

Review

Pharmacology and phytochemistry of *Coccinia indica*

U. A. Deokate* and S. S. Khadabadi

Government College of Pharmacy, Kathora Naka, Amravati-444604, (M. S.), India.

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***Coccinia indica* (Bimba, kanduri, Cucurbitaceae) is famous for its hypoglycemic and antidiabetic properties in Ayurvedic system of medicine. Other applications include the therapy of various conditions such as skin diseases and gonorrhoea. The present review highlights the phytochemistry and pharmacology of *Coccinia indica*. There are many patented formulations derived from *C. indica* which are now distributed increasingly all over the world. This has given rise to a concomitant increase in research on the phytochemical constituents and biological activity of *C. indica*.**

Key words: *Coccinia indica*, hypoglycemic, antidiabetic, saponins, terpenoids, sterol.

INTRODUCTION

Indian system of traditional knowledge that is, ayurveda is well known for its effective herbal treatments. There are about 7000 plant species are found in India. Although most of them have a long history in folk medicine, there is lack of written data on their efficacy and safety, esp. from human studies. Many of them are used to treat highly prevalent disorder diabetes mellitus (Mankil et al., 2006). Few of this common examples includes *Allium cepa*, *Allium sativum*, *Aloe vera*, *Coccinia indica*, *Caesalpinia bonducella*, *Eugenia jambolana*, *Mucuna pruriens*, *Murraya koeingii*, *Mormodica charantia*, *Swertia chirata*, *Syzygium cumini*, *Tinospora cordifolia* and *Trigonella foenum-graecum*. The present review highlights the phytochemistry and pharmacology of one of popular antidiabetic and hypoglycemic plant *C. indica*.

C. indica (Synonym: *Coccinia grandis*, *Coccinia cordifolia*) family Cucurbitaceae commonly called little gourd or Rantondli in Marathi, Bimba in Sanskrit and Kandutikibel in Hindi. It is indigenous to Bengal and other parts of India. *C. indica* grows abundantly all over India, Tropical Africa, Australia, Fiji and throughout the oriental countries. The plant has also been used extensively in Ayurvedic and Unani practice in the Indian subcontinent (Wealth of India, 1992). It has long tuberous fleshy roots, smooth and green fruits. Microscopy of root shows parenchyma, phelloderm, pericyclic fibers, stone cells, starch grains. TS of leaves show upper and lower

epidermis, ranunculaceae stomata, uniseriate multicellular trichomes.

PHARMACOLOGICAL REVIEW

Fresh juice of roots is used to treat diabetes; tincture of leaves is used to treat gonorrhoea, paste of leaves is applied to the skin diseases. Dried bark is a good cathartic. Leaves and stem are antispasmodic and expectorant. The fleshy green fruit is very bitter. Green fruit is chewed to cure sores on the tongue [Gupta and Variyar, 1964; Khan et al., 1980; Chandrasekar et al., 1989; Shibib et al., 1993; Platel et al., 1997].

Table 1 clears that this plant is exhaustively studied for its hypoglycemic, antidiabetic potential with different animal models from 1953 (Shakya, 2008; Venkateswaran et al., 2003; Venkateswaran and Pari, 2003; Yeh et al., 2003; Wasantwisut and Viriyapanich, 2003; Ajay et al., 2009; Mallick et al., 2009; Nahar et al., 1998). Indigenous people use various parts of the plant to get relief from diabetes mellitus. *C. indica* leaves showed that it depressed the activity of the enzyme glucose-6-phosphatase and possesses an antioxidant activity, which may be attributed to its protective action on lipid peroxidation and to the enhancing effect on cellular antioxidant defence contributing to the protection against oxidative damage in streptozotocin diabetes. Hypolipidemic activity was also studied but that was also in diabetic rats. Many clinical trial studies has also proven effectiveness and safety of this plant parts and derived

*Corresponding author. E-mail: deokate@yahoo.com.

Table 1. Pharmacological review of plant *C. indica*.

S/No.	Activity	Model	Plant part	Remark
1	Antidiabetic activity (Hossain et al., 1992)	Alloxan diabetic albino rats	95% ethanolic extracts	Found to be active.
2	Antidiabetic activity (Shakya, 2008).	Streptozotocin included diabetic rats	n-hexane extract	Found to be active.
3	Antidiabetic activity with testicular disorders (Mallick et al., 2007).	Streptozotocin Induced Diabetic Rat For Testicular Dysfunctions	Formulation Of <i>Musa paradisiacal</i> , <i>Tamarindus indica</i> , <i>Eugenia jambolana</i> and <i>Coccinia indica</i>	Found to be active.
4	Antidiabetic activity (Pari and Venkateswaran, 2003)	Normal and streptozotocin (STZ) diabetic rats	Leaves	Evaluated for effect on blood glucose, plasma insulin, cholesterol, triglycerides, free fatty acids, and phospholipids and fatty acid compound of total lipids in liver, kidney and brain.
5	Antidiabetic activity (Mukerji, 1953)	Alloxan diabetes in rabbits	Roots	Found to be active.
6	Antidiabetic activity (Kamble et al., 1998).	Normal and Streptozotocin-induced male diabetic rats	Leaves	Lowered blood glucose by depressing its synthesis, on the one hand though depression of the key gluconeogenic enzymes glucose-6-phosphatase and fructose-1,6- biphosphatase and on the other by enhancing glucose oxidation by the shunt pathway through activation of its principal enzymes G6PDH.
7	Hypoglycemic activity (Gupta, 1963)	Normal rats	Pectin isolated from the fruit	Glycogen synthetase activity was highly significant significant redn. in phosphorylase activity.
8	Hypoglycemic activity (Brahmachari and Augusti, 1963)		Water soluble Alkaloid fraction	Found to be active.
9	MOA of hypoglycemic activity (Kumar et al., 1993).	Glucose tolerance test	Alcoholic extarct of <i>Coccinia indica</i> (100 mg/kg.),	May be due to indirect stimulation of insulin secretion or to retardation of glucose absorption. Use of these drugs may prevent deterioration of pancreatic lesion in 8diabetics.
10	Hypoglycemic activity (Mukherjee et al., 1972).	Rabbits	Alcoholic and aqueous extract of root powder	Found to be active.
11	Clinical trial in type 2 diabetic patients (Azad et al., 1979).	Double- blind, placebo-controlled, randomized trial	Alcoholic extract of the herb	Have potential hypoglycemic action in patients with mild diadetes

Table 1. Contd.

12	Clinical trial in diabetic patients (Kuriyan et al., 2008).		Dried extract of whole plant	Ingredients present in the extract act like insulin, correcting the elevated enzymes G-6-P (ase), LDH in glycolytic pathway and restore the LPL activity in lipolytic pathway with the control of hyperglycemia in diabetes.
13	Antidiabetic activity (Singh et al., 1985)	Dog	Dried extract of whole plant	Found to be active.
14	Antioxidant activity (Venkateswaran and Pari, 2003).	Streptozotocin-diabetic rats	Ethanol extract of leaves	Found to be active.
15	Anti-inflammatory activity (Rao et al., 2004)	Carrageenin and histamine induced paw edema	fruit juice powder	Found to be active.
16	Antinociceptive activity (Rao et al., 2004).	Writhing induced by acetic acid in mice	Fruit juice powder	Found to be active.
17	Post- and pre- treatment anti-inflammatory activity (Niazi et al., 2009).	Carrageenan-induced paw oedema method	Aqueous extract of fresh leaves	Found to be active.
18	Analgesic activity (Niazi et al., 2009).	Tail flick model in rats	Aqueous extract of fresh leaves	Found to be active.
19	Antipyretic activity (Niazi et al., 2009).	Yeast- induced hyperpyrexia in rats	Aqueous extract of fresh leaves	Found to be active.
20	Larvicidal activity (Rahuman and Venkatesan, 2008).	Early fourth instar larvae of <i>Aedes aegypti</i> L. and <i>Culex quinquefasciatus</i> (say) (Diptera: Culicidae).	Hexane, ethyl acetate, petroleum ether, acetone and methanol extracts of the leaf <i>Citrullus colocynthis</i> , <i>Coccinia indica</i> , <i>Cucumis sativus</i> , <i>Momordica charantia</i> , and <i>Trichosanthes anguina</i> ,	Found to be active.
21	Hypolipidemic activity (Kumar et al., 1997).	Streptozotocin-diabetic rats	Ethanol extract of leaves.	Found to be active.
22	Hepatoprotective activity (Rao et al., 2003)	CCl ₄ induced hepatotoxicity in rats	Ethanol extract of fruits	Found to be active.
23	Antituberculosis activity (Mukerji and Gupta, 1958).	Experimental tuberculosis in Guinea pigs	Extract of fruit	No effect found.

Table 2. Phytochemical review of plant *C. indica*.

Plant part	Constituent reported
Roots (Vaishnav et al., 2001; Vaishnav and Gupta, 1996; Vaishnav and Gupta, 1995; Khastgir et al., 1958; Sucrow and Reimerdes, 1968)	Triterpenoid, saponin coccinoside – k(i). $C_{41}H_{66}O_{12}$ Flavonoid glycoside ombuin 3-o- arabinofuranoside 3- o- β - (α -l- arabinopyranosyl)-(1→2) – β -d-glucopyranosyl- (1→3)- β - hydroxylup – 20(29)- en-28- oic acid. Lupeol, β -amyrin, and β - sitosterol. Stigmast -7- en-3-one,
Fruits (Kundu and Ray, 1987; Basu and Ghosh, 1972; Bhakuni et al., 1962)	Taraxerone, taraxerol, and (24R)-24- ethylcholest- 5- en- 3 β - ol glucoside. B- carotene, lycopene, cryptoxanthin, and apo- 6'- lycopenal B- sitosterol and taraxerol
Aerial parts (Qudrat-i-Khuda et al., 1965; Dhargalkar and Guha, 1959)	Heptacosane Cephalandrol, $C_{29}H_{58}O$ tritriacontane $C_{33}H_{68}$ B- sitosterol alkaloids Cephalandrine a and Cephalandrine b.
Whole plant (Rahman et al., 1990)	Aspartic acid, Glutamic Acid, Asparagine, Tyrosine, Histidine, Phenylalanine And Threonine Valine Arginine

**Figure 1.** *C. indica* fruits.

formulations for antidiabetic effect. Anti-inflammatory, analgesic and antipyretic activity of fruit and leaves were studied and found to be significant.

**Figure 2.** *C. indica* leaves.

PHYTOCHEMICAL REVIEW

Plant contains saponins, flavonoids, sterols and alkaloids which are summarized in Table 2.

Saponin and flavonoid are found to be responsible for antidiabetic activity (Figures 1 and 2).

CONCLUSION

C. indica is a famous plant for its safe antidiabetic property. It proved the insulin stimulatory effect of *C. indica* leaves from existing b-cells in diabetic rats. It possesses hypoglycemic, antidiabetic, hypolipidemic, hepatoprotective, larvicidal, anti-inflammatory, analgesic and antipyretic activities. It is found to be devoid of antituberculosis properties. Various phytoconstituents reported in *C. indica* are cephalandrol, tritriacontane, lupeol, b-sitosterol, cephalandrine A, cephalandrine B, stigma-7-en-3-one, taraxerone and taraxerol. Terpenoids are found to be responsible for antidiabetic activity. Despite the broad use of *C. indica* in traditional medicine, very few systematic pharmacological and phytochemical studies are reported till date assessing its therapeutic properties.

REFERENCES

- Ajay SS, Bhaskar VH, Teotia D (2009). Quality standardization of *Coccinia indica* fruit. *Asian J. Chem.*, 21(7): 5351-5354.
- Azad KAK, Akhtar S, Mahtab H (1979). *Coccinia indica* in the treatment of patients with diabetes mellitus. *Bangladesh Med. Res. Council Bull.*, 5(2): 60-66.
- Basu K, Ghosh BK (1972). Chemical investigation of *Coccinia indica*. *Transactions of the Bose Res. Inst. (Calcutta)*. 35(2): 43-44.
- Bhakuni DS, Srivastava SN, Sharma VN, Kaul KN (1962). Chemical examination of the fruits of *Coccinia indica*. *J. Sci. Ind. Res.*, Section B: *Phys. Sci.*, 21B: 237-238.
- Brahmachari HD, Augusti KT (1963). Orally effective hypoglycemic principles from *Coccinia indica*. *J. Pharm. Pharmacol.*, 15(6): 411-412.
- Chandrasekar B, Mukherjee B, Mukherjee SK (1989). Blood sugar lowering potentiality of selected Cucurbitaceae plants of Indian origin. *Indian J. Med. Res.*, 90: 300-305.
- Dhargalkar IM, Guha SK (1959). Nutritional values of Indian vegetables. *J. Proc. Inst. Chemists*, (31): 109-112.
- Gupta SS, Variyar MC (1964). Experimental studies on pituitary diabetes. Iv. Effect of *Gymnema sylvestre* and *Coccinia indica* against the hyperglycaemic response of somatotropin and corticotropin hormones. *Indian J. Med. Res.*, 52: 200-207.
- Gupta SS (1963). Pituitary diabetes III. Effect of indigenous antidiabetic drugs against the acute hyperglycemic response of anterior pituitary extract in glucose fed albino rats. *Indian J. Med. Res.*, 51(4): 716-724.
- Hossain MZ, Shibib BA, Rahman R (1992). Hypoglycemic effects of *Coccinia indica*: Inhibition of key gluconeogenic enzyme, glucose-6-phosphatase. *Indian J. Exp. Biol.*, 30(5): 418-420.
- Kamble SM, Kamalakar PL, Vaidya S, Bambole VD (1998). Influence of *Coccinia indica* on certain enzymes in glycolytic and lipolytic pathway in human diabetes. *Indian J. Med. Sci.*, 52(4): 143-146.
- Khan AK, Akhtar S, Mahtab H (1980). Treatment of diabetes mellitus with *Coccinia indica*. *Brit. Med. J.*, 280(6220): 1044.
- Khastgir HN, Choudhuri SN, Gupta PS (1958). Roots of *Coccinia indica*. *J. Indian Chem. Soc.*, 35: 905-906.
- Kumar GP, Sudheesh S, Vijayalakshmi NR (1993). Hypoglycemic effect of *Coccinia indica*: Mechanism of action. *Planta Med.*, 59(4): 330-332.
- Kumar GP, Sudheesh S, Ushakumari B, Valsa AK, Vijayakumar S, Sandhya C, Vijayalakshmi NR (1997). A comparative study on the hypolipidemic activity of eleven different pectins. *J. Food Sci. Technol.*, 34(2): 103-107.
- Kundu S, Ray AB (1987). Chemical examination of *Coccinia indica* fruits. *J. Indian Chem. Soc.*, 64(12): 776-777.
- Kuriyan R, Rajendran R, Bantwal G, Kurpad AV (2008). Effect of supplementation of *Coccinia cordifolia* extract on newly detected diabetic patients. *Diabetes Care*, 31(2): 216-220.
- Mallick C, De Debasis, Ghosh D (2009). Correction of protein metabolic disorders by composite extract of *Musa paradisiaca* and *Coccinia indica* in streptozotocin-induced diabetic albino rat: An approach through the pancreas. *Pancreas*, 38(3): 322-329.
- Mallick C, Mandal S, Barik B, Bhattacharya A, Ghosh D (2007). Protection of testicular dysfunctions by MTEC, a formulated herbal drug in streptozotocin induced diabetic rat. *Biol. Pharm. Bull.*, 30(1): 84-90.
- Mankil J, Moonsoo P, Hyun CL, Yoon-Ho K, Eun SK, Sang KK (2006). Antidiabetic Agents from Medicinal Plants. *Curr. Med. Chem.*, 13: 1203-1218.
- Mukerji B, Gupta SK (1958). Indigenous drugs in experimental tuberculosis. *Chemotherapy Proceedings Symposium Lucknow*, pp. 90-101.
- Mukerji B (1953). Effect of *Coccinia indica* on alloxan diabetes in rabbits. *Indian J. Med. Sci.*, 7: 665-672.
- Mukherjee K, Ghosh NC, Datta T (1972). *Coccinia indica* as a potential hypoglycemic agent. *Ind. J. Exp. Biol.*, 10(5): 347-349.
- Nahar N, Mosihuzzaman M, Khan MD Shahinul H (1998). Determination of free sugars in plant materials having antidiabetic activity. *Dhaka Univ. J. Sci.*, 46(1): 167-170.
- Niazi J, Singh P, Bansal Y, Goel RK (2009). Anti-inflammatory, analgesic and antipyretic activity of aqueous extract of fresh leaves of *Coccinia indica*. *Inflammopharmacol.*, 17(4): 239-244.
- Pari L, Venkateswaran S (2003). Protective effect of *Coccinia indica* on changes in the fatty acid composition in streptozotocin induced diabetic rats. *Pharmazie*, 58(6): 409-412.
- Platel K, Srinivasan K (1997). Nahrung Plant foods in the management of diabetes mellitus. Vegetables as potential hypoglycemic agents. *41(2): 68-74.*
- Qudrat-i-Khuda M, Khaleque KA, Miah MAW (1965). Chemical investigations on *Cephalandra indica*. I. Constituents of dry aerial parts. *Dacca Sci. Res.*, (Dacca, Pakistan), 2(1/2): 27-231.
- Rahman M, Mahbubur C, Tofail A, Mosihuzzaman M (1990). Analysis of water- and alkali-soluble polysaccharides of *Coccinia indica* (Telakucha) plant. *J. Bangladesh Chem. Soc.*, 3(2): 199-204.
- Rahuman AA, Venkatesan P (2008). Larvicidal efficacy of five cucurbitaceous plant leaf extracts against mosquito species. *Parasitol. Res.*, 103(1): 133-139.
- Rao GM, Mohana VM, Rao CV, Rawat AKS, Mehrotra S (2003). Hepatoprotective effect of *Coccinia indica* against CCl₄ induced hepatotoxicity. *Nat. Prod. Sci.*, 9(1): 13-17.
- Rao GM, Rao V, Sudhakara M, Pandey MM, Rawat AKS, Sirwaikar A, Joshi AB (2004). Anti-inflammatory and antinociceptive activities of "*Coccinia indica* WA." fruit juice powder in animals. *Nat. Prod. Sci.*, 10(1): 20-23.
- Shakya VK (2008). Antidiabetic activity of *Coccinia indica* in streptozotocin induced diabetic rats. *Asian J. Chem.*, 20(8): 6479-6482.
- Shakya VK (2008). Antidiabetic activity of *Coccinia indica* in streptozotocin induced diabetic rats. *Asian J. Chem.*, 20(8): 6479-6482.
- Shibib BA, Khan LA, Rahman R (1993). Hypoglycaemic activity of *Coccinia indica* and *Momordica charantia* in diabetic rats: Depression of the hepatic gluconeogenic enzymes glucose-6-phosphatase and fructose-1,6-bisphosphatase and elevation of both liver and red-cell shunt enzyme glucose-6-phosphate dehydrogenase. *Biochem. J.*, 292(1): 267-270.
- Singh N, Singh SP, Vrat S, Misra N, Dixit K, Kohli RP (1985). A study on the anti-diabetic activity of *Coccinia indica* in dogs. *Ind. J. Med. Sci.*, 39(2): 27-29.
- Sucrow W, Reimerdes A (1968). 7-Sterols from Cucurbitaceae. *Biologie*, 23(1): 42-45.
- Vaishnav MM, Jain P, Jogi SR, Gupta KR (2001). Coccinioside-K, triterpenoid saponin from *Coccinia indica*. *Oriental J. Chem.*, 17(3): 465-468.
- Vaishnav MM, Gupta KR (1995). A new saponin from *Coccinia indica* roots. *Fitoterapia*, 66(6): 546-547.
- Vaishnav MM, Gupta KR (1996). Ombuin 3-O-arabinofuranoside from *Coccinia indica*. *Fitoterapia*, 67(1): 80.
- Venkateswaran S, Pari L (2003). Effect of *Coccinia indica* leaf extract on plasma antioxidants in streptozotocin-induced experiment diabetes in rats. *Phytother. Res.*, 17(6): 605-608.
- Venkateswaran S, Pari L (2003). Effect of *Coccinia indica* leaves on antioxidant status in streptozotocin-induced diabetic rats. *J. Ethnopharmacol.*, 84(2-3): 163-168.
- Venkateswaran S, Pari L, Suguna L, Chandrakasan G (2003). Modulatory effect of *Coccinia indica* on aortic collagen in streptozotocin-induced diabetic rats. *Clin. Exp. Pharmacol. Physiol.*, 30(3): 157-163.
- Wasantwisut E, Viriyapanich T (2003). Ivy gourd (*Coccinia grandis* Voigt, *Coccinia cordifolia*, *Coccinia indica*) in human nutrition and traditional applications. *World Rev. Nutr. Dietetics*, (91): 60-66.
- Wealth of India (1992). A dictionary of Indian raw materials and India products. *Raw Material*, New Delhi, 4, p. 312.
- Yeh GY, Eisenberg DM, Kaptchuk TJ, Phillips RS (2003). Systematic review of herbs and dietary supplements for glycemic control in diabetes. *Diabetes Care*, 26(4): 1277-1294.