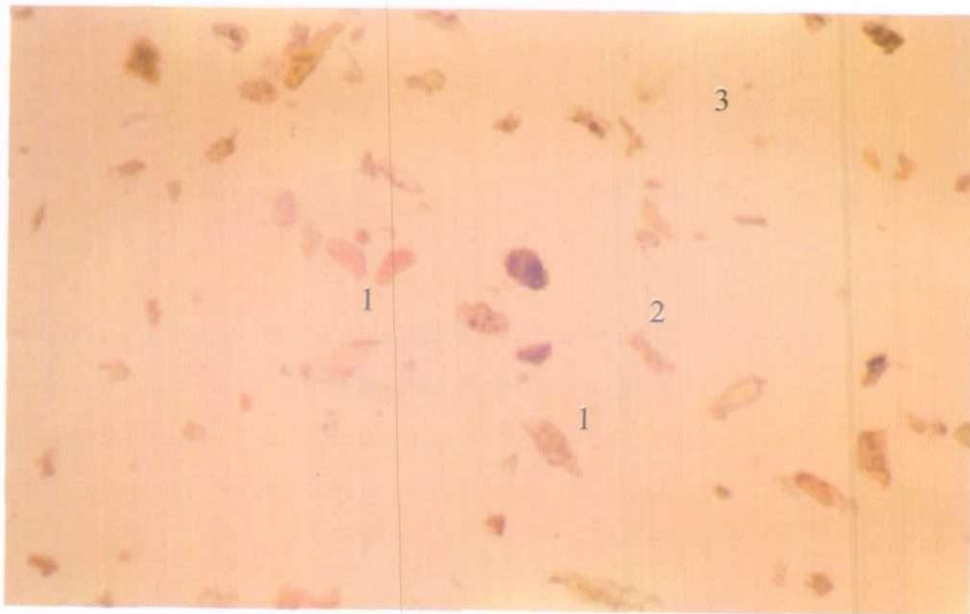


Fig – 4. Powdered plant material (leaf) under the microscope (10 x 4).
1. Parenchyma cells 2. Part of a vessel 3. Part of a fiber

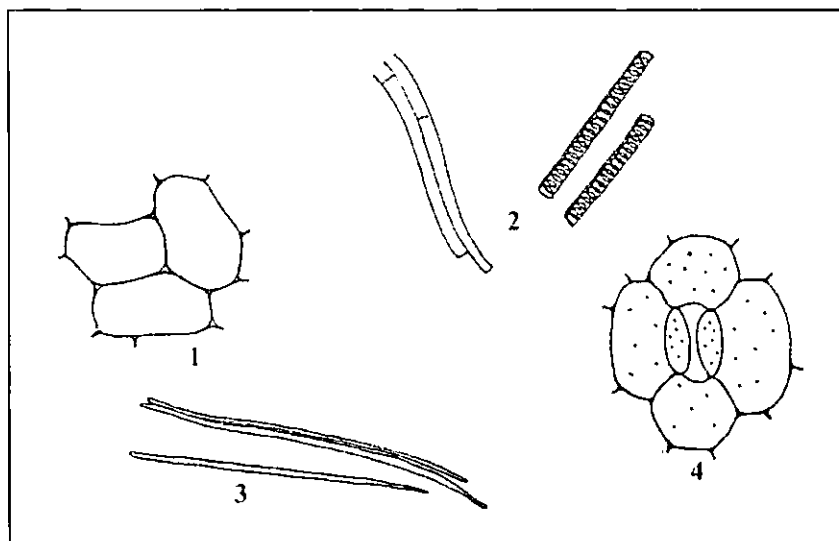


Fig - 5. Schematic diagram of powder microscopy

1. Parenchyma cells 2. Xylem vessels 3. Fibers 4. Stomata with subsidiary cells

Physico-chemical Analysis¹⁵⁶

Extractable matter

Crushed, dried plant material (about 4 g) was weighed to a glass-stoppered conical flask. Solvent (100 mL) was added, weighed, shaken well and allowed to stand for 1h. It was then boiled for 1h and cooled. The weight was readjusted with specified solvent and filtered. Filtrate (25 mL) was taken, solvent was evaporated and oven dried at 105 °C for 6h, cooled in a desiccator and weighed.

Total ash

Crushed, air dried plant material (about 4 g) was weighed to a previously ignited crucible. The material was ignited by gradually increasing the temperature to 550 °C until it was free from carbon. The crucible was cooled and weighed.

* These analysis were carried out by the authors at Industrial Technology Institute and the Dept. of Plant Sciences and Dept. of Zoology of University of Colombo.

Water soluble ash

Water (25 mL) was added to the crucible containing total ash, covered with a watch glass and boiled gently for 5 min. The insoluble matter was collected on an ashless filter paper and washed with hot water. The filter paper containing the insoluble matter was transferred to the original crucible and ignited for 15 min. at a temperature not exceeding 450 °C. Water soluble ash is the calculated difference in weight between the total ash and the residue remaining after treatment of the total ash with water.

Moisture content of the samples was estimated and all the calculations were done on dry weight basis.

Table 1. Physico-chemical parameters of *Aloe vera* leaf**

Physico-chemical parameter	Amount %
1. Water extractable matter	65.2 – 71.7
2. Ethanol extractable matter	51.4 – 55.3
3. Total ash	7.7 – 10.3
4. Water soluble ash	1.3 – 2.5

(Results are expressed as percentages on dry weight basis)

Thin Layer Chromatographic Profile**

Aloe vera water extract of leaves

Sample preparation : *A. vera* leaves (4 g) were boiled with water (100 mL) for 1h and the extract was filtered and evaporated to dryness. Eight microliters (8 μ L) of the diluted extract (65 mg in 3 mL) was spotted on TLC plate.

Absorbent : Silica gel GF₂₅₄

Solvent system : Ethyl acetate : Methanol : Chloroform (25 : 15 : 20)

Detection

Direct evaluation : UV₂₅₄ nm. R_f values - 0.24, 0.41

UV₃₆₆ nm. R_f values - 0.24, 0.41, 0.62, 0.74

Scanning : Densitometer at 254 nm (before spraying) and 450 nm (after spraying)

Spray reagent : Vanillin sulphate

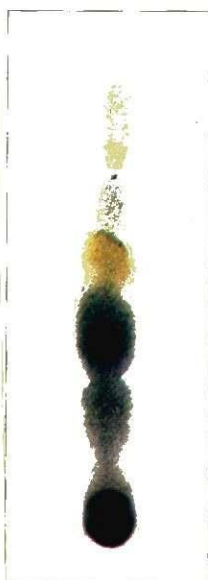


Fig - 4. TLC finger print profile of water extract of *Aloe vera* leaves

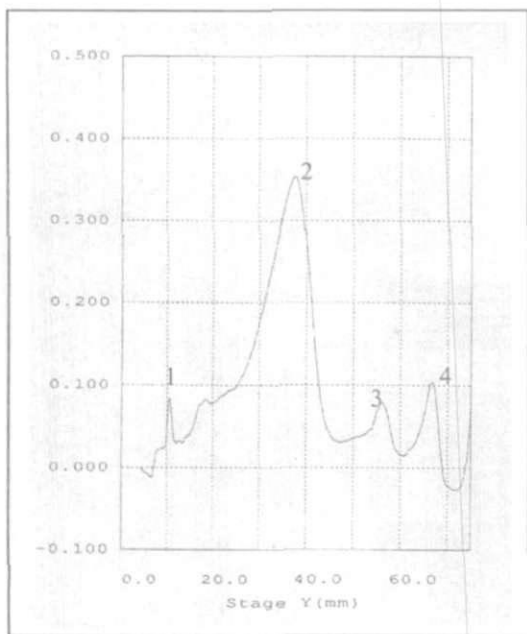


Fig-5. Densitogram of TLC finger print profile of water extract of *Aloe vera* leaves at 254 nm

Table 3. Description of densitogram (Fig-

6)

Peak no.	Y (mm)	Relative area %
1	10.54	4.08
2	18.36	6.09
3	37.65	69.61
4	65.06	11.95

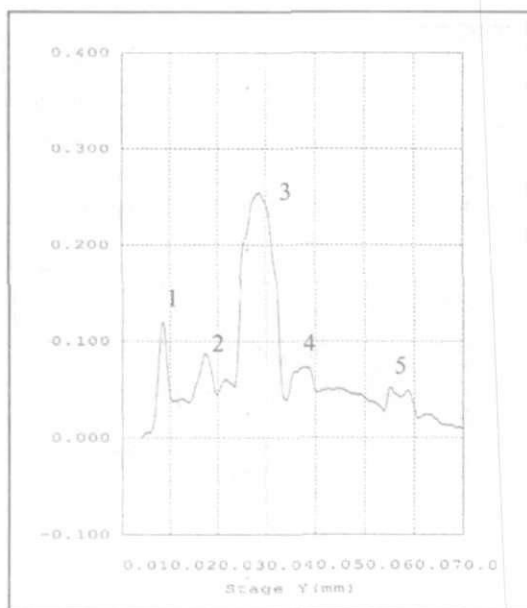


Fig - 6. Densitogram of TLC finger print profile of water extract of *Aloe vera* leaves at 450 nm

Table 3. Description of densitogram (Fig-6)

Peak no.	Y (mm)	Relative area %
1	8.52	7.12
2	12.50	8.64
3	17.46	47.08
4	28.66	9.16
5	55.38	5.84

***Aloe vera* ethanol extract of leaves**

Sample preparation : *A. vera* leaves (4 g) were boiled with 95% ethanol (100 mL) for 1h and the extract was filtered and evaporated to dryness. Eight microliters (8 μ L) of the diluted extract (67 mg in 3 mL) was spotted on TLC plate.

Absorbent : Silica gel GF₂₅₄

Solvent system : Ethyl acetate : Methanol : Chloroform (20 : 15 : 25)

Detection

Direct evaluation : UV 254 nm, R_f values - 0.15, 0.71, 0.95

UV 366 nm, R_f values - 0.15, 0.27, 0.42, 0.71, 0.95

Scanning : Densitometer at 254 nm (before spraying) and 450 nm (after spraying)

Spray reagent : Vanillin sulphate

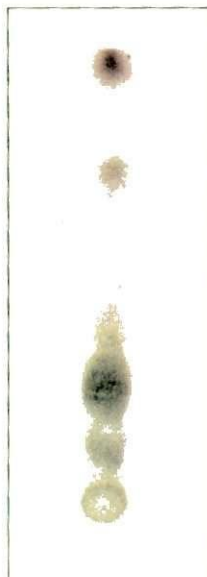


Fig – 7. TLC finger print profile of ethanol extract of *Aloe vera* leaves

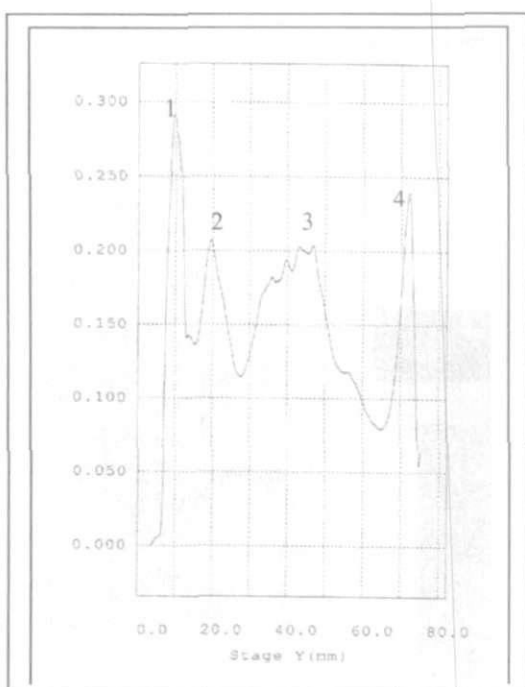


Table 4. Description of densitogram (Fig-8)

Peak no.	Y (mm)	Relative area %
1	9.75	24.61
2	19.66	24.55
3	43.13	41.53
4	72.10	20.25

Fig – 8. Densitogram of TLC finger print profile of ethanol extract of *Aloe vera* leaves at 254 nm

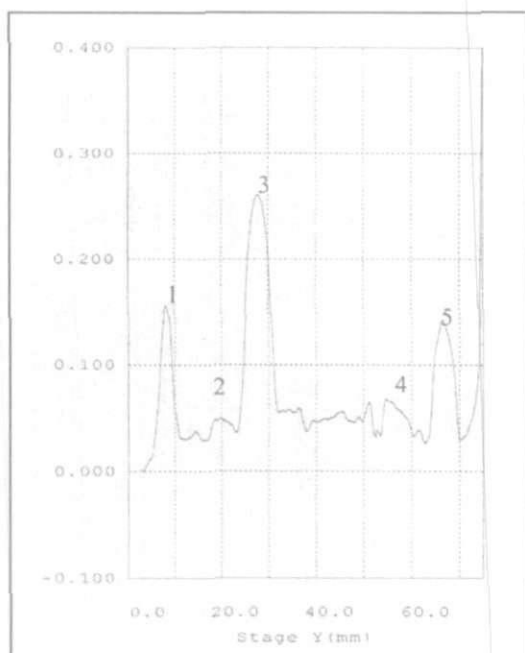


Table 5. Description of densitogram (Fig-9)

Peak no.	Y (mm)	Relative area %
1	8.16	14.95
2	19.87	6.54
3	27.76	38.7
4	55.11	6.54
5	66.45	12.91

Fig – 9. Densitogram of TLC finger print profile of ethanol extract of *Aloe vera* leaves at 450 nm

High Pressure Liquid Chromatographic Profile**

Aloe vera water extract of leaves

Sample preparation : *A. vera* leaves (4 g) were boiled with water (100 mL) for 1h and the extract was filtered and evaporated to dryness. The diluted extract (8 mg in 5 mL) was purified using Sep-pak C18 cartridge.

Injection volume : 20 μ L

Apparatus : Shimadzu LC – 10 ADvp pumps and Shimadzu SPD – M 10 Avp uv / vis photodiode array detector.

Column : Inertsil 5U ODS – 2 reverse phase column, (250 x 2.6 mm).

Solvent system : Acetonitrile : Water (30 : 70)

Flow rate : 1 mL/min

Detection : 254 nm

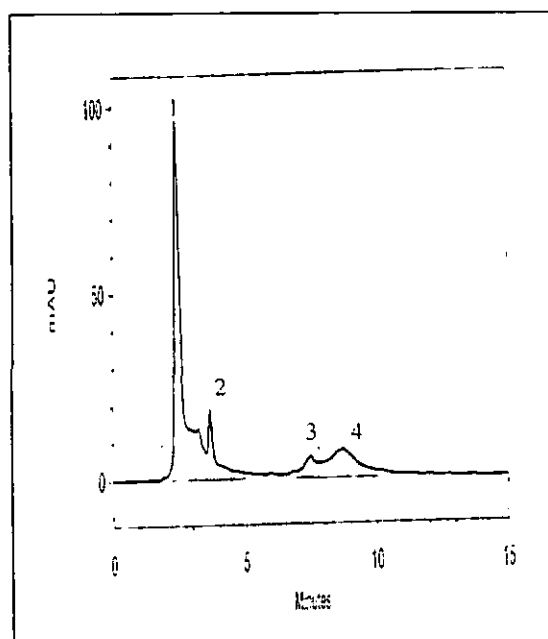


Table 6. Retention time of main peaks.

Peak no.	Retention time (min)	Relative area %
1	2.40	44.76
2	3.66	13.14
3	7.49	6.69
4	8.72	20.00

Fig – 10. HPLC finger print profile of water extract of *Aloe vera* leaves

Aloe vera ethanol extract of leaves

Sample preparation : *A. vera* leaves (4 g) were boiled with 95% ethanol (100 mL) for 1h and the extract was filtered and evaporated to dryness. The diluted extract (6 mg in 5 mL) was purified using Sep-pak C18 cartridge.

Injection volume : 20 μ L

Apparatus : Shimadzu LC – 10 ADvp pumps and Shimadzu SPD – M 10 Avp uv / vis photodiode array detector.

Column : Inertsil 5U ODS – 2 reverse phase column, (250 x 2.6 mm).

Solvent system : Methanol : Water (70 : 30)

Flow rate : 0.5 mL/min

Detection : 254 nm

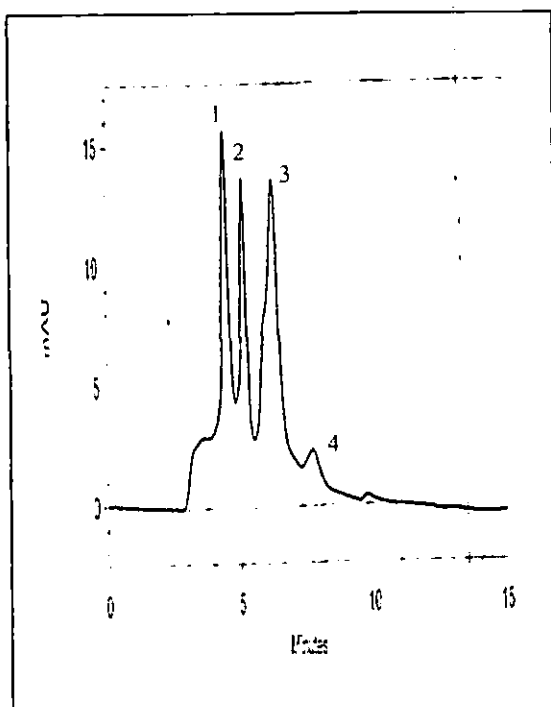


Fig – 11. HPLC finger print profile of ethanol extract of *Aloe vera* leaves

Table 7. Retention time of main peaks.

Peak no.	Retention time (min)	Relative area %
1	4.44	26.14
2	5.12	18.43
3	6.25	38.57
4	7.71	8.37

** These analysis were carried out by the authors at Industrial Technology Institute.

Phytochemistry

The major chemical constituents of *A. vera* are polysaccharides and water.

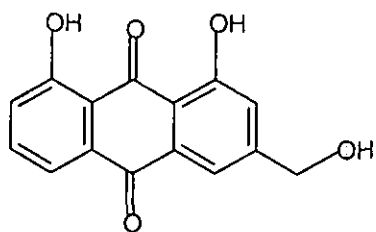
Following chemical constituents are also reported in *A. vera* plant. Iso-barbaloin and nataloin, homonataloin(7), α -cellulose, uronic acid, enzymes (oxidase, catalase, amylase, cellulase, proteinase), xylan, polyuronide, lignin, hydroxymethylanthraquinone, saponin, anthraquinone, polyphenols, triterpenoids, glucosamine, glucomannan, hexauronic acid, casanthranol I and II, aloetic acid, sapogenin, hecogenin, m-protocatechuic aldehyde, imidazole derivatives, erythritol, glycerol, hydrocinnamic acids, pectic acid, flavones, vitamins, minerals²².

Aloe leaf contains acemannan, aloe-emodin(1), aloesin, aloin, aloin A(5), aloin B(6), anthranol, monosaccharides (glucose, fructose, arabinose, mannose, galactose), barbaloin(2), amino acids (arginine, asparagine, histidine, leucine, isoleucine, threonine, valine, lysine), sterols (cholesterol(10), campesterol(11), β -sitosterol(12), lupeol(9)), β -carotene, chrysophanic acid, glutamic acid^{12,15,18,19,20,21}.

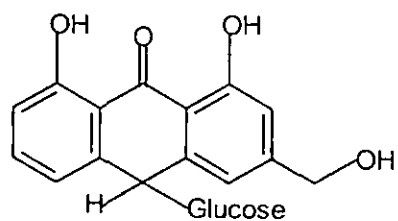
Leaf juice and the gel are parts of *A. vera* leaf. The following chemical constituents have been reported in these parts of leaf.

Aloe leaf juice contains aloe-emodin(1), anthranol, barbaloin(2), β -barbaloin, chrysophanol(4), p -coumaric acid(3), cystine, histidine, hydroxy proline polysaccharides^{11,12}.

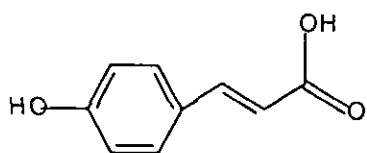
A. vera gel contains aloe-emodin(1), arachidic acid, arachidic acid methyl ester, arachidonic acid, behenic acid methyl ester, dehydro abetic acid methyl ester, 2(3H)benzothiazolone, cycloeicosane, cyclohexane, N-eicosane, N-docosane, N-heneicosane, freidooleanan-3-one, heneicosanoic acid methyl ester, lauric acid, linoleic acid, margaric acid, myristic acid, palmitic acid, palmitoleic acid methyl ester, pentadecanoic acid, stearic acid, tricosanoic acid methyl ester, N-octadecane, oleic acid methyl ester, oleic acid ethyl ester, uronic acid, enzymes (oxidase, catalase, amylase), galacturonic acid, monosaccharides (glucose, galactose, rhamnose, xylose, arabinose) lactate, salicylic acid, glucomannan, acetylated glucomannan and inorganic ions^{12,13,14,15,16,17}.



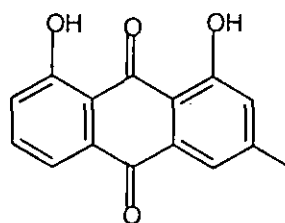
(1) Aloe-emodin



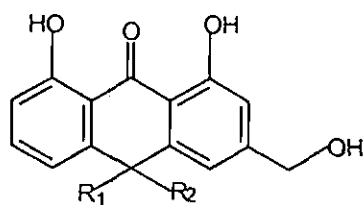
(2) Barbaloin



(3) *p*-Coumaric acid



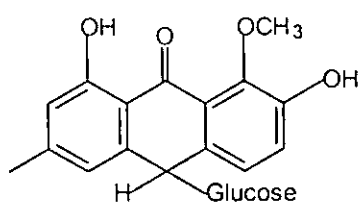
(4) Chrisophanol



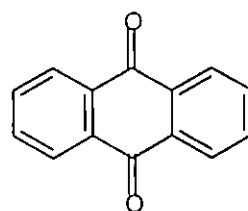
(5) Aloin A - $R_1 = H$, $R_2 = \text{Glucose}$

(6) Aloin B - $R_1 = \text{Glucose}$, $R_2 = H$

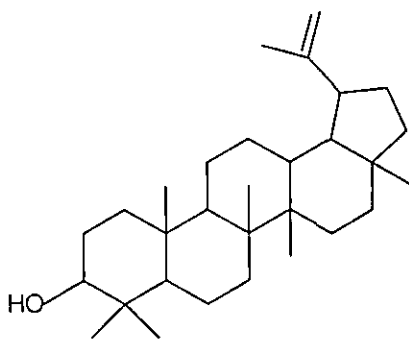
Fig – 12. Compounds present in *Aloe vera*



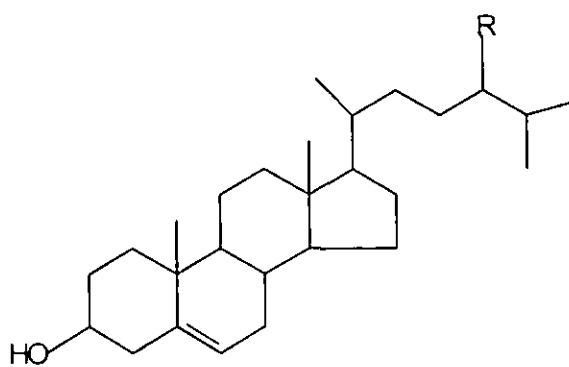
(7) Homonataloin



(8) Anthraquinone



(9) Lupeol



(10) Cholesterol R = H

(11) Campesterol R = CH₃

(12) β-Sitosterol R = CH₂-CH₃

Fig – 13. Compounds present in *Aloe vera*

Medicinal Uses

Uses described in pharmacopoeia and other traditional systems of medicine

According to the traditional systems of medicine *A. vera* has properties of alleviating the "Three Dosas"^{23,24,25,26}.

The plant is bitter, cooling^{3,5,23,24,27}, tonic^{5,27,28,29}, aphrodisiac^{5,27}, anthelmintic^{3,5,23,25}, alexiteric^{3,5,27}, fattening^{5,27}, aperient, ophthalmic³; useful in eye diseases^{1,3,5,23,30}, tumors^{5,23,25}, fever^{23,25,31}, fever due to bronchitis, vomiting, erysipelas, biliousness, leprosy⁵, skin diseases^{1,3,5,23,25}, asthma^{1,5}, jaundice^{5,23}, ulcers and pains^{5,23,24,25}. Aloe has diuretic properties and is useful in disuria and urinary disorders^{23,24,25,26}. It purifies blood^{23,24,25}, applied on wounds and good for painful inflammations^{23,24,25}.

A. vera has been used in diseases associated with the digestive system. It is purgative^{5,23,25,29,31,32,33,34}, carminative^{3,5,26,28} and laxative^{26,28,33,35,36}; used in dyspepsia^{1,3,5,23,37}, piles^{1,5,32,33} and rectal fissures^{32,33}; good for distention of abdomen and promotes downward movement of wind in stomach²⁶. It is said to improve appetite^{23,24,25}, promotes digestion^{24,226,28} and relieves constipation^{1,3,5,25,26,29,37}. It is given for enlargement of spleen and liver complaints^{3,5,23,24,25}. Aloe is emmenagogue^{3,5,28,29,32,37}. It produces pronounced pelvic congestion^{28,33} and used in menstrual suppressions and menstrual pains^{5,25,28,29}.

The fresh juice of leaves is cathartic^{1,5} and cooling^{28,29}, useful in fevers^{5,28}, treatment of abrasions, burns and skin irritations²⁸ and eye diseases^{1,5,25}.

The dried juice of *A. vera*, "bitter Aloe" or "mussabar" (titta abin in Sinhala) is a very useful drug in the Ayurvedic medicinal system. It is used in a large number of Ayurvedic preparations to treat tumors, spleen and liver troubles, flatus, dysurea, epilepsy, piles, jaundice, abdominal pain, inflammations, asthma, cough and uterine pain⁴. The dried juice is considered to be anthelmintic^{3,28}, cathartic^{28,29} (acting on large intestine), purgative^{3,32}, emmenagogue^{3,28,32}, stomachic^{28,32} and vermifuge²⁸.

The cooling mucilage is used as a poultice on inflammations caused by x-ray and other radiation burns. The pulp of leaves is said to possess wound healing activity and is used for intestinal ailments, sore throat and ulcers²⁸.

The root of the plant is given for colics²⁹. A decoction of the root is given for stomach cancer²⁸.

The flowers are anthelmintic, useful in biliousness and "vata"⁵.

Uses in folk medicine

Fresh Aloe leaf gel is applied on burns^{23,24,28}. It is kept on closed eyes for headaches with under eye pain²³. One teaspoonful of Aloe leaf gel at a time is given to dissolve blood clots²³. The ground mixture of Aloe leaf gel and green gram is eaten with jaggery to soothe the burning sensation in stomach²⁴. A mixture of Aloe leaf gel and passion fruit is applied on hair for dandruff²⁴.

A drink made of leaf pulp of Aloe and *Genipa americana* is a popular remedy for colds in USA³⁸. Leaf pulp is a refrigerant medicine in conjunctivitis, with small quantity of sugar-candy. Leaf pulp with addition of burnt alum is being used in ophthalmia. Tender pulp is eaten for rheumatism and also in menstrual suppression and piles²⁷.

Fresh juice has been used in fevers, liver problems, gonorrhea, methritis and external refrigerant application for local inflammations²⁷. Filtered juice of the mixture of *Terminalia chebula* seed powder and Aloe leaf gel have been put into the eye to prevent tearing²⁴. A plaster of the leaf or the leaf juice is said to be a folk remedy for tumors²⁸. The throat is washed with Aloe juice for sore throat²³. Leaf juice mixed with gingerly oil and boiled makes a fine hair-oil used in cases of sleeplessness. It is mixed with opium and applied to forehead to relieve headache²⁷. Turmeric with leaf juice is given in glandular enlargement and spleen diseases²⁷.

A decoction of the root is said to be a remedy for stomach cancer²⁸.

When sharp thorns pierce the skin, turmeric and scraped coconut are ground together, placed in the middle of the Aloe leaf, kept on warm charcoal and wrapped around the area²³.

A mixture of *Zingiber officinale*, *Mentha pulegium* (essential oil), *A. vera*, *Ipomoea purga*, *Glycyrrhiza glabra* and *Centella alba* is used for amenorrhea³⁹.

The water extract of Aloe is used against worms⁴⁰.

The plant is used as a contraceptive and as an abortifacient^{41,42,43,44}.

Tender leaves of Aloe mixed with cumin seed and sugar-candy is a remedy for dysentery with bloody stools²⁷.

In USSR, Aloe juice, Aloe emulsion and Aloe syrup with iron are in use. Aloe juice is used for treating burns, suppurative wounds and tropic ulcers. The thick syrup of Aloe with iron is effective against anaemia and the pulp of the leaves is used for bruises, sunburns, boils and carbuncles³⁷.

In Philippines the juice is applied to prevent the falling of hair and to cure baldness¹.

In Chinese medicine *A. vera* is used against intestinal parasites, as a laxative and to facilitate digestion. It is also used in cases of chronic constipation and amenorrhea, liver and eye troubles, against piles, infantile intestinal tuberculosis, convulsions, and epilepsy²⁶.

In western countries *A. vera* is used as a purgative, anti-gripe medicine in connection with other carminatives, laxative, to heal burns and wounds, cuts, sunburn, as a digestive, for hair and scalp care, skin care, varicose veins, psoriasis and eczematous rashes and as cleanser²⁶.

Aloe is also used in veterinary medicine²³.

Other Uses

A. vera gel is extensively used in shampoos, face creams, lotions, cleansers and ointments^{10,27,37}.

It is used as an ingredient in most major categories of food products including alcoholic beverages (bitters, liqueurs, vermouths) and nonalcoholic beverages, frozen dairy desserts, candy, baked goods, gelatins and puddings²⁸.

Bottled *A. vera* is available in the USA for internal consumption as a tonic¹⁰.

The plant is used as a repellent to insect pests (as a white ant repellent)⁴⁵.

Aloe leaf is cooked as a vegetable²³.

In home gardens it is grown as an ornamental plant.

Products Developed From *Aloe vera*

Cosmetics and toiletries - Facial moisturizers, Hand and body lotions, Cleansing agents, After bath moisturizers, Make-up removers, Suncare products, Shaving preparations, Hair care products, Mascara and eyelash conditioning products⁴⁶.

First aid applications - Topical analgesic / anaesthetic preparations, Anti-acne gel, Bedsore preparations, Antiviral / antibacterial topicals, Lip gel, Wet dressings⁴⁶.

Other products- *A. vera* gel, *A. vera* powder, oil extract, Shealoe (*A. vera* gel and Shea butter)⁴².

Products developed at Industrial Technology Institute

Cosmetics and toiletries - Shampoo, Body gel, Hand wash, Aloe moisturizer⁴⁷.

First aid applications - Burn creams⁴⁷

Food and beverages - Aloe jelly, Aloe drink⁴⁷.

Ayurvedic / Traditional Medicinal Preparations

A. vera is used as a constituent of the following Ayurvedic preparations.

Prameha gajankushaya, Ranahansa rasayana, Kumaryasava, Kumari taila, Nayanabharana tila, netrapatala kantiratna tila, Batu tila, bodhisatva tila, Svasavajra tila, kumarika vati, Rajapratvatani vati, Kumari paka, Chandrashura rasa, Pradarantaka rasa, Brihadchandrodaya makaradhwaja (Purna chandrodaya rasa), Hemagarbhapottali rasa, Kumarivati^{3,23,48}.

Activity Studies

Anti-inflammatory activity

Anti-inflammatory activity of *A. vera* has been investigated by a number of in vitro and *in vivo* studies^{49,50,51}. Continuing research into inflammation has shown that it is a complex process involving many biochemical pathways and a variety of agents and mediators. In particular three components are distinguished.

1. Vasoactive substances; agents causing dilation of blood vessels and opening of junctions between cells of the ultimate capillaries, produced by altering contractile elements in endothelial cells. These factors include vasoactive amines, bradykinin and also prostaglandins.
2. Chemotactic factors; these agents cause increased cell motility, especially of white blood cells (leucocytes) into stressed areas. These factors include several proteins and enzymes.
3. Degradative enzymes; these are hydrolytic enzymes breaking down tissue components.

It was also shown that Aloe gel contained both an inhibitory system and a stimulatory system that influenced both inflammatory and immune responses^{50,51,53,54}.

The gel relieved the inflammatory effect induced by carrageenan, albumin, gelatin, mustard and croton oil either by promoting prostaglandin synthesis or by increasing infiltration of leucocytes⁵¹. Some studies have shown that the gel exerts its anti-inflammatory activity through bradykinase activity⁵² and thromboxane B₂ and prostaglandin F₂ inhibition¹⁴. Some experiments have shown anti-lipoxygenase activity of *A. vera* extracts. This finding could explain the inhibitory effect of the *A. vera* in acute inflammation process⁵⁶. Three plant sterol compounds in *A. vera* gel reduced inflammation up to 37% in croton oil induced oedema in mice⁵⁷. The externally used aqueous extract of Aloe has shown anti-inflammatory effects^{58,59,60,61,62}.

Some experiments have shown *A. vera* extract has decreased inflammation due to adjuvant arthritis⁵².

Wound healing and burn treatment

A. vera gel has beneficial effects on wound healing^{49,60,63,64}. It has been found that externally used Aloe leaf juice can accelerate the wound healing process^{65,66,67}. Various mechanisms have been proposed for the alleged healing properties of *A. vera* gel. It is

possible that the gel may have more than one physiological effect. It is commonly suggested that there is a synergistic effect between the various components and the polysaccharide base¹⁰. Angiogenesis is an essential process in wound healing and some studies suggest that the gel contains angiogenic compounds⁶⁸. It is found that therapeutic effect of the gel includes the prevention of dermal injuries caused by burns, frostbite, electric injury and inadvertent arterial drug abuse⁶⁹. *In vitro* studies have demonstrated that *A. vera* gel promotes wound healing by directly stimulating the activity of microphages and fibroblasts⁷⁰. Increased collagen formation in wounded diabetes rats treated orally and topically with Aloe gel was demonstrated⁵². Healing and control of acute inflammation, distinct from chronic inflammation, was observed following gel treatment of excision and incision wounds in rats⁵². Acemannan, a complex carbohydrate isolated from Aloe leaves, has been shown to accelerate wound healing and reduce radiation induced skin reactions⁷¹. Acemannan can activate macrophages and macrophages play a considerable part in controlling microorganisms. It was shown that young active macrophages accelerated the rate of wound healing in aged rats⁵². A carbohydrate fraction of *A. vera* showed haematopoietic activity⁷².

Effects of *A. vera* on wounds caused by thermal burns and radiation burns have been studied^{52,73,74,75,76}. It was found that *A. vera* could inhibit the inflammatory processes following burn injury, as characterized by the reduction of leukocyte adhesion, as well as pro-inflammatory cytokines⁷⁷. *A. vera* gel on burn wounds in guinea pigs have shown promising effects of healing⁷⁸.

Antioxidant activity

Antioxidant potential of *A. vera* has been evaluated and it was suggested that the growth stage play a vital role in the composition and the antioxidant activity of *A. vera*⁷⁹. Studies have shown that the synergistic relation between the elements found in the leaf of *A. vera* could be a useful adjunct for maintaining the integrity of the antioxidant status⁸⁰. It was found that the supplementation of *A. vera* extract has preventive effects in attenuating oxidative damage in brain regions of streptozotocin induced diabetes rats via antioxidative mechanism⁸¹.

Anti-tumor activity

Anti-tumor activity of *A. vera* was observed⁵² and it was shown that Aloe juice⁸², the ethanol extract⁸³ and the polysaccharides isolated from Aloe^{84,85} possess anti-tumor activity. Studies have shown that Aloe-emodin has anti-tumor activity⁸⁶.

Anti-diabetes activity

Effect of Aloe gel on blood glucose level of diabetes patients has been investigated. Administration to non-insulin dependent human patients in small trials resulted in a sustained lowering of blood glucose level⁵². *A. vera* leaf gel and pulp extract showed the protective effect on mild damage on kidney tissues⁸⁷ and hepatotoxicity of liver⁸⁸ caused by type-II diabetes in rats. Studies have confirmed the ethnopharmacological use of *A. vera* in ameliorating the oxidative stress found in diabetic rats⁸⁹. The supplementation of *A. vera* showed the preventive effect in attenuating oxidative damage in brain regions of streptozotocin induced diabetic rats⁸¹. Hypoglycemic activity was studied in alcoholic extract, aqueous extract and the juice of Aloe^{90,91,92}. Anti-hyperglycaemic activity was also studied in Aloe sap and aqueous extract^{93,94}.

Radioprotective effects

Aloe polysaccharides showed radioprotective effects on non-tumor cells⁹⁵. Aloe polysaccharides exert radioprotective effects *in vitro* and *in vivo* through inhibition of apoptosis⁹⁶.

Effects on immune system

Immunomodulatory activity of bitter yellow juice of *A. vera* has been studied in mice bearing murine Lymphoma. In this study, restoration of immunological parameters in cellular immune response and phagocytosis and the stimulation of specific and non specific responses were observed⁹⁷. Immunomodulatory activity has been also observed in aqueous extract of *A. vera* gel⁹⁸. Externally used lyophilized extract of *A. vera* has shown immunostimulant activity⁹⁹. Polysaccharide fraction and Aloe aqueous extract have shown anticomplement activity^{100,101,102}. It has also acted as an immunological adjuvant on specific antibody production¹⁰² and enhanced the release of cytokines¹⁵⁹. Release of these cytokines stimulates the replication of fibroblast in tissue culture and enhances macrophage phagocytosis¹⁵⁹.

Effects on gastrointestinal function and ulcers

Dietary supplement of *A. vera* can suppress age-related increases in hepatic cholesterol¹⁰³. Aloe gel is effective in the treatment of peptic ulcers and has protective effects on gastric mucosa^{52,104,105}. *In vitro* studies provided support that it may have a therapeutic effect in inflammatory bowel disease¹⁰⁶. Preadministration of a water extract of whole *A. vera* leaves to rats, reversed the inhibition by blood ethanol of alcohol dehydrogenase and aldehyde dehydrogenase activities. It also reversed the increase of lactate/pyruvate ratio which could decrease NAD supply⁵².

Anti-microbial activity

Anti-bacterial activity of *A. vera* was observed against *Proteus vulgaris* and *Pseudomonas aeruginosa*⁸². The fresh juice showed activity against *Corynebacterium xerosis*, *Bacillus subtilis*, *Enterobacter* species, *Escherichia coli*, *Serratia marcescens*, *Staphylococcus aureus* and *Streptococcus pyogenes*^{107,108,109}.

Anti-viral activity against Distortion ringspot virus, Mild mosaic virus and Ringspot virus was observed in ethanol extract of *A. vera*¹¹⁰. Methanol extract of *A. vera* leaf showed antiviral activity against Herpes simplex 1 and 2 virus¹¹¹. Acemannan found in *A. vera* gel has shown to inhibit AIDS virus replication *in vitro* and an injectable form has been found to be of significant benefit in FIV-infected cats^{160,161}.

Fresh juice of leaves has shown anti-fungal activity against *Candida albicans* (yeast species)¹⁰⁸.

Effects on fertility and pregnancy

Ethanol and petether extract of Aloe showed anti-implantation activity when orally administered to female mice¹¹². Embryotoxic effects on pregnant rats have been studied in water extract, ethanol extract, benzene extract and ethanol : water (1:1) extract^{113,114}. Some studies have shown Aloe water extract and tincture possess uterine stimulant effects^{115,116}. Teratogenic effects were also studied in Aloe water extract^{43,117}.

Hypocholesterolemic activity

Water extract of *A. vera* has shown hypocholesterolemic activity in rats⁹¹. Studies have suggested that dietary Aloe supplementation suppresses free radical induced oxidative damage and age related increase in hepatic cholesterol¹⁰³.

Analgesic activity

Analgesic activity of *A. vera* was observed in ethanol and water extracts in mice^{61,118,119}. Externally used leaf pulp of Aloe has shown analgesic activity in human adults⁶⁴.

Stimulation effects

Fresh leaf juice of Aloe has the effect of stimulating the growth of hair⁶⁵. Tincture of Aloe possesses smooth muscle stimulant activity in isolated rat ileum¹²⁰.

Plant growth inhibition

Water extract of leaf and dried stem of Aloe possess plant germination inhibition effects¹²¹.

Anti-pyretic activity

Ethanol extract of Aloe has shown antipyretic effect in mouse¹¹⁸

The use of *A. vera* gel has also been described in veterinary medicine. It is also used to treat a variety of dental conditions¹⁰.

Safety Evaluation

Plant extract at oral doses of 100 and 200 mg/kg did not show any toxicity in rats. Prolonged use may severely affect the electrolyte balance and loss of potassium ultimately may reduce the laxative action and disturb cardiac rhythm in heart patients. Larger doses lead to accumulation of blood in the pelvic region and reflux stimulation of uterine muscles may bring about abortion or premature birth in late pregnancy. Toxic doses may also cause kidney damage. Active principles generally appear in milk during lactation. Due to these reasons the drug is contraindicated in pregnancy, lactation, kidney complaints and irritable bowel conditions. LD₅₀ in mice by *i.v.*, *i.p.* and oral routes were found to be 400 mg/kg, 800 mg/kg and 1600 mg/kg respectively³.

A. vera has protective effects against some toxicity effects of arsenic in rats¹²²

Patent Information

- Skin adhesive cosmetic for removing wrinkles, containing vitamins and Aloe extract¹²³
- Improved vitamin A palmitate composition for topical application to the skin¹²⁴
- Chlorine dioxide skin medicating compositions for preventing irritation¹²⁵
- Skin conditioner¹²⁶
- Skin enhancing glove containing *A. vera*¹²⁷
- Post hair removal skin care lotion¹²⁸
- Skin cosmetic¹²⁹¹³⁰
- Novel skin drug for external use¹³¹
- Skin and mucosal treatment formulation¹³²
- Anti-inflammatory and wound-healing topical skin preparations containing Aloe extract and ellagic acids¹³³
- Hair regenerator based on Aloe extract¹³⁴
- Aloe gel possessing anaesthetic activity¹³⁵
- Stabilized *A. vera* gel¹³⁶¹³⁷
- Analgesic gel composition for temporary relief from symptoms of arthritis¹³⁸
- Lip balm composition¹³⁹
- Medicated gel¹⁴⁰
- Pad for applying medicaments¹⁴¹
- Topical agent and method for the treatment of pseudofolliculitis barbae¹⁴²
- Aloe water preparation. (can be used in cosmetics for treating skin and hair)¹⁴³
- *A. vera* juice containing pellets and the process for preparation of pellets¹⁴⁴
- Hypoglycemic polysaccharides extraction from Aloe species¹⁴⁵
- Hand cleanser¹⁴⁶
- Wet sanitary pad for cleansing¹⁴⁷
- Detergents and soaps containing Aloe and herbs¹⁴⁸
- Shaving soap and aftershave gel¹⁴⁹¹⁵⁰
- Decongestant comprising zinc and vegetable oil¹⁵¹
- *A. vera* extracts¹⁵²
- Transdermal delivery system¹⁵³
- Preparation for regulating bowel function¹⁵⁴

- Antibacterial composition and method of using it¹⁵⁵
- Fibroblast proliferation promoting agent¹⁵⁶
- Separation of bitter components in Aloe extracts¹⁵⁷
- *A. vera* processed leather¹⁵⁸

Cultivation

Aloe vera Linn. Var. *littoralis* Koen. is the Aloe species found in Sri Lanka. *A. vera* *littoralis* grows on the poorest soil, has low water requirements and a tolerance to salinity. It can be grown satisfactorily on the seacoast of the dry zone with little agricultural input although financial return from it will be relatively low in comparison to other medicinal plants.

Propagation is carried out through root suckers. Plants may be placed in rows 1 meter apart, each row 1 meter broad containing 3 plants abreast, each row being 30 cm apart from the next row. This arrangement will facilitate harvesting. The preparation of soil need not exceed 20 cm in depth as the root system of this plant is very shallow. Intercultures may be required from time to time. Water should not be allowed to lodge near the plant as this plant is not water loving and rotting takes place easily when waterlogged.

The plant is ready for harvesting in one year, although best yields are obtained during the third year. Harvesting is carried out by lopping off the leaves of the mature plant.

The crude "Aloes" are derived from the juice of the plant. The juice is obtained when the leaf is cut transversely close to the stem and held vertically. The yellowish juice flows out from the pericycle associated with vascular bundles. This juice is very bitter in taste and has an unpleasant odour. The residual Aloe leaf may be used for the preparation of gel (Aloe Vera) by slitting the leaf longitudinally and removing the gel from the green outer covering of the leaf³².

References

1. Jayaweera, D.M.A. (1981). *Medicinal plants (Indigenous and Exotic) used in Ceylon*, National Science Council of Sri Lanka. Part III.
2. *WHO monographs on selected medicinal plants*, (1999). World Health Organization, Geneva. Vol. 1.
3. Sharma, P.C., Yelne, M.B. and Dennis, T.J. (2001). *Database on medicinal plants used in Ayurveda*, Central Council for Research in Ayurveda and Siddha, New Delhi. Vol. 1.
4. *Ayurveda Pharmacopoeia*, (1980). Department of Ayurveda, Colombo, Sri Lanka. Vol. 1, Part 2.
5. Kirtikar, K.R. and Basu, B.D. (1935). *Indian Medicinal Plants*, International Book Distributors, Booksellers and Publishers. Vol. IV, 2nd edition.
6. *British Pharmacopoeia*, (1993). Welsh office, Department of Health and Social Services for Northern Ireland. Vol. 1.
7. *The Pharmacopoeia of the United State of America*, (1960). Authority of the United States Pharmacopoeial Convention, Inc. 16th revision.
8. *European Pharmacopoeia*, (1997). The European Department of the quality of medicines within the Council of Europe, Strasbourg. 3rd edition.
9. Bose, K.C. (1984). *Pharmacopoeia Indica*. The Book Company Limited, College Square, Calcutta.
10. Grinlay, D. and Reynolds, T. (1986). The *Aloe vera* phenomenon; A review of the properties and modern uses of the leaf parenchyma gel. *Journal of Ethnopharmacology* 16: 117-151.
11. Mukerji, B. (1953). *The Indian Pharmaceutical Codex*. Indigenous drugs, Council of Scientific and Industrial Research, New Delhi., India. Vol. 1.
12. Waller, G.R., Magiafico, S. and Ritchey, C.R. (1978). A chemical investigation of *Aloe barbadensis*. *Proceedings of Oklahoma Academy of Science* 58:59.
13. Gowda, D.C., Neelisiddaiah, B. and Anjaneyalu, Y.V. (1979). Structural studies of polysaccharides from *Aloe vera*. *Carbohydrate Research* 72: 201-205.
14. Robson, M.C., Heggors, J. and Hagsrom, W.J. (1982). Myth, magic, witchcraft or fact? *Aloe revisited*. *Journal of Burn care and Rehabilitation* 3: 157-162.
15. Rowe, T.D. and Parks, L.M. (1941). A phytochemical study of *Aloe vera* leaf. *Journal of American Pharmaceutical Association (Science edition)* 30: 262-266.

16. Mandal, G. and Das, A. (1980a). Structure of the glucomannan isolated from the leaves of *Aloe barbadensis* Miller. *Carbohydrate research* 87: 249-256.
17. Yamaguchi, I., Mega, N. and Snada, H. (1993). Components of the gel of *Aloe vera* (L.) Burm. F. *Bioscience Biotechnology and Biochemistry* 57(8): 1350-1352.
18. Holdsworth, D.K. (1971). Chromones in Aloe species, part I, Aloisin A, C-Glucosyl-7-hydroxychromone. *Planta Medica* 19: 322-325.
19. Mary, N.Y., Christensen, B.V. and Beal, J.L. (1956). A paper chromatographic study of Aloe, Aloin and Cascara sagrada. *Journal of American Pharmaceutical Association (Science edition)* 45: 229-232.
20. Rauwald, H.W. (1987). New hydroxylations: the 'periodate-positive' substance from Cape Aloes and cinna oil esters from Curacoa Aloes. *Pharmacy Weekly (Scienced)* 9(4): 215.
21. Groom, Q.J. and Renolds, T. (1987). Barbalion in Aloe species. *Planta Medica* 53(4): 345-348.
22. *Pakistan Encyclopaedia Planta Medica*, (1986). A joint research project of Hamdard Foundation Pakistan and H. E. J. Research Institute of Chemistry. Hamdard Foundation Press, Hamdard centre, Pakistan. Vol. 2.
23. *Compendium of Medicinal plants, A Sri Lankan Study*, (2002). Department of Ayurveda. Vol. II.
24. Perera D.L. (Ed.), (2003). *Osuthuru Wagathuga*. Sri Lanka Conservation and Sustainable Use of Medicinal Plants, No.4, Woodland Avenue, Kohuwala.
25. *Osuthuru Visithuru*, (1994). Department of Ayurveda. Part III.
26. Bhagwan Desh and Junius, A.M.M. (1983). *Hand Book of Ayurveda*. New Delhi, India.
27. Agharkar, S.P. (1991). *Medicinal plants of Bombay presidency*, Scientific Publishers, 5, New Pali Rd, Jodhpur.
28. Duke, J.A. (1987). *CRC Handbook of Medicinal Plants*, CRC press, Boca Raton, Florida.
29. *Medicinal plants of India*, (1976); Indian Council of Medical Research, New Delhi. Vol. 1.
30. *Thalpatha Osumahima*, (2002). Department of Ayurveda, Bandaranaike Memorial Ayurveda Research Institute. Vol. III.
31. Shah, A.H., et al. (1989). Toxicity studies on six plants used in the traditional Arab system of medicine. *Phytotherapy Research* 3(1): 25-29.

32. Chopra, R.N., *et al.* (1982). *Chopras Indigenous Drugs of India*, Academic Publishers. 2nd edition.
33. *The Wealth of India, Raw materials*, (1948). Publication and Information Directorate, Council of Scientific and Industrial Research, New Delhi. Vol. 1.
34. *Medicinal plants in the South Pacific*. (1998). WHO regional publications, Western Pacific series, No. 19.
35. Mukerji, B. (1953). *The Indian Pharmaceutical Codex*, Indigenous drugs, Council of Scientific and Industrial Research, New Delhi, India. Vol. 1.
36. Bastien, J.W. (1983). Pharmacopoea of Qollahuaya Andeans. *Journal of Ethnopharmacology* 8(1): 97-111.
37. *The Wealth of India. Raw Materials*, (1985). Publication and Information Directorate, Council of Scientific and Industrial Research, New Delhi. Revised. Vol. 1: A.
38. Elvin-Lewis, M. and Lewis, W.H. (1983). The dental use of plants in Amazonia. *Odonto-Stomatologie Tropicale* 6(4): 179-186.
39. Anon, (1912). *More secret remedies, What they cost and what they contain*. British Medicinal Association, London.
40. Weninger, B., Haag-Berrurier, M. and Anton, R. (1982). Plants of Haiti used as antifertility agents. *Journal of Ethnopharmacology* 6: 67-84.
41. Woo, W.S., *et al.* (1981). A Review of research on plants for fertility regulation in Korea. *Korean Journal of Pharmacognosy* 12(3): 153-170.
42. Wong, W. (1976). Some folk medicinal plants from Trinidad. *Economic Botany* 30: 103-142.
43. El-Dean Mahmoud, A.A.G. (1972). *Study of indigenous (folkways) birth control methods in Alexandria*. Thesis MS, University of Alexandria, Higher Institute of Nursing.
44. Nath, D., *et al.* (1992). Commonly used Indian abortifacient plants with special reference to their teratologic effects in rats. *Journal of Ethnopharmacology* 36(2): 147- 154.
45. Jansz, E.R., Wijeyaratne, R., Silva, V. and Ratnayake, D., *Series on Sri Lankan Medicinal Plants of Economic value*, Booklet on. 3, Aloe. Section of Natural Products, Ceylon Institute of Scientific and Industrial Research, Colombo.
46. *Technical Bulletin*, Terry Laboratories, 390N. Wickham Road, Suite F, Melbourne, Florida, USA.

47. Industrial Technology Institute (CISIR), 363, Bhauddhaloka Mawatha, Colombo 7, Sri Lanka.
48. *Ayurveda Pharmacopoeia*, (1976). Department of Ayurveda, Colombo, Sri Lanka. Vol. 1, Part 1.
49. Davis, R.H., *et al.* (1994). Anti-inflammatory and wound healing activity of growth substances in *Aloe vera*. *Journal of American Paediatric Medical Association* **84**(2): 77-81.
50. Davis, R.H., *et al.* (1991). Isolation of a stimulatory system in an Aloe extract. *Journal of American Paediatric Medical Association* **81**(9): 473-478.
51. Davis, R.H., *et al.* (1991). The isolation of an active inhibitory system from an extract of *Aloe vera*. *Journal of American Paediatric Medical Association* **81**(5): 258-261.
52. Reynolds, T. and Dweck, A.C. (1999). *Aloe vera* leaf gel: a review update. *Journal of Ethnopharmacology* **68**: 3-37.
53. Fijita, K. and Teradaira, R. (1976). Bradykininase activity of Aloe extract. *Biochemical Pharmacology* **25**: 205.
54. Hegggers, J.P. and Robson, M.C. (1983). Prostaglandins and thromboxanes. In: Ninnemann. J.L. (Ed.), *Traumatic injury. Infection and Other Immunological Sequelae*. University Park Press, Baltimore.
55. Raine, T.J., *et al.* (1980). Anti- prostaglandins and anti-thromboxanes for treatment of frostbite. *American College of Surgeons Surgical Forum* **31**: 557-559.
56. Benzacova, L., *et al.* (2005). Antilipoxygenase activity and the trace element content of *Aloe vera* in relation to the therapeutical effect. *Ceskoslovenska Farmacie* **54**(1): 43-46.
57. Davis, R.H., *et al.* (1994). *Aloe vera*, hydrocortisone and sterol influence on wound tensile strength and anti-inflammation. *Journal of the American Paediatric Medical Association* **84**: 614-621.
58. Thompson, J.E. (1991). Topical use of *Aloe vera* derived allantion gel in otolaryngology. *Ear Nose and Throat Journal* **70**(2): 119.
59. Davis, R.H., Leitner, M.G. and Russo, J.M. (1987). Topical anti-inflammatory activity of *Aloe vera* as measured by ear swelling. *Journal of American Paediatric Medical Association* **77**(11): 610-611.
60. Davis, R.H., Kabbani, J.M. and Maro, N.P. (1987). *Aloe vera* and wound healing. *Journal of American Paediatric Medical Association* **77**(4): 165-169.

61. Davis, R.H., Leitner, M.G. and Russo, J.M. (1988). *Aloe vera*, a natural approach for treating wounds, edema and pain in diabetes. *Journal of American Paediatric Medical Association* **78(2)**: 60-68.
62. Davis, R.H. and Maro, N.P. (1989). *Aloe vera* and gibberellin anti-inflammatory activity in diabetes. *Journal of American Paediatric Medical Association* **79(1)**: 24-26.
63. Sayed, M.D. (1980). Traditional medicine in health care. *Journal of Ethnopharmacology* **2**: 19-21.
64. Lerner, F.N. (1987). Investigation of proteolytic enzymes, Aloe gel and iontophoresis on chronic and acute athletic injuries. *Chiropractic Sports Medicine* **1(3)**: 106-110.
65. El Zawahry, M., Hegazy, M.R. and Helal, M. (1973). Use of Aloe in treating leg ulcers and dermatoses. *International Journal of Dermatology* **12**: 68-73.
66. Bernes, T.C. (1947). The healing action of extracts of Aloe vera leaf on abrasions of human skin. *American Journal of Botany* **34**: 597A.
67. Northway, R.B. (1975). Experimental use of *Aloe vera* extracts in clinical practice. *Veterinary Medicine Small Animal Clinic* **70**: 89.
68. Moon, E.J., *et al.* (1999). A Novel angiogenic factor derived from *Aloe vera* gel: beta-sitosterol, a plant sterol. *Angiogenesis* **3(2)**: 117-123.
69. Heggors, J.P., Pelley, R.P. and Robson, M.C. (1993). Beneficial effects of Aloe in wound healing. *Phytotherapy Research* **7**: s48-s52.
70. Davis R.H., *et al.* (1994). Anti-inflammatory and wound healing of growth substances in *Aloe vera*. *Journal of the American Paediatric Medical Association*. **84**: 77-81.
71. Tizard, A.U. *et al.* (1995). Effect of Acemannan, a complex carbohydrate, on wound healing in young and aged rats. *Wounds, a compendium of clinical research and practice* **6**: 201-209.
72. Talmadge, J., *et al.* (2004). Fraction of *Aloe vera* L. inner gel, purification and molecular profiling of activity. *International Immunopharmacology* **4(14)**: 1757-1773.
73. Collins, C.E. and Collins, C. (1935). Roentgen dermatitis treated with fresh whole leaf of *Aloe vera*. *American Journal of Roentgenology* **33**: 396.
74. Loveman, A.B. (1937). Leaf of *Aloe vera* in treatment of roentgen ray ulcers: Report on two additional cases. *Archives of Dermatology and Syphilology* **36**: 838.

75. Rovatij, B. and Brennan, R.J. (1959). Experimental thermal burns. *Industrial Medicine and Surgery* **28**: 364.
76. Wright, C.S. (1936). *Aloe vera* in the treatment of Roentgen ulcers and telangiectasis. *Journal of American Medical Association* **106**: 1363-1364.
77. Duansak, D., Somboonwong, J. and Putumroj, S. (2003). Effect of *Aloe vera* on leukocyte adhesion and TNF-alpha and IL-6 levels in burn wounded rats. *Clinical Hemorheology Microcirculation* **29 (3-4)**: 239-246.
78. Rodriuez-Bibas, M.D., *et al.* (1988). Comparative evaluation of *Aloe vera* in the management of burn wounds in Guinea pigs. *Plastic and Reconstructive Surgery* **81(3)**: 366-389.
79. Hu, Y., Xu, J. and Hu, Q. (2003). Evaluation of antioxidant potential of *Aloe vera* (*Aloe barbadensis* Miller) Extracts. *Journal of Agricultural and Food Chemistry* **17; 51 (26)**: 7788-7791.
80. Saada, H.N., Ussama, Z.S. and Mahdy, A.M. (2003). Effectiveness of *Aloe vera* on antioxidant studies of different tissues in irradiated rats. *Pharmazie* **58(12)**: 929-931.
81. Parihar, M.S., *et al.* (2004). Susceptibility of hippocampus and cerebral cortex to oxidative damage in streptozotocin treated mice: prevention by extracts of *Withania somnifera* and *Aloe vera*. *Journal of Clinical Neuroscience* **11(4)**.
82. Suga, T. and Hirata, T. (1983). The efficacy of the Aloe plants: chemical constituents and biological activities. *Cosmetics and Toiletries* **98 (6)**: 105-108.
83. Estevez, A., Magdan, R. and Marquina, G. (1982). Chemical study and antitumor activity on Cuban plants, First Latin-American and Caribbean symposium on pharmacologically active natural products. Havana Cuba June 21-28 1980, Unesco 1982.
84. Im, S., *et al.* (2005). Identification of optimal molecular size of modified Aloe polysaccharides with maximum immunomodulatory activity. *International Immunopharmacology* **5(2)**: 271-279.
85. Wang, Z., *et al.* (2001). Study on anti-tumor effect and mechanism of Aloe polysaccharides. *Zhong Yao Cai* **24(5)**: 350-353.
86. Acevedo-Duncan, M., *et al.* (2004). Aloe-emodin modulates PKC isozymes, inhibit proliferation and induces apoptosis in U-373MG glioma cells. *International Immunology* **4(14)**: 1175-1184.

87. Bolkent, S., *et al.* (2004). Effect of *Aloe vera* (L.) Burn. fil. Leaf gel and pulp extracts on kidney in type-II diabetes rats models. *Indian Journal of Experimental Biology* **42(1)**: 48-52.
88. Can, A., *et al.* (2004). Effect of *Aloe vera* leaf gel and pulp extracts on the liver in type-II diabetes rat models. *Biological Pharmacy Bulletin* **27(5)**: 694-698.
89. Rajasekaran, S., Sivagnanam, K. and Subramaniam, S. (2005). Modulatory effects of *Aloe vera* gel extract on oxidative stress in rats treated with streptozotocin. *Journal of Pharmacy and Pharmacology* **57(2)**: 241-246.
90. Rajasekaran, S., *et al.* (2004). Hypoglycemic effect of *Aloe vera* gel on streptozotocin induced diabetes in experimental rats. *Journal of Medicinal Food* **7(1)**: 61-66.
91. Al Awadi, F. and Shoukry, M. (1988). The lipid lowering effect of anti-diabetes plant extracts. *Acta Diabetologica Latina* **25(1)**: 1-5.
92. Roman-Ramos, *et al.* (1991). Experimental study of hypoglycemic activity of some antidiabetes plants. *Archivos Investigation Medica (Mex)* **22(1)**: 87-93.
93. Ghannam, N., *et al.* (1986). The antidiabetes activity of Aloes: Preliminary clinical and experimental observations. *Hormone Research* **24**: 288-294.
94. Al-Awadi, F.M. and Gumaa, K.A. (1987). Studies on the activity of individual plants of an antidiabetes plant mixture. *Acta Diabetologica Latina* **24(1)**: 37-41.
95. Wang, Z.W., *et al.* (2005). Radioprotective effects of Aloe polysaccharides on three non tumor cell lines. *Ai Zheng* **24(4)**: 438-442.
96. Wang, Z.W., *et al.* (2004). Aloe polysaccharides exerted radioprotective effects *in vitro* and *in vivo* through an inhibition of apoptosis. *Journal of Radiation Research (Tokyo)* **45(3)**: 447-454.
97. Oranzo- Barocio, A., *et al.* (1999). Modulation of immune response of BALB/mice bearing lymphoma L5178Y treated with bitter yellow juice of *Aloe vera* (L) *in vitro*. *Russian Journal of Immunology* **4(1)**: 43-50.
98. t' Hart, L.A., Van Enkevort, P.H., Van Dijk, H., Zaat R., De Silva, K.T.D. and Labadie, R.P. (1988). Two functionally and chemically distinct immunomodulatory compounds in the gel of *Aloe vera*. *Journal of Ethnopharmacology*. **23**: 61-71.
99. Strickland, F.M., Pelley, R.P. and Kripke, M.L. (1994). Prevention of ultraviolet radiation induced suppression of contact and delayed hypersensitivity of *Aloe barbadensis* gel extract. *Journal of Investigative Dermatology* **102(2)**: 197-204.

100. 't Hart, L.A., Van Enckevort, P.H. and Labadie, R.P. (1987). Anionic polymers with anti-complementary activity from *Aloe vera* gel. *Pharmacy Weekly* (Sci. Ed.). 9(4): 223.
101. 't Hart, L.A., Van Enckevot, P.H. and Labadie, R.P. (1987). Analysis of two functionally and chemically different immunomodulators from *Aloe vera* gel. *Pharmacy Weekly* (Sci. Ed). 9(2): 157.
102. 't Hart, L.A., *et al.* (1989). Anti-complementary polysaccharides with immunological adjuvant activity from the leaf parenchyma gel of *Aloe vera*. *Planta Medica* 55(6): 509-512.
103. Lim, B.O., *et al.* (2003). Efficiency of dietary *Aloe vera* supplementation on hepatic cholesterol and oxidative status in aged rats. *Journal of Nutritional Science and Vitaminology* (Tokyo) 49(4): 292-296.
104. Kandil, A. and Gobran, W. (1979). Protection of gastric mucosa by *Aloe vera*. *Journal of Drug Research* (Egypt) 11: 191-196.
105. Kandil, A. and Gobran, W. (1982). Protection of gastric mucosa by *Aloe vera*. *Bulletin of Islamic Medicine* 2: 508-511.
106. Langmead, L., Makins, R.J. and Rampton, D.S. (2004). Anti-inflammatory effect of *Aloe vera* gel in human colorectal mucosa in vitro. *Aliment Pharmacology* 19(5): 521-527.
107. Lorenzetti, L.J., *et al.* (1964). Bacteriostatic property of *Aloe vera*. *Journal of Pharmaceutical Sciences* 53: 1287.
108. Levin, H., *et al.* Partial purification and some properties of an antibacterial compounds from *Aloe vera*. *Phytotherapy Research* 2(2): 67-69.
109. Hegggers, J.P., Pineless, G.R. and Robson, M.C. (1979). Dermaide Aloe / *Aloe vera* gel: comparison of the antibacterial effects. *Journal of American Medical Technology* 41: 293-294.
110. Khurana, S.M.P. and Bhargava, K.S. (1970). Effect of plant extracts on the activity of three papaya viruses. *Journal of General and Applied Microbiology* 16: 225-230.
111. Sydiskia, R.J. and Owen, D.C. (1987). Aloe emodin and other anthraquinones and anthraquinone like compounds from plant virucidal against herpes simplex virus. US Patent 4670265.
112. Bhaduri, B., *et al.* (1968). Antifertility activity of some medicinal plants. *Indian Journal of Experimental Biology* 6: 252-253.

113. Goswami, C.S. and Bokadia, M.M. (1979). The effects of extracts of *Aloe barbadensis* leaves on the fertility of female rats. *Indian drugs* **16**: 124-125.
114. Gupta, M.L., Gupta, A.K. and Bhargava, K.P. (1971). A study of antifertility effects of some indigenous drugs. *Indian Journal of Medical Research* **6**: 112-116.
115. Saha, J.C., Savini, E.C. and Kasinathan S. (1961). Ecobolic properties of Indian medicinal plants. Part 1. *Indian Journal of Medical Research* **49**: 130-151.
116. Boyd, L.J. (1928). The pharmacology of the homeopathic drugs 1. *Journal of American Institute of Homeopathy* **27**: 7.
117. Sethi, N., Nath, D. and Sing, R.K. (1989). Teratological evaluation of some commonly used indigenous antifertility plants in rats. *International Journal of Crude Drug Research* **27(2)**: 118-120.
118. Mohsin, A., et al. (1989). Analgesic antipyretic activity and phytochemical screening of some plants used in traditional Arab system of medicine. *Fitoterapia* **60(2)**: 174-177.
119. Gupta, R.A., Singh, B.N. and Singh, R.N. (1982). Screening of Ayurvedic drugs for analgesic activity. *Journal of Scientific Research in Plants and Medicines* **3**: 115-117.
120. Apisariyakul, A. (1984). Investigations from fractions isolated from Thai medicinal plants affecting on isolated rat ileum. Abstr. 10th conference of Science and Technology, Thailand Cheingmai Univ. Cheingmai Thailand, 450-451.
121. Chauhan, J.S., Singh, N.K. and Singh, S.V. (1989). Screening of higher plants for specific herbicidal principle active against Doder, *Cuscuta reflexa* Roxb. *Indian Journal of Experimental Biology* **27(10)**: 877-884.
122. Gupta, R. and Flora, S.J. (2005). Protective value of *Aloe vera* against some toxic effect of arsenic in rats. *Phytotherapy Research* **19(1)**: 23-38.
123. Jae-Hyun Bae and Ok-Yeon Kim (1998). Skin adhesive cosmetics for removing wrinkles, containing vitamins and Aloe extract. US Patent 5723138.
124. Lener, S. (1996). Improved A palmitate composition for topical application which achieves to the entire dermal membrane. US patent 5556887.
125. Alliger, H. and Roozdar, H. (1997). Chlorine dioxide skin medicating compositions for preventing irritation. US Patent 5616347.
126. Fukushima Makote (1989). Cosmetic skin conditioners containing aloenin and its salts. JP Patent 01287009.

127. Chou Belle, L. (2005). Skin-enhancing glove and method of manufacture. US Patent 6953582.
128. Ortiz, R. and Fernandez, V. (2004). Post hair removal skincare lotion. US Patent 6767534.
129. Miyamoto, T. (1986). Skin cosmetic. Application no. JP 61307394.
130. Bates, H.L. (1987). Cosmetic lotion, US Patent 4704280.
131. Egawa, M. (1989). Novel skin drug for external use. Application no. JP Patent 6453238.
132. Smothers, D.L. (2005). Skin and mucosal treatment formulation. US Patent 6949262.
133. Egawe, M., *et al.* (1990). Anti-inflammatory and wound healing topical skin preparation containing Aloe extract and elleric acids. JP Patent 02231408.
134. Nieto Burgos, C. (1991). Hair generator based on Aloe extract. Span Patent 2019828.
135. Coutts, B.C. (1979). Stabilized *Aloe vera* gel. JP Patent 79119018.
136. Cobble, H.H. (1975). Stabilized *Aloe vera* gel. US Patent 3892853.
137. Maughan, R.G., Poore, R.A. and Phan B.V. (2005). Product and process for stabilizing *Aloe vera* gel. US Patent 6869624.
138. Beck, F. (1991). Camphor and menthol based analgesic composition useful in providing a temporary relief from symptoms of arthritis. US Patent 5073366.
139. Lane, B. (1996). Lip balm composition. US Patent 5503825.
140. Schuiman, J.M. (1996). Medicated gel. US Patent 5503822.
141. Van, N.V. (1996). Pad for applying medicaments. US Patent 5480646.
142. Burns, M.J. (1995). Topical agent and method for the treatment of pseudofolliculitis barbae. US Patent 5435997.
143. Isnard, C. (1995). Aloe water preparation. US Patent 5427776.
144. Wunderlich, J.C. and Schick, U. (1995). *Aloe vera* juice containing pellets process for production thereof and the use thereof as pharmaceutical, cosmetic and peroral agent. US Patent 5387415.
145. Hikino, H. and Hayashi, T. (1985). Hypoglycemic polysaccharides extraction from Aloe species. JP Patent 60214741.
146. Tsujino Kaoru (1990). Thick liquid hand cleansing agents. JP Patents 02215899.
147. Takaoka Takio (1990). Wet sanitary pads for cleansing. JP Patent 02182999.

148. Morita Ysumasa (1989). Detergents and soaps containing aloe and herbs. JP Patent 01104700.
149. Smith, W.P. (1989). Post-foaming shave gel composition. European Patent 339634.
150. Mitchell, Jr.C. and Sanders, W. (2005). Shaving soap and aftershave gel and method of use thereof. US Patent 6893631.
151. Bates, H.L. (1989). Decongestant comprising zinc and vegetable oil. US Patent 4826683.
152. Maret, R.H. and Cobble, H.R. (1975). Extracts of *Aloe vera*. US Patent 3878197.
153. Jordan, F.L. (2005). Transdermal delivery system. US Patent 6946144.
154. Taylor and Allan, (2005). Preparation for regulating lower bowel function. US Patent 6916495.
155. Glick, F. and Puchalski, Jr.E. (2005). Antibacterial compositions and method of using same. US Patent 6927197.
156. Michimasu, H., *et al.* (1996). Fibroblast proliferation promoting agent. Application no. JP Patent 08208975.
157. Nagayoshi, S. (1990). Separation of bitter component in Aloe extract. Application no. JP Patent 2273435.
158. Litke, K.S., and Widdemer, J.D. (2005). Aloe Vera processed leather and leather gloves, garments, shoes and sandals made from Aloe Vera processed leather and a process for making Aloe Vera processed leather. US Patent 6953483.
159. Peng, S.Y., *et al.* (1991). Decreased mortality of norman murine sarcoma in mice treated with the immunomodulator, acemannan. *Molecular Biotherapy* 3: 79-87.
160. Kahlon, J., *et al.* (1991). Inhibition of AIDS virus replication by acemannan *in vitro*. *Molecular Biotherapy* 3: 127-135.
161. Yetes, K.M., *et al.* (1992). Pilot study of the effect of acemannan in cats infected with feline immunodeficiency virus. *Veterinary Immunology and Immunopathology* 35: 177-179.
162. *Quality Control Methods for Medicinal Plant Materials*, (1998). World Health Organization, Geneva.