Review Article

A review on kalanchoe pinnata (Crassulaceae)

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A B S T R A C T

The primary goal of this study is to offer preliminary data for drug discovery research using Kalanchoe pinnata, a heavenly plant that has a broad variety of active chemicals, including alkaloids, Phenols, Phenylpropanoids, Flavanoids, Triterpenoids, steroids, organic Salts. This plant was discovered to have a variety of pharmacological properties, including Antihypertensive activity, Hepatoprotective activity, Antimutagenic activity, Anti-ulcer activity, Uterine Contractility, Antidiabetic activity, Wound-healing activity, Antioxidant activity, Antimicrobial activity, Antileishmanial activity, Insecticidal activity, Antipyretic activity, Antilithic activity, Neuropharmacological, Immunosuppressive antibacterial activity, Cytotoxicity of testis. This study provides phytoconstituents and pharmacological activity of K. pinnata, a medicinal plant that may help researchers conduct more advanced qualitative research.

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1. Introduction

Medicinal plants have been valued for millennia as a rich source of medicinal substances for the prevention of illnesses and afflictions all over the globe.¹ Kalanchoe pinnata (Crassulaceae) is an erect, succulent perennial shrub that grows to be approximately 1.5 metres tall and reproduces both vegetatively and via seeds. It features tall hollow stems, black bell-like pendulous blooms, and newly dark green leaves that are scalloped and trimmed with red. This plant may be readily propagated by cutting stems or leaves.²,³

Phytochemicals may protect hominids against a wide range of diseases. Phytochemicals are nonnutritive plant combinations that provide caring, therapeutic, or disease-fighting properties. Plants generate these compounds to protect themselves; however, new research shows that certain phytochemicals may protect animals from more than only syndromes. In pods and sages, there are many phytochemicals, each with its own mechanism.⁴

1.1. Taxonomical classification⁵

1. Kingdom: Plantae — plants
2. Subkingdom: Tracheobionta — vascular plants
3. Division: Spermatophyta — seed plants
4. Subdivision: Magnoliophyta — flowering plants
5. Class: Magnoliopsida - dicotyledons
6. Subclass: Rosidae
7. Order: Rosales
8. Family: Crassulaceae — stonecrop
9. Genus: Bryophyllum
10. Species: B. pinnatum (Lam.) Oken.
1.2. Common names
Cathedral Bells, Air Plant (USA), Life Plant, Miracle Leaf, Goethe Plant, and Katakataka are some of the common names for this plant.

In the English-speaking Caribbean, it’s also known as "Wonder of the World." Mother Of Thousands, Herbe Mal Tete (Dominica), Never Dead, Parvu, Hoja Del Aire (Hawai‘i), Oliwa Ka Kahakai (Hawai‘i), Oliwa Ka Kahakai (Hawai‘i), Oliwa Ka Kahakai (Hawai‘i), and Oliwa Ka Kahakai (Bolivia).

1.3. Vernacular name
1. Sanskrit: Parnabeeja, Asthibhaksha
2. English: Air plant
3. Hindi: Zakhmhaiyat, Pathharchoor
4. Kannada: Gandukalinga, Kadu basale
5. Malayalam: Elamarunga
6. Tamil: Malaikalli, Ranakalli
7. Telugu: Ranapala
8. Marathi: Gayamari
9. Bengali: Koppatha, Pathar kuchi.

1.4. Synonyms
Bryophyllum calycinum Salisb., Kalanchoe pinnata (Lam.) Pers., Cotyledon pinnata Lam., and Sedum madagascaricum Clus.

1.5. Phytochemistry
The Phytoconstituents of plants shown in Table 1 and classified with compounds.

1.6. Worldwide ethnomedical uses
Worldwide traditional used of plant are shown in Table 2

1.7. Pharmacological activities
Pharmacological activities of Kalanchoe pinnata are shown in Figure 1 and discussed in details following are

1.8. Antihypertensive activity
This research examined the effects of aqueous leaf extract of K. pinnata on the blood pressure of anaesthetized cats, as well as the liver and kidney function of the rabbit. The findings indicated that the extract lowered the anaesthetized cat’s blood pressure somewhat and also mitigated the impact of adrenaline-induced hypertension. This research demonstrated the pharmacological foundation for the Igbo of Nigeria’s usage of K. pinnata to reduce blood pressure. However, the fact that the blood pressure decrease is minimal and the K. pinnata leaf extract may be harmful to humans precludes its usage as a blood pressure reducing agent.

1.9. Hepatoprotective activity
In the folk medicines of India’s Bundelkhand region, the juice of fresh leaves is extremely useful for curing jaundice. The ability of the leaves’ juice and the ethanolic extract of the marc left over after expressing the juice to protect against CC14-induced hepatotoxicity was evaluated in rats. In vitro, in vivo, and histology studies revealed that the test chemical was hepatoprotective. The juice was shown to be more effective than the ethanolic extract.

1.10. Antimutagenic activity
Antihistamine and antiallergic effects are found in the plant. Methanol extract from the leaves has also been found to inhibit histamine receptors (H1) in the ileum, peripheral vasculature, and bronchial muscle, thus guarding against chemically induced allergic reactions and death by selectively inhibiting histamine receptors in the lungs. Quercetin-3-o—Larabinopyranosyl (12) Obaseiki- Ebor et al. found that organic solvent extracts of leaves inhibited His — to His + reverse-mutations caused by ethyl methanesulphonate acting on S. typhimurium TA100 or TA1002, as well as reversions caused by 4nitro-o-phenylenediamine and 2-aminofluorene in TA98. The alkaloidal/water soluble and acid fractions have no apparent antimutagenic efficacy.

1.11. Anti-ulcer activity
In this research, the incidence of ulceration and mean basal and histamine induced stomach acid production were significantly reduced in a dose-dependent manner, supporting its usage as an anti-ulcer drug in traditional medicine.

1.12. Antidiabetic activity
The presence of zinc in the plants may indicate that they could be useful in the treatment of diabetes caused by insulin dysfunction. The antinociceptive effect of the herb’s aqueous leaf extract was tested in mice using the ‘hot-plate’ and ‘acetic acid’ pain models. Fresh egg albumin-induced pedal oedema and streptozotocin-induced diabetes mellitus were used to test the plant extract’s anti-inflammatory and anti-diabetic properties in rats. In mice, the aqueous leaf extract had substantial antinociceptive effects (P0.05-0.001) against thermally and chemically generated nociceptive pain stimuli.

In rats, the plant extract reduced fresh egg albumin-induced acute inflammation and caused substantial hypoglycemia (P0.05-0.001). The herb’s flavonoids, polyphenols, triterpenoids, and phytosterols are thought to be responsible for the plant’s antinociceptive, anti-inflammatory, and antidiabetic effects. It is thought to have antinociceptive and anti-inflammatory properties.
Table 1: Phytoconstituents of kalanchoe pinnata

| Classification | Plant part | Compound |
|---------------|-----------|----------|
| Phenols phenylpropanoids and flavanoids | Aerial parts of plants | Syringic acid, caffeic acid, 4-hydroxy-3-methoxy-cinnamic acid, 4-hydroxybenzoic acid, p-hydroxycinnamic acid, p-hydroxybenzoic acid, ferulic acid, protocatechuic acid, phosphoenolpyruvate, protocatechuic acid, astragalin, 3,6,9-trihydroxy-4', 5, 6, 9-tetrahydroxyflavone, friedelin, epigallocatechin-3-O-syringate, luteolin, rutin, kaempferol, quercetin, quercetin-3L-rhamnoside-L-arabinose furanoside, quercetin-3-O-diarrabinoside, kaempferol-3-glucoside, quercetin-3-O-α-L-arabinosyl(1→2)β-L-rhamnoside. |
| | Leaves | astragalin, 3,8-dimethoxy-4,5,7-trihydroxyflavone, friedelin, epigallocatechin-3-O-syringate, luteolin, rutin, kaempferol, quercetin, quercetin-3L-rhamnoside-L-arabinose furanoside, quercetin-3-O-diarrabinoside, kaempferol-3-glucoside, quercetin-3-O-α-L-arabinosyl(1→2)β-L-rhamnoside. |
| Triterpenoids | Whole plant | 12β-amyrin, 12β-amyrin acetate, 13β-amyrin, 13β-amyrin acetate, bryophollenone, bryophollone, taraxerol, Ψ-taraxasterol, pseudo taraxasterol, 18β-oleanane, friedelin, glutinol. |
| Steroids | Aerial parts | 12β-sitosterol, bryophyllol, bryophyln B (Antitumor), bryophyllin A (bryotoxin C, bufadienolide I, 3, 5-orthocacetate) with potent cytotoxicity, a insecticidal bufadienolide bryophyllin C and bersaldegenin-3-acetate, bryotoxin A, bryotoxin B, bersaldegenin-1, 3, 5-orthocacetate, campsterol, 24-ethyl-25-hydroxycholesterol, isofernasterol, clionasterol, codisterol, peposterol, 22-dihydrobrassicasterol, clerosterol, patuletin, 3-O-(4-O-acetyl-α-L-rhamnopyranosyl)-7-Orhamnopyranoside patuletin. |
| Organic salt | Leaves | Complexed between a hydro amino acid and malic acid (1:2) |

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Table 2: Traditional uses of kalanchoe pinnata.

Leaf extract is taken in empty stomach is used in the treatment of urinary bladder stones and fewer in children’s. India For diarrhea The leaves juice is used against cough, dysentery, Leaf juice externally applied to scabies and leucoderma and leaf decoction applied over cuts to stop bleeding. For abdominal discomfort, boils, bruises, cholera, cuts, diabetes, diarrhea, dysentery, flatulence, headaches, kidney stones, indigestion, insect bites, scabies, sores, urinary insufficiency, wounds. 15,16 For eye infections, headaches, inflammation, menstrual disorders, pimples, wounds. 17,18 For aches, burns, childbirth, colds, coughs, fever, headache, pain, respiratory infection. 17,18 For bacterial infections, boils, broken bones, bronchitis, cancer (lymphoma), conjunctivitis, coughs, earaches. 17,18 For eye infections, epilepsy, erysipelas, fever, gas, headache, heartburn, inflammation, intestinal problems, migraine, nausea, skin problems, sores, ulcers, urethritis. 17,18 For asthma, chest colds, earaches, headaches, sores, strains, tumors. 17,18 For chicken pox, fevers, stomachache Mexico For arthritis, asthma, bruises, burns, constipation, diabetes, earaches, headaches, malnutrition, migraines, nephritis, paralysis, respiratory infections, rheumatism, sprains, swelling, ulcers, wound, and to induce vomiting of blood. 19 For antibacterial and anti-inflammatory. 20 For fever, headache, heartburn, inflammation, intestinal problems, migraine, nausea, skin problems, sores, ulcers, urethritis. 17,18 For asthma, chest colds, earaches, headaches, sores, strains, tumors. 17,18 For chicken pox, fevers, stomachache For arthritis, asthma, bruises, burns, constipation, diabetes, earaches, headaches, malnutrition, migraines, nephritis, paralysis, respiratory infections, rheumatism, sprains, swelling, ulcers, wound, and to induce vomiting of blood. 19 For antibacterial and anti-inflammatory. 20

India For eye infections, headaches, inflammation, menstrual disorders, pimples, wounds. 17,18
Mexico For aches, burns, childbirth, colds, coughs, fever, headache, pain, respiratory infection. 17,18
Nicaragua For bacterial infections, boils, broken bones, bronchitis, cancer (lymphoma), conjunctivitis, coughs, earaches. 17,18
Peru For asthma, chest colds, earaches, headaches, sores, strains, tumors. 17,18
South America For arthritis, asthma, bruises, burns, constipation, diabetes, earaches, headaches, malnutrition, migraines, nephritis, paralysis, respiratory infections, rheumatism, sprains, swelling, ulcers, wound, and to induce vomiting of blood. 19
USA For antibacterial and anti-inflammatory. 20
Elsewhere For fever, headache, heartburn, inflammation, intestinal problems, migraine, nausea, skin problems, sores, ulcers, urethritis. 17,18
Vietnam For asthma, chest colds, earaches, headaches, sores, strains, tumors. 17,18
by reducing the release, synthesis, and/or production of inflammatory cytokines and mediators such as prostaglandins, histamine, polypeptide kinins, and others.25

1.13. Uterine contractility
The phytotherapeutic tocolytic action of in human myometrium in vitro against the standard betamimetic, fenoterol Contractility was assessed in strips of term myometrium biopsied at caesarean section and subjected to escalating concentrations of B.Pinnatum vs +/- oxytocin 1 U/l in 14 women. The suppression of spontaneous contraction in the result state was concentration dependant. B. Pinnatum raised contraction frequency by 91% at constant amplitude while inhibiting oxytocin-stimulated contractions by 20% at constant amplitude but with a somewhat lower frequency. Fenoterol reduced contraction by 50% with a substantial reduction in frequency.26

1.14. Wound-healing activity
The ethanolic extract of K. pinnata demonstrated substantial wound-healing activity by reducing the size of the affected region and oedema at the wound site. This may be because steroidal glycosides and phenolic antioxidants are present.27 Water, petroleum ether, and alcoholic extracts of the plant were shown to have the ability to cure wounds in a research. Water extract was likewise shown to be more active than the other two extracts in the research.28

1.15. Antioxidant activity
The leaves were reported to show maximum scavenging effects than stems and the ethanolic extract showed more total phenolic and flavonoid content than other extracts. The high amount of phenols and flavonoids in the extracts may be the reason for their high antioxidative activity (Bhatti et al., 2012). The phenolic constituents have the ability to interact with the transition metals even in lipid phase and chelate them by filling their aqua-coordination sites and generating metal-coordinated insoluble complexes. Inhibition of lipid auto-oxidation could be attributed to the ability of phenolics to stabilise the radicals through generation of stabilised phenoxyl radicals by directly scavenging peroxyl radicals.29

1.16. Antitumour activity
Experiments were carried out on mice by causing tumour development in the peritoneal cavity. The plant’s methanolic and aqueous extracts were used as medicines in precise doses. As a tumor-suppressing drug, these extracts reduced the amount of ascitic fluid and halted tumour development. As a result, the extracts were said to have antitumor action.30

The plant’s anticancer properties was investigated. The intake of the aqueous extract for N-diethylnitrosamine (DENA)-induced hepatocarcinogenesis in rats reduced liver damage, according to the findings. The protective effect may be attributed to antioxidant and antiperoxidative properties, as well as the capacity to rectify anomalies in lipid and lipoprotein metabolism by increasing the activity of a few lipid metabolising enzymes. DENA's metabolism in the liver tends to produce free radicals, disrupting antioxidant status and eventually contributing to oxidative stress and carcinogenesis, which is why it’s classified as a significant environmental hepatocarcinogen. Histopathological study of DENA-treated rats’ liver sections revealed severe centrilobular necrosis and vacuolisation. The aqueous extract neutralised free radicals, decreased necrosis, and protected hepatocytes against DENA's carcinogenic effects.31

1.17. Antiviral activity
The anticancer and anti-HPV properties of the plant’s chloroform extract were investigated in a research. When the extract fractions were applied to cancer cell lines, they inhibited the production of viral proteins, limiting both viral and tumour development. The Epstein-Barr virus is a herpes virus that infects human B-lymphocytes, causing tumours to develop32

1.18. Antimicrobial & antifungal activity
The antibacterial activity of K. pinnata was investigated, and it was shown that the methanolic extract had a higher inhibition rate. Bacteria that cause skin infections may also cause respiratory illnesses, food poisoning, wound infections, abscesses, osteomyelitis, endocarditis, pneumonia, and other problems when they enter the body. As a result, the produced extract may fight such illnesses and save the lives of those who are afflicted. The information provided may be utilised to create a commercial antibacterial and antifungal cream.33

1.19. Antileishmanial activity
Leishmaniasis is caused by protozoans of the genus leishmania. Mice infected with leishmania amazonensis were administered an aqueous extract of K. pinnata. Following the experiment, a few findings were recorded, including the lesions shrinking in size and the parasitical load in the affected region. Continuous treatment with the extract not only slowed the development of the bacteria, but also stopped them from spreading. This technique may potentially be used to treat visceral leishmaniasis, according to some.34
1.20. Insecticidal activity

The methanolic extract of K. pinnata yielded two bufadienolides. The inclusion of the 1, 3, 5-orthoacetate moiety of the bufadienolides was shown to be responsible for the isolated compounds’ significant insecticidal action against silkworm third instar larvae. 35

1.21. Antipyretic activity

The impact of plant extract on hyperthermic conditions in experimental animals was shown. Brewer’s yeast was used to cause pyrexia in rats. When K. pinnata hydroalcoholic extract was given to laboratory specimens, it lowered body temperature, demonstrating its antipyretic activity. This action may be due to the presence of flavonoids in the extract. 36

1.22. Antilithiatic activity

Calcium oxalate stones develop when oxalate excretion in the urine is decreased. On the basis of medical prophylaxis, fresh juice produced from the leaves of K. pinnata was given to patients with stones in their bodies. The juice, when consumed on a regular basis, successfully dissolved the stones, regardless of their location, nature, or prior treatments. The amount of urine voided increased, demonstrating the juice’s diuretic properties. It also aided in the reduction of oxalate excretion while boosting citrate excretion. According to this research, the juice may have antilithiatic effects. 3

2. Neuropharmacological

The aqueous leaf mine has a sedative effect on the central nervous system (CNS). The CNS treatment of rats with 50–200 mg/kg was shown to cause a significant reduction in locomotory activity in a dose-dependent manner, with no ptosis at these dosages. Similarly, there was a significant defeat of management and reduction in muscular tone in animals given intraperitoneally with aqueous extract in a dose-dependent manner in the chimney, ascending, and inclined screen tests. The findings show significant changes in universal behaviour patterns, a reduction in impulsive lethality, and a dose-dependent potentiation of pentobarbitone-induced sleeping duration. 37

3. Immunosuppressive

In mice, an aqueous extract of grasses produced a substantial reserve of cell-mediated and humoral immunological responses. The spleen cells of mice pre-treated with herbal extract had a limited capacity to proliferate in vitro in response to both mitogen and antigen. By almost completely eradicating the Delayed-type hypersensitivity reaction, the in vitro and topical techniques of directing were the most effective. The intraperitoneal and verbal methods, respectively, decreased the response by 73 percent and 47 percent of controls. The precise antibody responses to ovalbumin were also summarized in a meaningful way by handling. As a result, the immunosuppressive properties of the aqueous extract of leaves are preserved. The lymphoproliferative tests were used to categories the immunosuppressive components found in Panphuti. A cleaned fraction (KP12SA) derived from the ethanolic extract was 20 times more effective than the crude extract in inhibiting murine lymphocyte explosion. As a consequence, the provides evidence that basil’s saturated fatty acids have a substantial effect on lymphocyte proliferation, elucidating its immunosuppressive effect in vivo. 38

![Fig. 1: Pharmacological activities of kalanchoe pinnata](image)
3.1. Antibacterial activity

Bacillus subtilis, Staphylococcus aureus, Pseudomonas aeruginosa, Klebsiella aerogenes, Escherichia coli, Klebsiella pneumoniae, and Salmonella typhi are among the bacteria that cause typhoid fever and other bacterial illnesses, according to reports. Antibacterial activities of the infusion and methanolic extracts against S. aureus ATCC 13709, E. coli ATCC 9637, Bacillus, P. aeruginosa, K. pneumonia, and S. typhi were investigated using the agar diffusion process, as well as adjacent to S. aureus, E. coli, S. typhi, Klebsiella spp., and P. aeruginosa using. \(^{39,40}\)

3.2. Cytotoxic to cattle

The effects of cardiac glycoside poisoning in calves fed flower heads of hybrid Bryophyllum species were studied, and it was discovered that for each plant (except Bryophyllum tubiflorum), two calves were given a single dosage of 20 g wet weight per kg body weight.

The findings of the calf toxicity experiment, together with the quantities of bufadienolide found in the plants, indicate that brzyotoxins A, B, and C are most likely to blame for the illness. \(^{41}\)

3.3. Herbal tonic

Ascorbic acids, riboflavin, thiamine, and niacin are all abundant in the plant. Natural ascorbic acid is essential for body function, which includes the proper synthesis of intercellular components such as collagen, bone matrix, and tooth dentine throughout the body. As a result, the clinical symptoms of scurvy, such as bleeding from the mouth, gastrointestinal tract, anaemia, and joint aches, may be linked to the relationship between ascorbic acid and proper connective tissue metabolism. \(^{42}\)

3.4. Cytotoxicity of testis

The cytotoxic impact of an ethanolic extract of B. pinnatum leaf on cells of the rat testis was investigated at two dosages (100 mg/kg and 200 mg/kg) given orally for eight weeks. When compared to the control group, which showed intact normal histological features of the testes, the seminiferous tubules shrank and intracellular spaces were seen within the epithelium at a dose of 100 mg/kg, and a higher dose (200 mg/kg) showed a marked increase in intracellular spaces within the germinal epithelium and a reduction of spermatozoa. \(^{43}\)

4. Conclusion

The plants are well-known and have long been believed to be medicinal. It is hoped that the comprehensive information provided in this study on the plant’s phytochemical components and different biological characteristics of extracts and constituents would provide an impetus to evaluate the plant’s application in medicine. The pharmacological potentials of K. pinnata which is extremely useful to researcher to know more about this important plant.

5. Source of Funding

None.

6. Conflict of Interest

None.

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References

1. Joseph B, Priya RM, Helen P, Sujatha S. Bio-active compounds in essential oil and its effects of antimicrobial, cytotoxic activity from the Psidium guajava (L.) Leaf. J Adv Biotechnol. 2010;9:10–4.
2. Okwu DE, Nnamdi FU. Two new flavonoids from Bryophyllum pinnatum and their antimicrobial Activity. J Chem Pharm Res. 2011;3(2):1–10.
3. Gahlaut A, Pawar SD, Mandal TK, Dabur R. Evaluation of clinical efficacy of Bryophyllum pinnatum Salisbury. for treatment of lithiasis. Int J Pharm Sci. 2012;4(4):505–7.
4. Pathak D, Alam K, Rohilla H, Rai AR, Agrawal A. Phytochemical investigation of Boerhavia diffusa and Rorograph paniculata: A comparative study. Int J Pharm Sci. 2012;4:975–1491.
5. Some magnetic is land plants; 2014. Available from: http://www.somemagneticislandplants.com.au/index.php/plants/396-bryophyllum-pinnatum.
6. Bryophyllum_pinnatum; 2014. Available from: http://www.keyserver.lucidcentral.org/weeds/data/030308000607490a8d040605030c0f01/media/Html/Bryophyllum_pinnatum.htm.
7. Gurudeva MR. Botanical and vernacular names of south Indian plants; 2001. p. 245246.
8. Chunekar KC, Pandey GS. Bhavanishaska. Varanasi: Chaukhambha Bharathi Academy; 2010. p. 107.
9. Environmental Weeds of Australia; 2014. Available from: http://keyserver.lucidcentral.org/weeds/data/030308000607490a.
10. Gaind K, Gupta R, Alkanes, Alkanols T, Kalanchoe S, Pinnata. Triterpenes and sterols of kalanchoe pinnata. Phytochemistry. 1972;11:1500–2.
11. Mukul. Kalanchoe Pinnata(Patharchur); 2009. Available from: http://findmeacure.com/2009/03/25/kalanchoe-pinnata/.
12. Supratman U, Fujita T, Akiyama K, Hayashi H. New insecticidal bufadienolide, Bryophyllin C from Kalanchoe pinnata. Biosci Biotechnol Biochem. 2000;64(6):1310–2. doi:10.1271/bbb.64.1310
13. Akinpelu DA. Antimicrobial activity of Bryophyllum pinnatum leaves. Fitoterapia. 2000;71(2):193–4. doi:10.1016/s0367-326x(99)00135-5
14. Costa SS, Souza M, Ibrahim T, Melo GO, Almeida AP, Guette C, et al. Kalanchoe dimilate, an anti-inflammatory salt from Kalanchoe brasiliensis. J Nat Prod. 2006;79(5):815–8. doi:10.1021/np050477z
15. Prabal S, Mihin D, Dutta M, Dhrupad C. Documentation and Traditional Herbal Knowledge of Khamptis of Arunachal Pardesh. Indian J Tradit Knowledge. 2008;5(3):438–42.
BMC Complement Altern Med leaf on human cervical cancer cells.

et al. Anticancer property of Bryophyllum pinnata (Lam.) Oken. Int J Res Biol Sci Swiss Albino mice.

Bryophyllum calycinum Salisb. against Ehrlich ascites carcinoma in stems of Kalanchoe pinnatum. Int J Green Pharm comparison of antioxidant activities of various extracts of leaves and J Nat Remedies extract of Kalanchoe pinnata Lam. leaf-A preliminary study. J Pharm Pharmacol. 2011;5(1):83–92.

Yadav NP, Dixit VK. Hepatoprotective activity of leaves of Kalanchoe pinnata Pers. J Ethnopharmacol. 2003;86(2-3):197–202.

Adesanwo JK, Raji Y, Olaleye SB, Onasawo SA, Fadare OO, Ige OO. Antitulcer Activity of Methanolic Extract of Bryophyllum pinnatum in Rats. J Biol Sci. 2007;7(2):409–12.

Ojewole J. Antinociceptive, anti-inflammatory and antidiabetic effects of Bryophyllum pinnatum (Crassulaceae) leaf aqueous extract. J Ethnopharmacol. 2005;99(1):13–9.

Gwendenberger B, Rist L, Huch R, Mandach NV. Effect of Bryophyllum pinnatum leaf versus fenoterol on uterine contractility. Eur J Obstet Gynecol Reprod Biol. 2004;112(3):64–71.

Nassis CZ, Haebsch EM, Giesbrecht AM. Antihistamine activity of Bryophyllum Calycinum. Braz J Med Biol Res. 1992;25(9):929–36.

Ajeet Pal Singh, Amar Pal Singh, Harjinder Singh, Ameeta Singh, Rakesh K. Singh. Antimicrobial potential of ethanolic extract of Bryophyllum pinnatum on the micro anatomy of drug and pharmaceutical industries. Afr J Biotechnol. 2012;10(12):15–15.

Nayak BS, Marshall JR, Isitgor O. Wound healing potential of ethanolic extract of Kalanchoe pinnata Lam. leaf-A preliminary study. Indian J Exp Biol. 2010;48(6):572–6.

Khan M, Patil PA, Shobha JC. Influence of Bryophyllum pinnatum on uterine contractility. J Pharm Pharmacol. 2011;5(1):83–92.

Bhatti M, Kamboj A, Saluja AK, Jain UK. In vitro evaluation and comparison of antioxidant activities of various extracts of leaves and stems of Kalanchoe pinnatum. Int J Green Pharm. 2012;6:340–347.

Devbhu D, Gupta JK, Devbhu P. Studies on antitumour activity of Bryophyllum calycinum Salish. against Ehrlich ascites carcinoma in Swiss Albino mice. J Pharma Sci Tech. 2012;2(1):31–3.

Afzal M, Kazmi I, Khan R, Singh R, Chauhan M, Bisht T, et al. Bryophyllum pinnatum: A review. Int J Res Biol Sci. 2012;2(11):143–9.

Mahata S, Maru S, Shukla S, Pandey A, Mugesh G, Das BC, et al. Anticancer property of Bryophyllum pinnata (Lam.) Oken. leaf on human cervical cancer cells. BMC Complement Altern Med. 2012;10(12):15–15.

Pattewar SV, Patil DN, Dahikar SB. Antimicrobial potential of extract from leaves of Kalanchoe pinnata. Int J Pharm Sci Res. 2013;4(12):4577–80.

Muzitano MF, Falcão C, Cruz EA, Bergonzi MC, Bilia AR, Vinci M. Oral metabolism and efficacy of Kalanchoe pinnata flavonoids in a murine model of cutaneous leishmaniasis. Planta Med. 2009;75(4):307–11.

Supratman U, Fujita T, Akiyama K, Hayashi H. New insecticidal bufadienolide, bryophyllin C, from Kalanchoe pinnata. Biosci Biotechnol Biochem. 2000;64(6):1310–12.

Biswas D, Mondal TK. Evaluation of anti-pyretic activity of hydroalcoholic extract of Kalanchoe pinnata leaves against yeast-induced pyrexia in rat. Int J Innovat Pharm Sci Res. 2015;3:483–92.

Salahdeen HM, Yemitan OK. Neuropharmacological effects of aqueous leaf extract of Bryophyllum pinnatum in mice. Afr J Biomed Res. 2004;7(2):164–71.