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**ALOE BARBEDANSIS MILL- A REVIEW OF ITS ETHNOBOTANY,
PHYTOCHEMICAL AND PHARMACOLOGICAL PROFILE**

**Amit Kumar Singh, Kuldip Kumar, Vaishnavee Gupta, Rahul Kumar, Triveni,
Kishu Tripathi***

Smt.Vidyawati College of Pharmacy, Jhansi (UP) 284121, India.

ABSTRACT

Aloe vera is species of succulent plants belonging to family Liliaceae. It is easily available, cheap, multipurpose, mucilaginous plant. Various herbal formulation of *Aloe vera* are available in market like herbal soap, face wash, shampoo, cream etc Moreover, *Aloe vera* has been reported to have various pharmacological activities like wound healing, anti-inflammatory, antioxidant, analgesic, antibacterial, antiviral, antidiabetic, anti-hypercholesteremic etc. Present review article will provide a guideline for new researchers.

Keywords: *Aloe barbedansis*, Ghritakumarika, Antioxidant, Anxiolytic.

INTRODUCTION

Aloe vera (*Aloe barbedansis* miller) is a plant, belonging to family of Liliaceae and is mostly succulent with a whorl of elongated leaves. The name is derived from the Arabic words "aloe" which means "bitter" referring to the taste of the liquids present in leaves[1]. Plants derived natural products such as flavonoids, terpenoids and steroids etc. have received considerable attention in recent years due to their diverse pharmacological properties such as antitumor, antioxidant [2]. *Aloe vera* has been medicinally used for several thousand years in many cultures from Egypt, Greece, Rome, China and India [3]. Different bioactive constituent medicinal value of *Aloe vera* treat different type of disease [4]. *Aloe vera* is such a plant which is known to process a wide spectrum of medicinal and therapeutic properties scientifically established by the modern scientific practioner [5]. *Aloe vera* has been promoted for large variety of conditions such as diabetes mellitus, hyperlipidemia, inflammation, treatment of acne etc. and has come to play a prominent role as a contemporary folk remedy [6]. Several herbal preparations that can enhance the body's immune system status are extensively being used in the indigenous system of medicine. There are upsurges in the clinical usages of indigenous drugs as they are free from serious side effect [7].

Probably native to North Africa along the upper Nile in the Sudan, and subsequently introduced and naturalized in the Mediterranean region, most of the tropics and warmer areas of the world, including Asia, the Bahamas, Central

America, Mexico, the southern United States of America, southeast Asia and the West Indies [8].

TAXONOMICAL HIERARCHY

Besides the usual classification medicinal plants can be classified according to the parts used, habits, habitat, therapeutic value etc. But the botanical classification is the most comprehensive and scientific classification which is following:

Kingdom	Plantae
Subkingdom	Viridaplantae
Phylum	Lilliacae
Class	Magnoliopsida
Order	Asparagales
Family	Xanthorrhoeaceae
Genus	Aloe
Species	barbedansis

VERNICULAR NAMES

English:	Indian Aloe, Barbados aloe
Hindi	:Ghikunwar, Ghikumari
Kannada	:Kathaligida
Bengali	:Ghritakumari
Marathi	:Korpad
Telugu	:kalabanda
Tamil	:Kattalai
Gujarati	:Kumarpathu
Malayalam	:Kattuwal
Urdu	:Musabbar, Ailiva, Siber

Synonymes: Kumari, Grihakanya, Ghritkumarika, Kanya
Plants parts used: Leaf, Leaf-juice, dried juice of leaf

MORPHOLOGY



Leaves large, succulent, subulate, sessile, 20-50 cm long and 5-10 cm wide. Apex in the form of a sharp and acute spine. Both the surface are strongly cuticularized. Dried leaf juice dark chocolate brown to black in colour and of irregular masses. Odour characteristic and taste very bitter. In transverse section pericycle cells are thin walled and large containing yellow fluid. The fluid under microscope shows crystals in the form of innumerable needles varying in size and shape. Powder of dried juice when mounted in glycerol lactophenol and examined under microscope, shows innumerable crystalline, yellowish brown to chocolate coloured particles of varying shape and size [8].

DISTRIBUTION

The plant aloe is as old as human civilization and its properties for various purposes have been well documented. The genus is found in Tropical and South Africa, Malagasy and Arabic and introduced in other places for ornamental and medicinal purposes. *Aloe vera* is a succulent plant, almost sessile perennial plant with

multiple tuberous roots and many fibrous supporting roots and many fibrous roots penetrate into the soil. Each leaf is 30-50 cm long and 10 cm broad at base and margin is indented [9].

Phytochemistry

The chief constituents are hydroxyanthraquinone-barbaloin (a mixture of aloin A and B, the distereoisomeric 10-C glucoside of aloe emodin anthrone) and γ -hydroxyaloin isomers. The other constituents include aloin, chrysophanol, chromone derivative-aloeresin B with its p-coumaroyl derivative aloeresin A and C and the glycon aloesone [8]. Leaves identified their constituents of bioactive compounds (viz., glycosides, anthraquinone, phlobatannins, carbohydrates, alkaloids, terpenes, saponins, tannins, steroids, flavonoids), which believe to be responsible for their medicinal use [10].

Folk remedies and traditional uses

Traditionally the aloe leaf juice is most useful for healing of fractures and bones. Plants have been documented in Ayurvedic for antihelminthic, apparent carminative, deobstruent, diuretic, ophthalmic and alexeteric, juice used in dyspepsia, amenorrhoea, burn, colic hyperdenosis, hepatopathy, skin disease, tumor etc.

Toxicity study

Plants extract at a dose of 100 and 200 mg/kg did not show any toxicity in rats. Prolonged use may severely affect the electrolyte balance. Loss of potassium may ultimately reduce the laxative action and disturb cardiac rhythm in heart patients. Longer use may cause accumulation of blood in pelvic region and reflex stimulation of uterine muscle and may bring about abortion [8]. Cytotoxicity of *Aloe vera* gel crude extract on vero cells was determined by calculation of CC50 which is 3238 μ g/ml [22].

Table 1. Class of compound according to their amount in percentage

Class of Compound	Group of component	Amount
Resin wax and Fatty acid	Flavonoids, Phenolic acid and Ester	45-55%
Wax and Fatty acid	Bee wax and plant origins	23-35%
Essential oil	Volatiles	10%
Pollen	Protein (16 Free amino acid > 1%) arginine and proline together total of 45%	5%
Other organics and minerals	14 trace minerals, Iron and zinc most common Ketone, lactone, Quinone most common, vitamins, sugar	5%

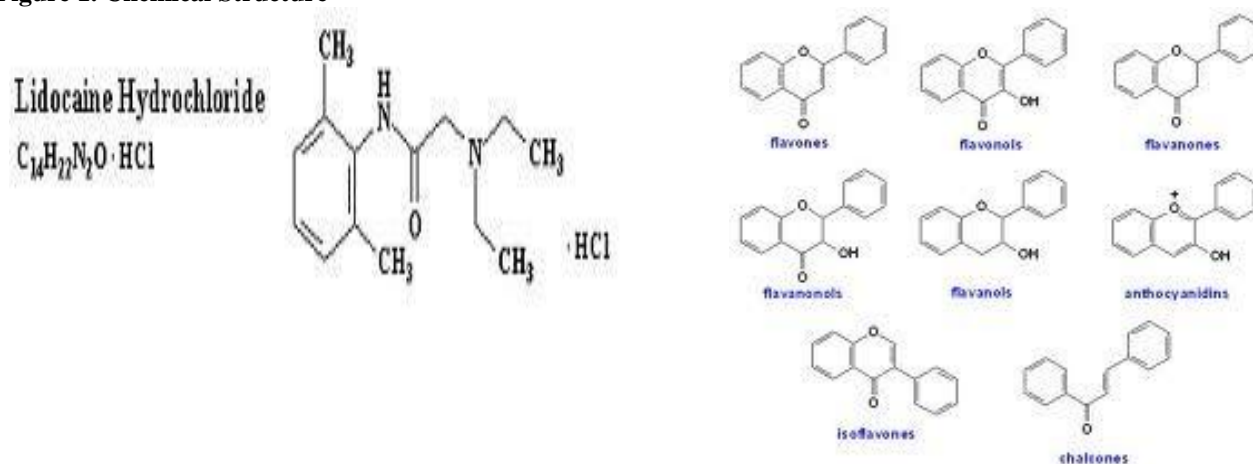
Table 2. Various traditional uses are of *Aloe barbadensis*

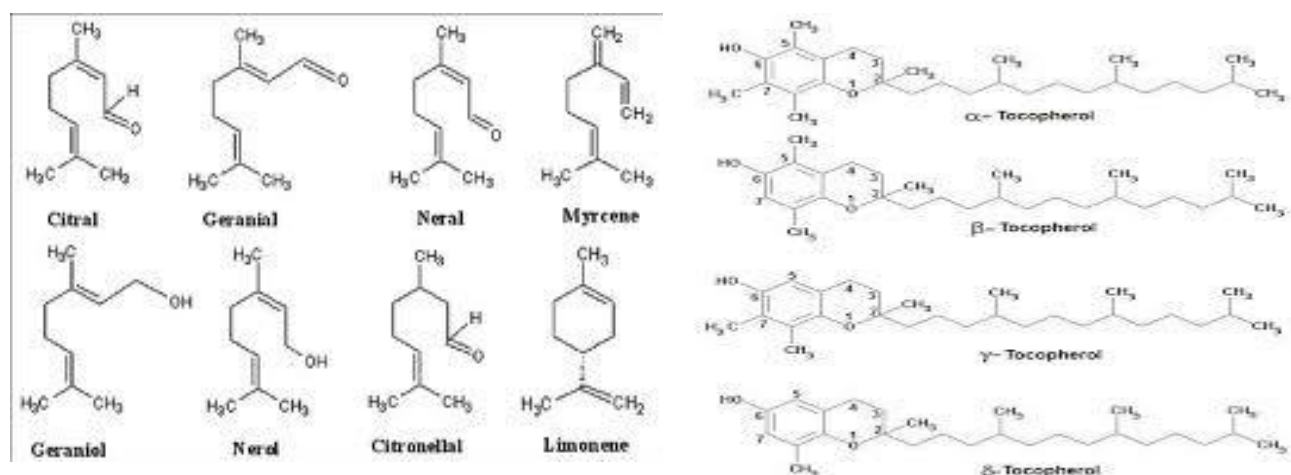
Plant Part used	Activity	Active Constituent	Reference
Leaf juice	Photo protective effect	Resin and Polysaccharides	[11]
Leaf juice	Wound healing	Glycoprotein	[12]
Leaf juice	Burn wound healing	Mannose 6-Phosphate	[13]
Leaf juice	Nitric oxide production	Arginine	[14]
Leaf juice	Cosmetic ingredients	Aloinodides, aloin	[15]
Leaf juice	Reduce obesity	Glycosides, mannoses, fructose	[16]
Leaf juice	Cooling, constipation, hepatopathy, Skin disease, dysmenorrhoea	Aloin, aloin	[8]
Leaf juice	Anti-tumor	Flavonoids, triterpenoids, steroids	[2,17]

Leaf juice	Inflammation,Diabeties	Saponin, sterol	[18]
Leaf juice	Multi drug resistance	Anthraquinon	[19]
Leaf juice	Synergistic effect in bacteria	Aloin	[20]
Leaf juice	Hemorrhagic shock	Aloin	[21]

Table 3. Pharmacological study of *Aloe barbedansis*

Pharmacological study	Active Constituent	Reference
Antioxidant activity	Polysaccharide, flavonoids	[23]
Analgesic activity	Lignin, Saponin, Anthraquinon	[24]
Analgesic and anti-inflammatory activity	Mallic acid, Acylated carbohydrate	[25,56]
In vitro antibacterial activity	Mannans, Polymannans, Anthraquinons, C-Glycosides, Lectin	[26]
Antimicrobial activity	Phenol, lignin Saponin, Sterol, Amino Acids,flavonoid	[27,45,52]
Antitumor activity	Flavonoids, Triterpenoids, Steroids	[28]
Antiviral activity	Monosaccharides(Mannose-6-Phosphate)	[29]
Anxiolytic activity	Mannose, Glucose,	[30]
Burn wound healing	Mannose -6- Phosphate	[31,40]
Diabetic control	Triglyseride	[32,41]
DNA degradation activity	Aloin, Aloe emodin	[33]
Degenerative CNS disorder activity	Phenol, Flavonoids	[34]
Immunomodulatory activity	Polysaccharides', Glycoprotein	[35]
Increase Nitric oxide production activity during inflammation	Arginine	[36]
Photo protective activity	Resin, Polysaccharides	[37]
Toxicological activity	Polysaccharides, mallic acid.	[38]
CNS activity	Chlorides	[39]
Antifungal activity	Sugar, lignin, Anthraquinon, Saponin.	[42,44]
Hepatoprotective activity	Glycoside, Aloe emodin, Glucose, Mannose	[43,48,50]
Antifungal and antioxidant activity	Lignin, Anthraquinon	[44]
Anticarcinogenic activity	Flavonoids, terpinoids, Steroids	[46]
Hypoglycemic and atherogenic activity	Phenol, flavonoids	[47]
Genotoxic and antixenotoxic effect	Polysaccharides,	[49]
Hypocholestromic effect	Flavonoids	[51]
Antinociceptive activity	Flavonoids, Salicylates	[53]
Antiseptic ulcer activity	Calcium Carbohydrate, Phosphate	[54]
Protective nephrotoxic activity	Flavonoids	[55]
In vitro antileishmanic activity	Tannins, Flavonoids	[57]
Haemopoitic activity	Flavonoids	[58]

Figure 1. Chemical Structure



CONCLUSION

Phytochemical and pharmacological investigation carried out in the plant reveals its multi-disciplinary usages. The plant was found to be very useful to reduce obesity, anti-inflammatory, analgesic, antioxidant, hepato protective, antitumor property, etc. Several investigators have reported the plants as valuable antimicrobial, antidiabetic, antifungal, antipeptic ulcer, antileishmanial etc, and also active against other plants pathogens. According to the proper documentation the plant Aloe do

not show the toxic effect in animal in normal amount but it is used in combination with other plant herb and biological preparation like Panchagavya to show the synergistic effect with combination to treating the various ailments. Thin layer chromatography (TLC) profile of *Aloe Vera* extract was carried out to confirm the presence of different phytoconstituents different combination of solvent system of varying polarity has been used for the optimization of solvent system in high performance liquid chromatography [59].

REFERENCES

1. Bashir A, Saeed B, Mukhid TY, Jehan N. Comparative study of antimicrobial study of *Aloe vera* extract and antibiotic again isolated from skin infection. *African Journal of Biotechnology*, 10(19), 2011, 3835-3840.
2. Naveena, BharatBK, Selvasubramanyam. Antitumor activity of *Aloe vera* against ehrlich associated carcinoma in swice mice. *Int.Jorn of Pharma and Bioscience*, 2(2), 2011, 400-409.
3. Abd-Alla Hi, Abu-Gabel NS. Antiviral activity of *Aloe vera*. *Arch Pharm Res*, 35(8), 2012, 1347-1354.
4. Vijay R, Krishna Reddy. Antioxidant activity of *Aloe vera* due to medicinal property. *Journ of Phytotherapy and Pharmacology*, 1(6), 2012, 01-13.
5. Dutta B. A study of patenting activity in *Aloe vera*. *Journ of Intellectual Property right*, 7, 2002, 330-341.
6. Bahram AT, Daryoush M, Study of Oxidative stress activity. *World Journal of Zoology*, 7(3), 2012, 192-199.
7. Atul NC, Santosh K, Subal B. Study of Immunomodulatory activity. *Int Journ of applied Biology and Pharmaceutical tech*, 2(1), 2011, 19-22.
8. The Wealth of India Raw Material: A Dictionary of Indian Raw Material and Industrial Products, Council of Scientific Indian Research New Delhi, 2, 1952, 56-57.
9. Nilanjana Das, Chatopadhyay RN. Commercial cultivation of *Aloe vera*. *Natural product Rediance*, 3(2), 2004, 85-87.
10. Ejoba Raphael. Phytochemical constituent of some leaves extract of *Aloe vera* and *Azadirachta indica*. *Global advance journal of environ Science and Taxicology*, 1(2), 2012, 14-17.
11. Daud FS, Kulkarni SB. Comparative evaluation of photo protective effect of *Aloe vera* on UV Damage in different Asian hair types. *Indian Journal of Natural Product and Resources*, 2(2), 2010, 179-183.
12. Robert HD, Mark GL, Joseph MR, Mogon ES. Wound healing oral and tropical activity of *Aloe vera*. *Jorn of American Pediatric Medical Asso*, 79, 1989, 559-562.
13. Ratree M, Nothern C, Surachet N. Efficacy of *Aloe vera* for burn wound healing. *Science direct*, 33, 2007, 713-718.
14. Sarkar D, Dutta A, Das M, Sarkar K, Manal C, Chaterjee C. Nitric oxide production activity during inflammation by *Aloe vera*. *Ind Jour of Pharmacology*, 37, 2005, 371-375.
15. Gauri B, Neha J, Farhat D. A Valuable multicosmetic ingredient of of *Aloe vera*. *Int.J.Med.Arom*, 1, 2011, 338-341.
16. Shin S, Kim S, Kong H. Dietary *aloe vera* reduce obesity. *PubMed*, 2, 2002, 139-142.
17. Agrawal RC, Sonam P. Evaluation of anticarcinogenic activity of aloe vera in swice albino mice. *Int Journ of scientific and Research Publication*, 3, 2013, 01-06.
18. Davish RH, Maro NP. Anti-inflammatory activity in diabetic rat. *J Am Pedatric Med Asso*, 79, 1989, 24-26.
19. Asima Banu, Satyanarayan BC. Efficacy of fresh *Aloe vera* gel again multiple drug resistance in bacterial infected leg ulcer. *Australas Med J*, 05, 2012, 305,309.
20. Ghamari F, Ghaffari SM, Salim M, Moosavi-Mohavedi F, Ferivar F, Johri A. Synergistic effect of *Aloe vera*. *PubMed*, 2012,

21. Lu J Xia O, Genq ZL, Liu D. Effect of Haemorrhagic shock on rat. *Zhonghuawai Keza zhi*, 50, 2012, 655-658.
22. Kievam Z, Moloud A.Z., kohzad S. Antiviral activity of *Aloe vera* against herp simplest virus-2. *African Journal of Biotech*, 6, 2007, 1770-1775.
23. Yun Hu, Jaun Xu, qihui Hu. Evaluation of antioxidant activity. *J Agri.Food Chem*, 51, 2003, 7788-7791.
24. Ghosh AK, Banerjee M, mandal TK, mishra Akhilesh. A study on analgesic effect and adverse effect in wistar rats. *Pharmacologyonline*, 1, 2011, 1098, 1108.
25. Mwale M, masika PJ. Analgesic and Antiinflammatory activity of *Aloe vera*. *AJPP*, 4, 2010, 291-297.
26. Irshad S, Butt M, younush H. Invitro antibacterial activity of *Aloe vera*. *Int.Res.Journ of Pharmaceuticals*, 01, 2011, 59-64.
27. Bashir A, Saeed B, Mujahid TY, Jehan N. Comparative study of antimicrobial of *Aloe vera* and antibiotic again skin infection. *A.J.of Biotech*, 10, 1990, 3835-3840.
28. Naveena, Bharath BK, Selusubramanyam. Antitumor activity of *Aloe vera*. *Int.Journ of pharma and Bio Science*, 2(2), 2011, 400-409.
29. Zandi K, Zedesh MA, Rastian Z. antiviral activity of *Aloe vera* against herp. *A.J.of Biotech*, 6(15), 2007, 1770-1773.
30. Sultana N, Najam R. Anxiolytic activity of *Aloe vera*. *Pakistan Journ of pharmacology*, 29, 2012, 07-13.
31. Maenthisang R, Chaiyakunapruk N, nirutrapan S, The efficacy of burn wound healing of *Aloe vera*. *Science Direct*, 33, 2010, 713-718.
32. Ken jones. Glycemic control by *Aloe vera*. *Diebitic.Org*, 2007, 01-06.
33. Naqui S, Ullah MS, Hadism. DNA Degradation activity by *Aloe vera* in presents of copper ions. *Idian Journ of Biochen ans Biophysis*, 17, 2010, 161-165.
34. Ozosovn, Chanoken. Degradation activity by *Aloe vera*. *Oxi Med Cell*, 2(2), 2009, 96-10.
35. Chandu AN, Kumar S, Bhattacharjee C, Devnath S. Study on Immunomodulatory activity by *Aloe vera*. *Int Journ of Applied Bio and pharmaceutical tech*, 2(1), 2011, 19-22.
36. Sarkar D, Dutta A, Das M, Sarkar K, Chaterjee M. Effect of *Aloe vera* on nitric oxide production by macrophages during inflammation. *Indian J Pharmacol*, 37, 2005, 371-375.
37. Daud FS, Kulkarni SB. Phytochemical effect of hair damage by *Aloe vera*. *Ind Journ of Natural Product and Resources*, 02, 2008, 179-183.
38. Stemcamp V, Stevart MJ. Medicinal value and Toxicological effect of *Aloe vera*. *Openup research paper*, 2007,
39. Madhusudhan N, basha PM, Rai P, Ahmed F, Prasad GP. Role on CNS express by natural fluid exposer by *Aloe vera*. *Ind Journ of Exp Bio*, 48, 2010, 430-436.
40. Harith yadav KC, Ravi J, Basha SI, Deshmukh GR, Gujja R. Wound healing activity of *Aloe vera* in rodent. *Int Journ of Pharma and Bio Science*, 3(2), 2012, 63-72
41. Enas ali kamel Muhammad. Antidiabetic, antihypercholestromic and antioxidant activity of *Aloe vera* in alloxan induce diabetic rats. *Australian journ of basis and applied Science*, 5(11), 2011, 1321-1327.
42. Sitara V, Hussan N, Naseem j. Antifungal activity of *Aloe vera* gel again plant pathogen fungi. *Pak.Journ of Bot*, 3(4), 2011, 2231-2233.
43. Bahram AT, Daryous M. *Aloe vera* leaf extract attenuate oxidative extract oh hepatic tissue. *world Journ of zoology*, 07(3), 2012, 192-199.
44. Tin A.khaing. Evaluation of antifungal and antioxidant activity of *Aloe vera*. *World academic of science, eng and Tecn*, 51, 2011609-611.
45. Thiruppathi, Ramasubramanyam S, shivkumar v, thirumalli A. Antimicrobial activity of *Aloe vera* again pathogen microorganism. *J.Bio.Science and researchpublication*, 1, 2010, 251-258.
46. Agrawal RC, Paney S. Evaluation of anticarcinogenic activity of *Aloe vera* in mice. *Int.Journ of Science and Research*, 3(2), 2013, 01-06.
47. Gupta A, Sethi J, Sood S, Dahiya K, Singh Gajynder, Gupta R. Evaluation of hyperglycemic and atherogenic effect of *Aloe vera*. *IJCP*, 8(3), 2011, 01-07.
48. Sharma HD. Hepatoprotective effects of *Aloe vera* against carbon tetra chloride induce hepatotoxicity. *IJRPBS Online*, 3, 1119-1124.
49. Kayraldiz A, Yayuz Kochman A. The Genotoxic and antigenotoxic effect of *Aloe vera* in vivo and in vitro. *Turk j boil*, 34, 2010, 335-346.
50. Sharma YK, Singh H, Mehta BL. Hepatpprotective effect of *Aloe vera* herb in patient having tuberculosis treatment. 3(4), 2004, 391-396.
51. Chandrakar M, Palekar S, Chirade S, Hafiz AM. Hypocholestremic effect of *Aloe vera*. *Asian Journ of exp Sci*, 22(3), 2008, 295,298.
52. Shrinu B, Vikram B, Raol V, Kalakumar B, Rao TM, Reddy AG. Screening of antimicrobial activity of *Aloe vera* and *Withania somnifera*. *JOCPR*, 4(11), 2012, 4800-4803.
53. Shuraki M.R, Mirshikari H, Shahraki H. Antinociceptive activity of *Aloe vera* in fractise feed rats. *Basic and clinical neuro science*, 1(3), 2013, 39-43.
54. Chauhan O, Gehlat A, Rathod R, Chjaudhary R. Antipeptic ulcer activity of *Aloe vera*. *Pak Journ of physio*, 6(2), 2000, 29-31.

55. Chaterjee P, Mikherjee A, Nandy. Protective nephrotoxic effect of *Aloe vera*. *Assian pacific Journ of tropical Biomed*, 2012, S1751-S1762.
56. Vazaques B, Avila G, Segura D, Escalente B, Antiinflammatory activity of *Aloe vera* gel. *Journ of Ethanopharmacology*, 55, 2005, 69-75.
57. Dutta A, Mandal G, Chaterjee M. Anleishmanic activity of *Aloe vera* leaf exudates. *Glycoconj J*, 24(1), 2007, 81-86.
58. Osonuga OA, Osonuga OY, Osonuga AA. Haematological activity of antiretroviral drugs in rats. *Asian Journ of Med Sci*, 1, 2010, 41-44.
59. Patel DK, Patel K, Dhanabal SP. Phytochemical standardization of *Aloe vera* extract by HPTLC Technique. *Journ of acute disease*, 2012, 47-50.