

A review on chemical and biological properties of *Cayratia trifolia* Linn. (Vitaceae)

Dinesh Kumar, Sunil Kumar, Jyoti Gupta, Renu Arya, Ankit Gupta¹

Division of Pharmacognosy and Phytochemistry, Institute of Pharmaceutical Sciences, Kurukshetra University, Kurukshetra, ¹G.G.S. College of Pharmacy, Yamuna Nagar, Haryana, India

Submitted: 09-04-2011

Revised: 18-07-2011

Published: 23-12-2011

ABSTRACT

Cayratia trifolia Linn. Domin Syn. *Vitis trifolia* (Family: Vitaceae) is commonly known as Fox grape in English; Amlabel, Ramchana in Hindi and Amlavetash in Sanskrit. It is native to India, Asia and Australia. It is a perennial climber having trifoliate leaves with 2-3 cm long petioles and ovate to oblong-ovate leaflets. Flowers are small greenish white and brown in color. Fruits are fleshy, juicy, dark purple or black, nearly spherical, about 1 cm in diameter. It is found throughout the hills in India. This perennial climber is also found in the hotter part of India from Jammu and Rajasthan to Assam extending into the peninsular India upto 600 m height. Whole plant of *Cayratia trifolia* has been reported to contain yellow waxy oil, steroids/terpenoids, flavonoids, tannins upon preliminary phytochemical screening. Leaves contain stilbenes (piceid, reveratrol, viniferin, ampelopsin). Stem, leaves, roots are reported to possess hydrocyanic acid, delphinidin and several flavonoids such as cyanidin is reported in the leaves. This plant also contains kaempferol, myricetin, quercetin, triterpenes and epifriedelanol. Infusion of seeds along with extract of tubers is traditionally given orally to diabetic patients to check sugar level of blood. Paste of tuberous is applied on the affected part in the treatment of snake bite. Whole plant is used as diuretic, in tumors, neuralgia and splenopathy. Its climbers wrapped around the neck of frantic bullock and poultice of leaves are used to yoke sores of bullock. The bark extract shows the antiviral, antibacterial, antiprotozoal, hypoglycemic, anticancer and diuretic activity. This article focuses on the upgraded review on chemical and biological properties of *Cayratia trifolia* Linn. and triggers further investigation on this plant.

Key words: Biological, *Cayratia trifolia*, chemical, review

INTRODUCTION

According to World Health Organization, traditional medicine is defined as diverse health practices, approaches, and knowledge and believes incorporating plant, animal and/or mineral-based medicines, spiritual therapies, manual techniques and exercises applied singularly or in combination to maintain well-being as well as to treat, diagnose or prevent illness.^[1,2] More than 35,000 plant species are being used in various human cultures around

the world for medicinal purposes.^[3] Crude drugs are usually the dried parts of medicinal plants (roots, stem wood, bark, leaves, flowers seeds, fruits and whole plants, etc.) that form the essential raw materials for the production of traditional remedies in various systems of medicines like Ayurveda, Siddha, Unani, Homeopathy, Tibetan, etc.

Cayratia trifolia Linn. Domin syn. *Vitis trifolia* Linn. (Family: Vitaceae) is a native of India, Asia and Australia.^[4] It is a perennial climber, found in the hotter parts of India from Jammu and Rajasthan to Assam, Tripura and West Bengal extending into peninsular India up to 600 m.^[5,6]

Synonyms

Cayratia trifolia is also known by various synonyms^[7-11] such as:
Vitis trifolia Linn.
Cissus carnosus Lamk.
Vitis carnosus (Lamk.) Wall.ex M. Lawson
Cissus trifolia (Linn.) K. Schaum
Cayratia carnosus (Lamk.) Gagnep

Local names

Different vernacular names^[6] of *Cayratia trifolia* have been reported in Table 1.

Address for correspondence:

Dr. Dinesh Kumar,
 Division of Pharmacognosy and Phytochemistry,
 Institute of Pharmaceutical Sciences, Kurukshetra
 University, Kurukshetra-136119 Haryana, India
 E-mail: dineshbarbola@yahoo.co.in

Access this article online

Quick Response Code:



Website:
www.phcogrev.com

DOI:
 10.4103/0973-7847.91117

Taxonomical hierarchy

The taxonomical hierarchy^[4,13] of *Cayratia trifolia* has been mentioned in Table 2.

Botanical distribution

Cayratia trifolia is a weak herbaceous climber, woody at base, stem is more or less succulent, compressed and densely. Leaves are trifoliolate with petioles 2-3-cm long. Leaflets are ovate to oblong-ovate, 2-8-cm long, 1.5-5-cm wide, pointed at the tip. Flowers are small greenish white 2.5mm, and brown on solitary cymes in leaf axils.^[12,14,15] Fruits are fleshy, juicy, dark purple or black, nearly spherical and about 1 cm in diameter [Figure 1]. Seeds are triangular, apex rounded, ventral holes and ribs obtuse along margin, slightly raised.^[17]

Geographical distribution

Cayratia trifolia is known as kalit--kalit in Philippines where it is found at low altitudes. It is also found from India to southern China, through the Malaya to the Moluccas and the Caroline Islands. It also found throughout the hilly regions in India.^[17,18] This perennial climber also grows wildly in Jammu, Rajasthan, Assam, Tripura and West Bengal extending into peninsular India up to 600 m.^[6] This plant is also distributed in Bangladesh, Burma, Ceylon, Combia, Indonesia, Laos, Makaysia, Malacca, Pakistan, Thailand and Vietnam.^[19,20] It is found in tropical and subtropical areas of Asia, Africa, Australia and Island of the Pacific Ocean.^[21]

Chemical constituents

This plant also contains kaempferol, myricetin, quercetin, triterpenes and epifriedelanol.^[22] Whole plant of *Cayratia trifolia* has been reported to contain yellow waxy oil, steroids/terpenoids, flavonoids, tannins.^[6] Leaves contain stilbenes such as piceid, reveratrol, viniferin and ampelopsin.^[35] Stem, leaves and roots are reported to possess hydrocyanic acid and delphinidin. Several flavonoids such as cyanidin are reported in the leaves.^[23,24] Its seeds and fruits showed presence of cyanogenic compounds. Fruits also contain calcium oxalate responsible for severe irritation in the mouth.^[6]

Ethnomedicinal uses

Whole plant is used as diuretic and is also useful in tumors, neuralgia and splenopathy, leucorrhea,^[6,16] astringent.^[31] Leaves, root and seeds are used as poultice to ulcers and boils.^[8,14,16] Fermentation of hot decoction of leaves and root is used as diaphoretic^[8] and recommended in high fever^[16]. Sap of stems and juice of leaves are used as aphrodisiac.^[23] Root is used to reduce anemic condition, stomachic diseases, as an astringent^[27] and paste as an antidote in snake bite, also in complained of carencules.^[3,6,25,26] Extract of tuber along with infusion of *Cayratia trifolia* seeds is given orally to diabetic patients to check sugar level of blood whereas powder of tuberous root is taken orally with the milk for the early recovery of fractured bone.^[25,26] Leaves are Rubifacient,^[29] used to stop bleeding of injuries.^[29,30] Root bark reduces the muscular pain.^[31]

Table 1: Vernacular names of *Cayratia trifolia*

Language	Vernacular names
Assam	<i>Ghepeta-lat, Chepeta lota</i>
Bengali	<i>Amla-lata</i>
English	<i>Fox-grape</i>
Gujrati	<i>Khat-khatumbo</i>
Hindi	<i>Amal-bel, Ramchana, Teen panya kand, Amar chatioo, khkata-limba, Tamnaya, Gidardrak</i>
Karnataka	<i>Heggoli</i>
Malyalam	<i>Sorivalli</i>
Marthi	<i>Ambat-vel</i>
Punjabi	<i>Armal-bel</i>
Sanskrit	<i>Amlavetash, Atyamlaparni, Gandiran.</i>

Table 2: Taxonomical hierarchy of *Cayratia trifolia*

Taxonomical hierarchy	Names
Domain	<i>Eukaryota</i>
Subkingdom	<i>Viridaeplantae</i>
Kingdom	<i>Plantae</i>
Phylum	<i>Tracheophyta</i>
Subphylum	<i>Euphylllophytina</i>
Infraphylum	<i>Radiatopses</i>
Class	<i>Magnoliopsida</i>
Subclass	<i>Rosidae</i>
Suborder	<i>Vitanae</i>
Order	<i>Vitales</i>
Family	<i>Vitaceae</i>
Subfamily	<i>Vitoideae</i>
Genus	<i>Cayratia</i>
Species	<i>trifolia</i>



Figure 1: Major phytochemical constituents of *H. spinosa*

Therapeutic uses

Paste of *Cayratia trifolia* is applied locally by the tribal's for early cure of wounds and edema^[9,25]. Roots are grounded with black peeper and applied as poultice on boils.^[15,29,31] Root paste is mixed with coconut oil and applied as decoction for 3 days.^[29] Leaf paste of *Gymnema sylvestris* and *Cayratia trifolia* is applied locally in eczema.^[32]

Pharmacological uses

The 50% ethanolic extract of the plant (excluding root) in a preliminary biological screening showed gross behavioral effect and hypothermia. The bark extract showed 40-59.9% inhibition of potato virus. The plant is reported to have antibacterial, antifungal, antiprotozoal, hypoglycemic, anticancer and diuretic actions.^[6]

Veterinary uses

Poultice of leaves are used for yoke sores of bullock and also used to cure swelling, injury and infection.^[16,29,31] Climbers are wrapped around the neck of a frantic bullock.^[17,31]

Non-medicinal uses

Fruits are edible, pleasantly acidic in taste.^[8] Stem bark is used to make net and ropes.^[34]

IN VIVO AND IN VITRO RESEARCH AND PHARMACOLOGICAL ACTIONS

Antioxidant activity

The powdered plants were continuously extracted with petroleum ether, chloroform, ethyl acetate and methanol. The crude extract of ethyl acetate and methanol were tested for their biological activity including antioxidant activity by scavenging effect on DPPH (1,1-diphenyl-2-picryl hydraryl) radicals. The crude extract of *Cayratia trifolia* showed the ED₅₀ values of 10.24 and 11.36 g/ml, respectively.^[36]

Antimicrobial activity

Crude extract of this plant was tested in preliminary biological screening for their antimicrobial activity against *Escherichia coli*, *Bacillus subtilis*, *Micrococcus luteus* and *P. oxalium*. Pre-cleaned extract was also investigated for their ability to inhibit protein kinase and tyrosine-specific protein kinase of epidermal growth factor.^[37]

Anticancer activity

A large variety of phytochemical constituents that have been reported from natural product research has been proven successfully as anticancerous agent. The finding from the study reveals that methanolic extract is more potent than aqueous extract in exerting antineoplastic effect in both cell lines as evident by a dose dependent decrease in cell growth. The effect was analysed at different concentration level ranging from 50 to 500 µg/ml. Delphinidin and cyaniding which are anthocyanin and showed antiproliferative and proapoptotic properties in gastric adenocarcinoma and were also found to be protective against esophageal cancer in rodents.^[38]

Neuroprotective effect

The dietary supplementation with resveratrol significantly reduced plaque formation in animal brains, a component of Alzheimer and other neurodegenerative disease.^[39] In mice, oral resveratrol produced large reductions in brain plaque in the hypothalamus(-90%), Striatum(-89%) and redial cortex (-48%) section of the brain in humans. In humans it is theorized that oral doses of resveratrol may reduce β-amyloid plaque associated with aging changes in the brain.^[40-43]

Anti-inflammatory effect

In a rat model of carrageenan-induced Paw edema, resveratrol inhibited both acute and chronic phases of the anti-inflammatory process.^[44]

Cardioprotective effects

- It inhibits the vascular cell adhesion molecular expression.^[45,46]
- Inhibition of vascular smooth muscle cell proliferation^[47-49]
- Stimulation of endothelial nitric oxide synthase activity^[50-52]
- Inhibition of platelet aggregation^[53-55]

Antidiabetic effect

It possesses hypoglycemic and hypolipidemic effect in both Streptozotacin-induced diabetes rats and STZ-Nicotinamide-induced diabetes rats. Other diabetic animal model studies by different researches have also demonstrated the antidiabetic effect of resveratrol.^[56-61]

Antiviral effect

It inhibits herpes simplex virus types 1 and 2 replication by inhibition of an early step in virus replication cycle. *In vivo* studies in mice shows that resveratrol inhibits or reduce HSV replication in the vagina and limits extra-vaginal disease.^[62-64] Studies also show that resveratrol inhibits varicella-Zoster virus, certain influenza viruses, human cytomegalovirus. Furthermore, resveratrol synergistically enhances the anti-HIV-1 activity of several anti-HIV drugs.^[65-70]

CONCLUSIONS

Cayratia trifolia Linn. is a medicinally important plants and used in the treatment of various diseases in Indian system of medicine. This paper provides valuable information about plant. Such information may serve as a base for new pharmacognostical, phytochemical, pharmacological, toxicological and clinical research.

ACKNOWLEDGMENTS

The authors are thankful to UGC, New Delhi for financially supporting the study [F.No. 39-955/2010].

REFERENCES

1. Lewington A. Medicinal plants and plant Extracts: A review of their importation into Europe. Cambridge, UK: Traffic

- International; 1993.
2. Maurya R, Gupta CM. Traditional herbs for modern medicine. *Tech Monitor* 2006;43:23-36.
 3. WHO. Quality control guidelines for medicinal plant materials. Geneva: World Health Organization; 2008.
 4. Purushothama S, Viswanath S, Kunhikannan C. Economic valuation of extractive conservation in a tropical deciduous forest in Madhya Pradesh, India. *J Trop Eco* 2001;41:61-72.
 5. The Wealth of India: A Dictionary of India Raw Material and Industrial Products. New Delhi, India: CSIR; 2004;3:399-400.
 6. Gupta AK, Sharma M. Review on Indian Medical Plants. New Delhi, India: ICMR; 2007;5:879-82.
 7. Sesagiriravuru R. Flora of Srikakulam district, Andhra Pradesh. *J Ind Bot Soc India* 1986;147.
 8. Nazimuddin S, Qaiser M. Flora of Pakistan. Available from: <http://www.efolra.org>. 17.
 9. Chen Z, Ren H, Wen J. Vitaceae, Flora of China. (Beijing) and Missouri Bot. Garden (USA): Sci Press; 2010;12:33,115,173.
 10. Drury H. Handbook of Indian flora: Being a guide to all flowering plants. Trivandrum, India: Travancore Sircar Press 1864; 1: 175.
 11. Chaudhary AB. Forest Plants of Eastern India. New Delhi, India: Ashish Publishing House; 1993. p. 180.
 12. Garden CA, Bennet HW. The Toxic Plant of Western Australian Path. *West Aust News Paper*; 1956.
 13. Bradacs G. Ethnobotanical survey and Biological screening of medical plants from Vanuatu Ph.D. dissertation. Germany: University of Regensburg; 2008. p. 171.
 14. Vardana R. Direct use of medical plant and their identification. New Delhi, India: SARUP and Sons; 2008;1:177.
 15. Pulliah T. Encyclopedia of World Medical Plants. India: Regency; 2006;1:492.
 16. Gaur RD, Sharma J. Plants Used in Traditional Healthcare of live stock by Gujjar community of Sub Himalayan tracts, Uttarakhand, India. *IJNPR* 2010;2:243-8.
 17. Tutul E, Uddin MD. Z. Angiospermic flora of Ructia Sal Forest (Bangladesh). *Bangladesh J Plant Taxon.* 2010;17:33-45.
 18. Manjuhatta BK, Krishna V, Pullaiah T. Flora of Davanagere district, Karnataka, India: Regency; 2004. p. 94.
 19. Soejima A, Wen J. Phylogenetic analysis of Grape Family (Vitaceae) based on three chloroplast markers. *Am J Bot* 2005;93:278.
 20. Lee CC, Houghton P. Cytotoxicity of plants from Malaysia and Thailand used traditionally to treat Cancer. *J Ethnopharmacol* 2005; 237-43.
 21. Defilippis AR, Maina LS. The Palauan and Yap Medical Plant Studies of Masayoshi okabe. *Atoll Res Bull* 1988:17.
 22. Munchen, Staatssamml. Protabase Record Display 1953;1:352.
 23. Grubben GJ, Denton OA. Plant Resources of Tropical Africa. Vol. 2:Vegetables. Backhuys. 2004. p. 166.
 24. Throton WBC. Krakatau. Harvard University Press. 1997: 121, 155.
 25. Choudhary K, Singh M. Ethnobotanical Survey of Rajasthan-An Update. *Am Euras J Bot* 2008;1:38-45.
 26. Swarnkar S, Katewa SS. Ethnobotanical observation on tuberous plants from tribal areas of Rajasthan (India). *Ethnobotanical leaflets* 2008. p. 647,660.
 27. Khare CP. Indian Medicinal Plants. An illustrated of Dictionary. Springer: Verlag, Berlin; 2007. p. 132.
 28. Board N. Compendium of Medical Plant. Delhi, India: National Institute of Industrial Research; 2005. p. 43.
 29. Patil VM. Ethnobotany of Nasik District, Maharashtra. Delhi, India: Daya Publishing House; 2006. p. 103,119,340,386,413.
 30. Azam MN, Hassan AI, Ismal M, Islam MN, Haque MZ, Jahan R, et al. An Ethanopharmacological survey of Daulatdia ghat area, Kushtia District (Bangladesh), used for treatment of "HARD To CURE" diseases. University of Ottawa, Canada: OGIRC/CICMR Second Joint Conference; 2010. p. 43.
 31. Patil DA, Pawar S. Ethnobotany of Jalgaon District, Maharashtra. New Delhi, India: Daya Publishing House; 2006. p. 100,486,513,516,549.
 32. Jain A, Katwal SS. Some therapeutic Uses of Biodiversity among the tribals of Rajasthan. *Indian J Tradit Knowl* 2008;7:256-62.
 33. Rahmatullah M, Mollik MA, Jilani MA, Hossain MA, Hossain SH, Rahman MM, et al. Medicinal plant used by folk medicinal practitioners in three villages of Natore and Rajshahi district, Bangladesh. *Adv Nat Appl Sci* 2010;4:132-8.
 34. Ayyanar M, Lgnacimuthu S. Plants used for non-medicinal purposes by the tribal people in Kalakad, Mundan-Thurai Tiger Reserve Southern India. *Indian J Tradit Knowl* 2010;9:515-8.
 35. Arora J, Roat C, Goyal S, Ramawat KG. High Stilbenes accumulation in root culture of *Cayratia trifolia* (L.) Domin grown in shake flask. *Acta Physiol Plant* 2009;31:1307-11.
 36. Homhua S, Tonggok P, Bonjim J. Evaluation of Biological activities of Crude Extracts from *Cratogeomys formosum* (Jack.) Dyer. and *Cayratia trifolia* L. Domin young shoots. *J Ubon Rajathanee Uni* 2007;9:54-60.
 37. Nick A. Biological Screening of Traditional Medicinal Plant from Papua New Guinea and subsequent phytochemical investigation of *Dillenia Papuana*, Doctoral Theses (Diss. ETH No. 11231). Zürich: Swiss Federal Institute of Technology; 1995. p. 2-5.
 38. Rejitha G, Das A. Cytotoxic effect of *Cayratia camosa* leaves on Human Breast Cancer Cell Lines. *Int J Cancer Res* 2009;5:115-22.
 39. Anekonda TS. Resveratrol-A Boon for treating Alzheimer's disease? *Brain Res Rev* 2006;52: 316-26.
 40. Sharma M, Gupta YK. Chronic treatment with trans resveratrol prevents intracerebroventricular streptozotocin induced cognitive impairment and oxidative stress in rats. *Life Sci* 2002;71:2489-98.
 41. Kumar P, Padi SS, Naidu PS, Kumar A. Effect of resveratrol on 3-nitropropionic acid-induced biochemical and behavioural changes: Possible neuroprotective mechanisms. *Behav Pharmacol* 2006;17:485-92.
 42. Yang YB, Piao YJ. Effects of resveratrol on secondary damages after acute spinal cord injury in rats. *Acta Pharmacol Sin* 2003;24:703-10.
 43. Sinha K, Chaudhary G, Gupta YK. Protective effect of resveratrol against oxidative stress in middle cerebral artery occlusion model of stroke in rats. *Life Sci* 2002;71:655-65.
 44. Gentilli M, Mazoit JX, Bouaziz H. Resveratrol decreases hyperalgesia induced by carrageenan in the rat hind paw. *Life Sci* 2001;68:1317-21.
 45. Ferrero ME, Bertelli AE, Fulgenzi A, Pellegatta F, Corsi MM, Bonfrate M, et al. Activity *in vitro* of resveratrol on granulocyte and monocyte adhesion to endothelium. *Am J Clin Nut* 1988;68:1208-14.
 46. Rotondo S, Rajtar G, Manarini S, Celardo A, Rotilio D, Gaetano G de, et al. Effect of trans-resveratrol, a natural polyphenolic compound, on human polymorphonuclear leukocyte function. *Br J Pharmacol* 1998;123:1691-9.
 47. Haider UG, Roos TU, Kontaridis MI, Sorescu BD, Griendling KK, Vollmar AM, et al. Resveratrol inhibits angiotensin II- and epidermal growth factor-mediated Akt activation: Role of Gab1 and Shp2. *Mol Pharmacol* 2005;68:41-8.
 48. Wang Z, Chen Y, Labinskyy N, Hsieh TC, Ungvari Z, Wu JM. Regulation of proliferation and gene expression in cultured human aortic smooth muscle cells by resveratrol and standardized grape extracts. *Biochem Biophys Res Commun* 2006;346:367-76.

49. Poussier B, Cordova AC, Becquemin JP, Sumpio BE. Resveratrol inhibits vascular smooth muscle cell proliferation and induces apoptosis. *J Vasc Surg* 2005;42:1190-7.
50. Duffy SJ, Vita JA. Effects of phenolics on vascular endothelial function. *Curr Opin Lipidol* 2003;14:21-7.
51. Wallerath T, Deckert G, Ternes T, Anderson H, Li H, Witte, K, *et al.* Resveratrol, a Polyphenolic Phytoalexin Present in Red Wine, Enhances Expression and Activity of Endothelial Nitric Oxide Synthase. *Circulation* 2002;106:1652-8.
52. Chen CK, Pace-Asciak CR. Vasorelaxing activity of resveratrol and quercetin in isolated rat aorta. *Gen Pharmacol* 1996;27:363-6.
53. Olas B, Wachowicz B. Resveratrol, a phenolic antioxidant with effects on blood platelet functions. *Platelets* 2005;16:251-60.
54. Stef G, Csiszar A, Lerea K, Ungvari Z, Veress G. Resveratrol inhibits aggregation of platelets from high-risk cardiac patients with aspirin resistance. *J Cardiovasc Pharmacol* 2006;48:1-5.
55. Wang Z, Zou J, Huang Y, Cao K, Xu Y, Wu JM. Effect of resveratrol on platelet aggregation *in vivo* and *in vitro*. *Chin Med J* 2002;115:378-80.
56. Lagouge M, Argmann C, Gerhart-Hines Z, Meziane H, Lerin C, Daussin F, *et al.* Resveratrol improves mitochondrial function and protects against metabolic disease by activating SIRT1 and PGC-1 α . *Cell* 2006;127:1109-22.
57. Deng JY, Hsieh PS, Huang JP, Lu LS, Hung LM. Activation of Estrogen Receptor Is Crucial for Resveratrol-Stimulating Muscular Glucose Uptake via Both Insulin-Dependent and Independent Pathways. *Diabetes* 2008;57:1814-23.
58. Palsamy P, Subramanian S. Resveratrol, a natural phytoalexin, normalizes hyperglycemia in streptozotocin-nicotinamide induced experimental diabetic rats. *Biomed Pharmacother* 2008; 62:598-605.
59. Sharma S, Anjaneyulu M, Kulkarni SK, Chopra K. Resveratrol, a polyphenolic phytoalexin, attenuates diabetic nephropathy in rats. *Pharmacology* 2006;6:69-75.
60. Schmatz R, Mazzanti CM, Spanevello R, Stefanello N, Gutierrez J, Corrêa M, *et al.* Resveratrol prevents memory deficits and the increase in acetylcholinesterase activity in streptozotocin-induced diabetic rats. *Eur J Pharmacol* 2009;610:42-8.
61. Penumathsa SV, Thirunavukkarasu M, Zhan L, Maulik G, Menon VP, Bagchi D, *et al.* Resveratrol enhances GLUT-4 translocation to the caveolar lipid raft fractions through AMPK/Akt/eNOS signalling pathway in diabetic myocardium. *J Cell Mol Med* 2008;12:2350-61.
62. Docherty JJ, Fu MM, Stiffler BS, Limperos RJ, Pokabla CM, DeLucia AL. Resveratrol inhibition of herpes simplex virus replication. *Antiviral Res* 1999;43:145-55.
63. Docherty JJ, Fu MM, Hah JM, Sweet TJ, Faith SA, Booth T. Effect of resveratrol on herpes simplex virus vaginal infection in the mouse. *Antiviral Res* 2005;67:155-62.
64. Docherty JJ, Smith JS, Fu MM, Stoner T, Booth T. Effect of topically applied resveratrol on cutaneous herpes simplex virus infections in hairless mice. *Antiviral Res* 2004;61:19-26.
65. Docherty JJ, Sweet TJ, Bailey E, Faith SA, Booth T. Resveratrol inhibition of varicella-zoster virus replication *in vitro*. *Antiviral Res* 2006;72:171-7.
66. Guan WD, Yang ZF, Liu N, Qin S, Zhang FX, Zhu YT. *In vitro* experimental study on the effect of resveratrol against several kinds of respiroviruses(in Chinese). *Zhong Yao Cai* 2008;31:1388-90.
67. Palamara AT, Nencioni L, Aquilano K, De Chiara G, Hernandez L, Cozzolino F, *et al.* Inhibition of influenza A virus replication by resveratrol. *J Infect Dis* 2005;191:1719-29.
68. Li YQ, Li ZL, Zhao WJ, Wen RX, Meng QW, Zeng Y. Synthesis of stilbene derivatives with inhibition of SARS coronavirus replication. *Eur J Med Chem* 2006;41:1084-9.
69. Evers DL, Wang X, Huong SM, Huang DY, Huang ES. 3,4',5-Trihydroxy-trans-stilbene (resveratrol) inhibits human cytomegalovirus replication and virus-induced cellular signaling. *Antiviral Res* 2004;63:85-95.
70. Heredia A, Davis C, Redfield R. Synergistic inhibition of HIV-1 in activated and resting peripheral blood mononuclear cells, monocyte-derived macrophages, and selected drug-resistant isolates with nucleoside analogues combined with a natural product, resveratrol. *J Acquir Immune Defic Syndr* 2000;25:246-55.

How to cite this Article: Kumar D, Kumar S, Gupta J, Arya R, Gupta A. A review on chemical and biological properties of *Cayratia trifolia* Linn. (Vitaceae). *Phcog Rev* 2011;5:184-8.

Source of Support: UGC, New Delhi for financially supporting the study [F.No. 39-955/2010]., **Conflict of Interest:** None declared