

Cucurbitane-Type Triterpenoids from the Blood Glucose-Lowering Extracts of *Coccinia indica* and *Momordica balsamina* Fruits

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ABSTRACT

Context: Few vegetables that are commonly consumed in India as part of diet have been claimed for their antidiabetic potential. **Objective:** The present study was aimed at evaluating preventive effects of cucurbit vegetables namely, *Coccinia indica* and *Momordica balsamina* belonging to family *Cucurbitaceae* in diabetic hyperglycemia. **Materials and Methods:** The fruits of *M. balsamina* and *C. indica* were extracted with chloroform and fractionated with hexane to prepare an extract rich in moderately polar components. These extracts were used for evaluating the effect of these cucurbits in nicotinamide/streptozotocin-induced type 2 diabetes. Streptozotocin–nicotinamide-induced diabetic animals were orally treated with chloroform extract of fruits (250 mg/kg BW) given daily for a week separately. **Results:** Both the extracts reduced fasting blood glucose significantly ($P < 0.05$ versus diabetic control) when estimated on seventh day of treatments. Pretreatment with fruit extracts for 7 days also blunted the OGTT (oral glucose tolerance test) curve. Results indicated that *C. indica* and *M. balsamina* fruits possess beneficial effects in diabetes by lowering elevated blood glucose level. Six cucurbitane-type triterpenoids were isolated from bioactive extracts of *C. indica* (1-3) and *M. balsamina* (4-6). The structures of these compounds were elucidated on the basis of spectroscopic data analysis. **Conclusion:** The study concludes that the inclusion of *C. indica* and *M. balsamina* fruits in food can be useful for newly diagnosed diabetic patients or high risk group of population for glycemic control.

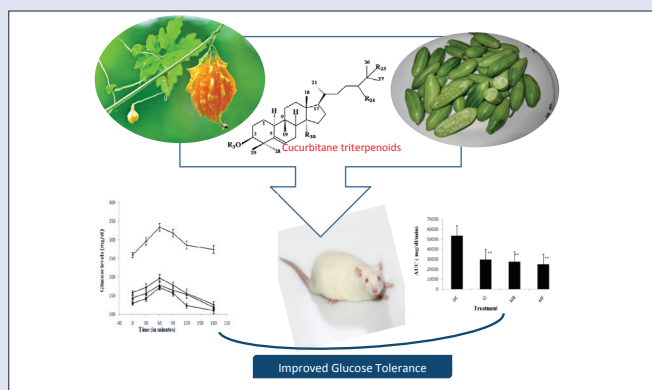
Key words: *Coccinia indica*, *Momordica balsamina*, cucurbitacins, hyperglycemia, streptozotocin, nicotinamide

SUMMARY

“Cucurbitane-type triterpenoids from the blood glucose-lowering extracts of *Coccinia indica* and *M. balsamina* fruits”

- The beneficial effects of chloroform extracts of vegetal cucurbits namely *C. indica* (Ivy gourd) and *M. balsamina* (Balsam apple) in streptozotocin-nicotinamide (STZ-NA)-induced diabetic rats has been evaluated.
- The isolation and characterization of six cucurbitacins from bioactive extracts of *C. indica* (Coccinoside A, B, and C) and *M. balsamina* (cucurbit-5, 7-dien-3 β -ol, cucurbita-5-en-3 β -ol-3-O- β -d-glucopyranoside, and cucurbit-5-en-3 β -ol-3-O- β -d-glucopyranosyl-(4'→1'')-O- β -d-glucopyranoside) have been reported for the first time.

- The study concludes that the inclusion of *C. indica* and *M. balsamina* fruits in food can be useful for newly diagnosed diabetic patients or high risk group of population for glycemic control.



Abbreviation used: C: *indica* (*Coccinia indica*), M: *balsamina* (*Momordica balsamina*), Kbr: Potassium bromide, FTIR: Fourier transform infrared spectroscopy, COSY: Correlated Spectroscopy, DEPT: Distortionless Enhancement by Polarization Transfer, DMSO: Dimethyl sulfoxide, TMS: tetramethylsilane, ESI-MS: Electrospray Ionization mass spectrometry, TLC: thin layer chromatography, STZ-NA: Streptozotocin-nicotinamide, CMC: carboxy methyl cellulose, BW: body weight, ANOVA: analysis of variance, AUC: area under curve

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INTRODUCTION

Diabetes mellitus is a serious metabolic illness worldwide. Defect in the carbohydrate metabolism and continuous efforts of physiological systems to correct this imbalance have been reported to pose overexertion on the endocrine system leading to the deterioration of endocrine control. Variation in carbohydrate metabolic enzymes due to deterioration of endocrine control leads to hyperglycemia.^[1] Some risk factors associated with diabetes such as elevated plasma glucose concentrations in the fasting state and after an oral glucose load, overweight, and sedentary lifestyle are potentially reversible. At present, most of the diabetes-related research involves drug therapy rather than preventive or nondrug interventions.^[2] In a study, incidence of diabetes was reduced by 58% with the lifestyle intervention and 31% by metformin.^[3] Few vegetables that are commonly consumed in India as part of diet have been claimed their antidiabetic potential especially those from family

Cucurbitaceae. As a part of lifestyle modification, dietary intervention focused on consumption of food plants can be an interesting approach in ameliorating the effects of this metabolic disorder to certain extent.

Cucurbitaceae is a family of about 110 genera and 640 species found abundantly in tropics and subtropics. Its members are mostly herbs

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climbing by tendrils with abundant sap and a very rapid growth. The main genera include *Cucurbita* (five species), *Cucumis* (25 species), *Citrullus* (three species), *Luffa* (six species), *Bryonia* (four species), and *Momordica* (45 species). The blood glucose-lowering efficacy of few members of *Cucurbitaceae* family has been assessed by researchers worldwide as reviewed by us previously.^[4]

Coccinia indica Wight and Arn. has long been traditionally used in diabetes care.^[5,6] Phytoconstituents reported from *C. indica* include lupeol, β -amyrin, cucurbitacins, taraxerol, β -carotene,^[7-9] steroidal saponins,^[10] flavonoids,^[11] pectin,^[12] and polyphenol.^[13] *Momordica balsamina* Linn. is a highly reputed medicinal herb used especially in the management of hyperglycemia in diabetes.^[14,15] Phytoconstituents such as momordicin^[16] and cucurbitanes^[17-19] have been reported earlier from the plants.

The current study involves the evaluation of beneficial effects of chloroform extracts of vegetable cucurbits namely *C. indica* (Ivy gourd) and *M. balsamina* (Balsam apple) in streptozotocin-nicotinamide (STZ-NA)-induced diabetic rats. In this model, diabetes is induced by administration of STZ-NA combination at low doses.^[20,21] The model is ideal for screening of dietary herbs against moderate hyperglycemia. The article also describes the isolation and characterization of six cucurbitacins from bioactive extracts of *C. indica* (1-3) and *M. balsamina* (4-6), being reported for the first time from these plants.

MATERIALS AND METHODS

General

Melting points were determined on a Perfit apparatus and are uncorrected. The infrared (IR) spectra were measured in KBr (potassium bromide) pellets on a Bio-Rad FT-IR spectrometer (SpectraLab Scientific Inc. 38 McPherson St. Markham (Greater Toronto Area) Ontario L3R 3V6 Canada North America E-mail: Technical Support: support@spectralabsci.com). Ultraviolet (UV) spectra were obtained in methanol with a Lambda Bio 20 spectrometer (Perkin-Elmer, Rotkreuz, Switzerland). ¹H (400 MHz), ¹³C (100 MHz), COSY, and DEPT NMR spectra were recorded on Bruker spectrometer (Bruker AXS, Karlsruhe, Germany). DMSO-*d*₆ (dimethyl sulfoxide) or CDCl₃ (Sigma-Aldrich, Bangalore, India) were used as solvent and TMS (tetramethylsilane) as an internal standard. ESI-MS (electrospray ionization mass spectrometry) analyses were performed on a Waters Q-TOF Premier (Micromass MS Technologies, Manchester, United Kingdom) Mass Spectrometer. Column chromatography separation was carried out on silica gel (60–120 mesh; Merck, Mumbai, India). Preparative TLC (thin layer chromatography) was performed on silica gel (Merck, 200–300 mesh). Precoated silica gel plates (Silica gel 60 F254; Merck) were used for analytical TLC visualized by exposure to iodine and UV radiations. STZ was obtained from Sigma-Aldrich Chemicals (St. Louis, Missouri, USA). NA was obtained from Central Drug House (P) Ltd., New Delhi (India) and metformin was obtained from Franco-Indian Pharmaceuticals Pvt. Ltd., Mumbai, India. All other chemicals used were of analytical grade from S.D. Fine chemicals, Delhi, India.

Raw materials and preparation of extracts

Fresh fruits of *M. balsamina* were procured from vegetable market in Moga (Punjab) and *C. indica* fruits were obtained from local vegetable market in New Delhi, India. The fruits were authenticated by Dr. H.B. Singh, Taxonomist, National Institute of Science Communication and Information Resources (NISCAIR), New Delhi, India. Voucher specimens were deposited in NISCAIR, Delhi. Fruits were sliced, shade dried, and grounded to yield the coarse powder. The powder of dried fruits (2 kg each) was defatted with petroleum ether and then extracted with chloroform. The extract was then concentrated under reduced

pressure on a Rotavapor (Buchi, Switzerland) to yield a viscous residue (110 g for *C. indica* and 130 g for *M. balsamina*).

Antihyperglycemic activity

Male albino Wistar rats (3 months old, 150–200 g BW), raised in the animal house of Hamdard University, New Delhi, were used (Animal ethics permission number 173/CPCSEA/755). They were housed in the colony cages (five rats per cage), at ambient temperature and allowed free access to water. Animals were fed with standard rodent diet. Diabetes was induced by STZ-NA combination in overnight fasted rats. STZ was dissolved in 50 mM citric acid buffer (pH 4.5) for intraperitoneally administration (45 mg/kg BW). NA was dissolved in physiological saline and intraperitoneally administered (110 mg/kg BW), 20 min before STZ administration. One week later, mildly diabetic rats with fasting blood glucose (FBG) levels at least 150 mg/dL were selected for further study.

Subchronic antidiabetic activity

The rats were randomized into four groups comprising of five animals each. Diabetic control rats, received orally 1 mL/kg BW vehicle (0.3% CMC (carboxy methyl cellulose) in distilled water). Metformin-treated diabetic rats received metformin daily (15 mg/kg BW p.o.) in vehicle 1 week after STZ-NA treatment and continued for 7 days. *C. indica* and *M. balsamina* treated diabetic rats received chloroform extracts of respective fruits (250 mg/kg BW, p.o.) in vehicle, 1 week after STZ-NA treatment and continued daily for 7 days. Blood was withdrawn from tail vein after 0, 3 and 7 days and FBG was measured.

Oral glucose tolerance test

Oral glucose tolerance test was performed after overnight fasting of rats on seventh day of treatment. Vehicle (0.3% CMC in distilled water), *C. indica* fruit extract in vehicle (250 mg/kg BW, p.o.), *M. balsamina* fruit extract in vehicle (250 mg/kg BW, p.o.), and metformin (15 mg/kg BW) were administered orally to respective groups (*n*=5). Glucose (2 g/kg BW, p.o.) was fed to all groups after 60 min of treatment. Blood was withdrawn from the tail vein at 0, 30, 60, 90, 120, and 180 min and blood glucose was measured. Blood glucose concentration was estimated using an Accu-Check active (Roche Diagnostics GmbH, Sandhoferstrasse 116, DE-68305 Mannheim, GERMANY, <http://www.roche.de/diagnostics/index.htm>), based on the glucose oxidase method.

Statistical analysis

Observed data were represented as mean \pm S.D. For statistical analysis of the data, group means were compared by one-way analysis of variance ANOVA (analysis of variance) followed by Dunnett's test for multiple comparison. A difference in the mean values of *P* < 0.05 was considered significant statistically.

Isolation of compounds

Melting points were determined on a Perfit apparatus and are uncorrected. The IR spectra were measured between 4000 and 650 cm⁻¹ in KBr pellets on a Bio-Rad FT-IR spectrometer (Spectra Lab Scientific Inc.). UV spectra were obtained in methanol from 200 to 400 nm with a Lambda Bio 20 spectrometer (Perkin-Elmer). ¹H (400 MHz), ¹³C (100 MHz), COSY, and DEPT NMR spectra were recorded on Bruker spectrometer (Bruker AXS) with acquisition and processing parameters as follows: prob hd 5 mm, pulprog zg 30, td 65536, SWH 120.230 Hz, FIDRES 0.183399 Hz, AQ 2,7263477 sec, RG 912, DW 41, 600 usec, DE 6.00 usec, TE 297.9 K, D1 1.00 sec, P1 10.90 usec, SI 32768. DMSO-*d*₆ or CDCl₃ (Sigma-Aldrich, Bangalore, India) were used as solvents, and TMS as an internal standard. ESI-MS analyses were performed on a Waters Q-TOF Premier Mass Spectrometer (Micromass MS Technologies). Column chromatography separations were carried out on silica gel (60–120 mesh; Merck). Preparative TLC was performed on silica gel (Merck,

200–300 mesh). Precoated silica gel plates (Merck, Silica gel 60 F254) were used for analytical TLC visualized by exposure to iodine and UV radiations. STZ was obtained from Sigma–Aldrich chemicals (St. Louis, Missouri, USA). NA was obtained from Central Drug House (P) Ltd., New Delhi (India) and metformin was obtained from Franco–Indian Pharmaceuticals Pvt. Ltd., Mumbai, India. All other chemicals used were of analytical grade from S.D. Fine chemicals, Delhi, India.

For isolation of the compounds each extracted viscous residue was dissolved in a minimum amount of methanol and adsorbed on silica gel (60–120 mesh) for column for preparation of slurries separately. The slurries were air-dried and chromatographed over silica gel columns individually packed in petroleum ether. Each column was eluted successively with petroleum ether, mixture of petroleum ether and chloroform (9:1, 3:1, 1:1, and 1:3), pure chloroform and finally the mixture of chloroform and methanol (99:1, 98:2, 96:4, 95:5, 97:3, and 9:1). Various fractions were collected separately and matched by TLC to check homogeneity. Similar fractions (having the same R_f values) were combined and crystallized. The isolated compounds were recrystallized to isolate compounds 1–3 (from *C. indica*) and 4–6 (from *M. balsamina*) as outlined below:

Compound 1

Elution of the column with chloroform–methanol (19:1) gave a pale yellow amorphous powder of **1**, recrystallized from chloroform–methanol (1:1); R_f 0.35 (chloroform–methanol, 9:1); m.p. 265–267°C; UV λ_{\max} (MeOH): 207 nm (log ϵ 4.1); FTIR (fourier transform infrared spectroscopy) ν_{\max} (KBr): 3391, 2919, 2850, 1735, 1639, 1538, 1465, 1378 cm^{-1} ; ^1H and ^{13}C NMR (DMSO- d_6): [Table 2]; ESI-MS m/z (rel. int.): 664 $[\text{M}]^+ \text{C}_{38}\text{H}_{64}\text{O}_9$ (5), 316 (22), 163 (35), 143 (13), 124 (32).

Compound 2

Elution of the column with chloroform–methanol (9:1) yielded a yellow powder of **2**, recrystallized from acetone–methanol (1:1); R_f 0.72 (chloroform–methanol, 9:1); UV λ_{\max} (MeOH): 218 nm (log ϵ 3.1); FTIR ν_{\max} (KBr): 3366, 2919, 2850, 1739, 1627, 1538, 1467, 1376, 1261, 1169 cm^{-1} ; ^1H and ^{13}C NMR (DMSO- d_6): [Table 2]; ESI-MS m/z (rel. int.): 676 $[\text{M}]^+ \text{C}_{39}\text{H}_{64}\text{O}_9$ (4), 316 (17), 163 (14), 126 (22), 99 (19).

Compound 3

Further elution of the column with chloroform–methanol (9:1) gave light brown powder of **3**, recrystallized from chloroform–methanol (1:1); R_f 0.53 (benzene–chloroform–methanol, 5:4:1); m.p. 270–272°C; UV λ_{\max} (MeOH): 212–nm (log ϵ 4.2); FTIR ν_{\max} (KBr): 3411, 2918, 2850, 1724, 1636, 1561, 1439, 1246, 1024 cm^{-1} ; ^1H and ^{13}C NMR (DMSO- d_6): [Table 2]; ESI-MS m/z (rel. int.): 701 $[\text{M}-\text{H}]^+ \text{C}_{41}\text{H}_{65}\text{O}_9$ (4), 316 (10), 182 (9), 163 (7), 123 (82).

Compound 4

Elution of the column with chloroform yielded a pale yellow waxy compound **4**, recrystallized from chloroform; R_f 0.58 (benzene–chloroform–methanol, 5:4:1); IR ν_{\max} (KBr): 3438, 2928, 2854, 1641, 1445, 1382, 1215, 1072, 1019, 928, 932, 758 cm^{-1} ; ^1H NMR (CDCl_3): δ 6.13 (1H, d, $J=5.6$ Hz, H-6), 5.86 (1H, d, $J=5.6$ Hz, H-7), 3.81 (1H, dd, $J=5.8, 8.9$ Hz, H-3 α), 1.28 (3H, brs, Me-28), 1.25 (3H, brs, Me-19), 1.16 (3H, d, $J=6.1$ Hz, Me-21), 1.04 (3H, brs, Me-29), 0.82 (3H, d, $J=6.6$ Hz, Me-26), 0.79 (3H, d, $J=6.3$ Hz, Me-27), 0.75 (3H, brs, Me-18), 0.70 (3H, brs, Me-30), 0.65 (3H, s, Me-28); ^{13}C NMR (CDCl_3): δ 76.30 (C-3), 143.72 (C-5), 118.89 (C-6), 135.01 (C-7), 136.92 (C-8), 16.39 (C-18), 18.65 (C-19), 26.12 (C-21), 29.48 (C-26), 27.49 (C-27), 23.61 (C-28), 32.33 (C-29), 17.59 (C-30); ESI-MS m/z (rel. int.): 426 $[\text{M}]^+ \text{C}_{30}\text{H}_{50}\text{O}$ (3).

Compound 5

Elution of the column with chloroform–methanol (9:1) afforded a yellow amorphous powder of **5**, recrystallized from chloroform–methanol (9:1); R_f 0.45 (chloroform–methanol, 1:1); UV λ_{\max} (MeOH): 211 nm; IR ν_{\max} (KBr): 3425, 3390, 3260, 2957, 2850, 1644, 1445, 1377, 1161, 1074, 1028, 887 cm^{-1} ; ^1H NMR (DMSO- d_6): δ 5.22 (1H, m, H-6), 5.18 (1H, d, $J=7.1$ Hz, H-1'), 4.85 (1H, m, H-5'), 4.85 (1H, m, H-5'), 4.40 (1H, dd, $J=7.1, 8.0$ Hz, H-2'), 3.66 (1H, dd, $J=5.5, 8.9$ Hz, H-3), 3.56 (1H, m, H-3'), 3.43 (1H, m, H-4'), 3.12 (1H, d, $J=10.5$ Hz, H₂-6'a), 3.09 (1H, d, $J=10.3$ Hz, H₂-6'b), 1.28 (3H, brs, Me-28), 1.20 (3H, brs, Me-19), 0.98 (3H, d, $J=6.6$ Hz, Me-21), 0.95 (3H, brs, Me-29), 0.81 (3H, d, $J=6.5$ Hz, Me-26), 0.78 (3H, d, $J=6.4$ Hz, Me-27), 0.74 (3H, brs, Me-18), 0.51 (3H, brs, Me-30); ^{13}C NMR (DMSO- d_6): δ 76.27 (C-3), 42.19 (C-4), 148.23 (C-5), 118.42 (C-6), 57.36 (C-17), 15.61 (C-18), 22.52 (C-19), 23.73 (C-21), 25.71 (C-26), 26.03 (C-27), 21.77 (C-28), 29.48 (C-29), 18.22 (C-30), 103.61 (C-1'), 74.51 (C-2'), 68.76 (C-3'), 65.03 (C-4'), 76.17 (C-5'), 60.85 (C-6'); ESI-MS m/z (rel. int.): 590 $[\text{M}]^+ \text{C}_{36}\text{H}_{62}\text{O}_6$ (2).

Compound 6

Elution of the column with chloroform–methanol (9:1) produced yellow crystals of **6**; R_f 0.62 (benzene–chloroform–methanol, 5:4:1); UV λ_{\max} (MeOH): 212 nm; IR ν_{\max} (KBr): 3528, 3438, 3315, 2918, 2849, 1732, 1640, 1445, 1381, 1197, 1072, 1019, 987, 932, 720 cm^{-1} ; ^1H NMR (DMSO- d_6): δ 5.29 (1H, brs, H-6), 5.78 (1H, d, $J=7.3$ Hz, H-1'), 5.73 (1H, d, $J=7.1$ Hz, H-1''), 4.81 (1H, m, H-5'), 4.76 (1H, m, H-5''), 4.60 (1H, m, H-4''), 4.52 (1H, m, H-4'), 4.49 (1H, m, H-2'), 4.46 (1H, m, H-2''), 4.38 (1H, m, H-3'), 4.33 (1H, m, H-3''), 3.98 (1H, dd, $J=5.5, 9.3$ Hz, H-3 α), 3.20 (4H, brs, H₂-6, H₂-6''), 1.31 (3H, brs, Me-19), 0.97 (3H, d, $J=6.3$ Hz, Me-21), 1.34 (3H, brs, Me-28), 0.92 (3H, brs, Me-29), 0.84 (3H, d, $J=6.2$ Hz, Me-26), 0.78 (3H, d, $J=6.3$ Hz, Me-27), 0.72 (3H, brs, Me-18), 0.63 (3H, brs, Me-30); ^{13}C NMR (DMSO- d_6): δ 76.86 (C-3), 138.72 (C-5), 120.46 (C-6), 50.51 (C-17), 16.10 (C-18), 20.58 (C-19), 23.20 (C-21), 24.39 (C-26), 25.14 (C-27), 21.35 (C-28), 29.30 (C-29), 18.13 (C-30), 103.25 (C-1'), 91.64 (C-1''); ESI-MS m/z (rel. int.): 752 $[\text{M}]^+ \text{C}_{42}\text{H}_{72}\text{O}_{11}$ (2).

RESULTS AND DISCUSSION

STZ is a well known diabetogenic agent exerting cytotoxic action on pancreatic β -cells, whereas NA is given to partially protect these cells against STZ mimicking type 2 diabetes response.^[22,23] The results of subchronic administration of chloroform extracts of fruits of *C. indica* and *M. balsamina* for a week separately to diabetic rats are given in Table 1. It was observed that untreated diabetic rats showed a consistent increase in fasting blood glucose (FBG) when estimated over a week. The fruit extracts of *C. indica* and *M. balsamina*, each given orally daily for a week at a dose of 250 mg/kg BW, showed a significant reduction in FBG ($P < 0.05$ versus diabetic control) when estimated on the seventh day of treatment. Both the extracts not only prevented reduced elevated blood glucose but kept FBG of experimental animals to near basal level comparable to metformin (15 mg/kg BW, p.o.). There was more significant decrease in FBG with metformin ($P < 0.01$). Figure 1 depicted the effect of oral glucose overload (2 g/kg BW, p.o.) in diabetic rats pretreated with *C. indica* and *M. balsamina* fruit extracts (250 mg/kg BW, p.o.) given daily for 7 days separately. The fruit extracts blunted the effect of glucose challenge as signified by about 25 mg/dL increase in peak blood glucose level after 1 h of glucose overload, compared with about 75 mg/dL increment in diabetic control animals at the same time level. Both the extracts and metformin normalized blood glucose to basal level by the third hour of glucose challenge. Figure 2 provides an easy visual comparison of the effect of extracts and metformin with respect to the area under the curve (AUC) in glucose overload-induced hyperglycemia.

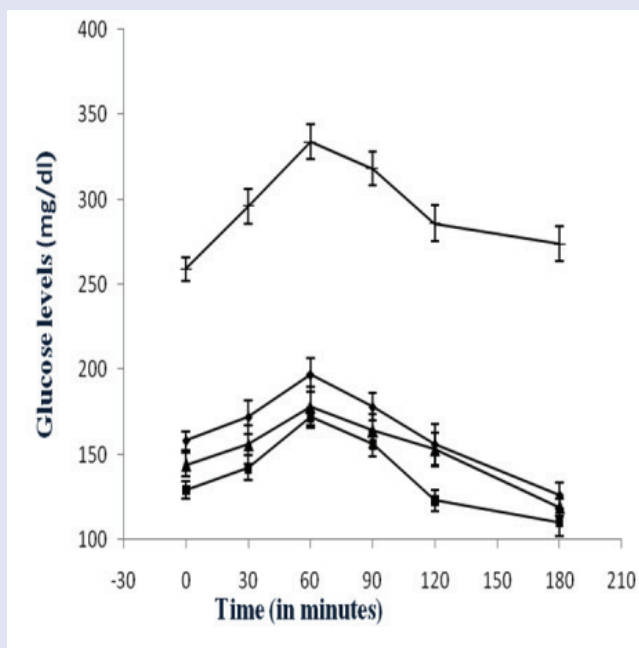


Figure 1: Effect on the glucose tolerance in mild diabetic rats after oral glucose overload.

(▲) Diabetic control (1 mL/kg BW of 0.3% CMC in distilled water, p.o.); (■) pretreated with metformin (15 mg/kg BW, p.o.) for seven days; (◆) pretreated with chloroform extract of *Coccinia indica* fruits (250 mg/kg BW) for seven days; (▲) pretreated with chloroform extract of *Momordica balsamina* fruits. Values are expressed as of mean \pm S.D.; $n = 5$ in each group.

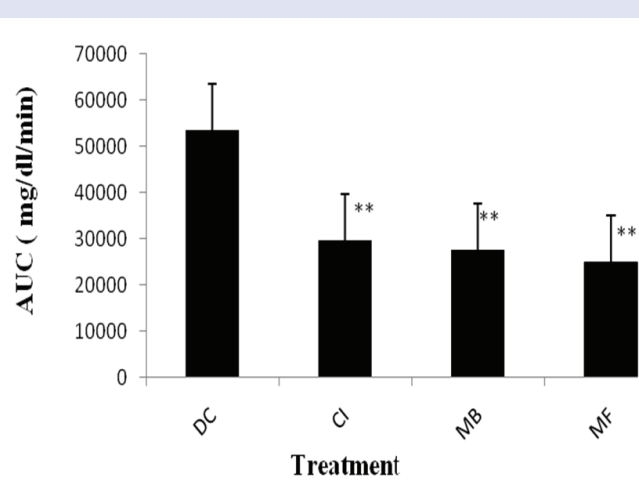


Figure 2: Representation of area under the curve (AUC) of blood glucose levels after OGTT in STZ-NA-induced mild diabetic rats.

DC: Diabetic control (1 mL/kg BW of 0.3% CMC in distilled water, p.o.); CI: pretreated with chloroform extract of *Coccinia indica* fruits (250 mg/kg BW) for seven days; MB: pretreated with chloroform extract of *Momordica balsamina* fruits (250 mg/kg BW) for seven days; MF: pretreated with metformin (15 mg/kg BW, p.o.) for seven days. Values are expressed as of mean \pm S.D.; $n = 5$ in each group. ** $P < 0.01$, significant versus DC.

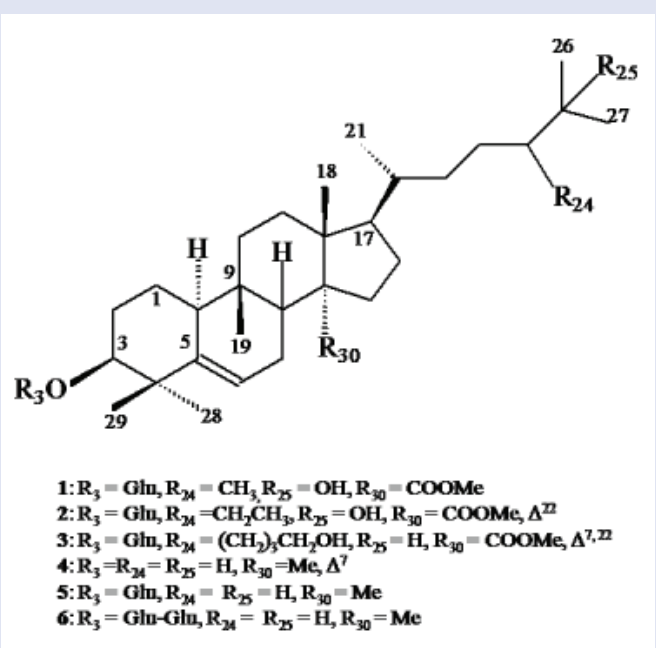


Figure 3: Chemical structures of compounds 1-6.

The bioactive extracts were subjected to column chromatography over silica to yield compounds 1-3 from *C. indica* and 4-6 from *M. balsamina*. The structures compounds [Figure 3] were elucidated on the basis of detailed spectroscopic data analysis.

Compound 1, was obtained as pale yellow amorphous powder from chloroform-methanol (19:1) eluants. Its FTIR spectrum exhibited absorption bands characteristic for hydroxyl group (3391 cm^{-1}), ester group (1735 cm^{-1}), and unsaturation (1639 cm^{-1}). Its ESI MS displayed a molecular ion peak at m/z 664 consistent with the molecular formula $\text{C}_{38}\text{H}_{64}\text{O}_9$ of a triterpenoid alglycosidic ester. The fragment ion peaks at m/z 163 [$\text{C}_6\text{H}_{11}\text{O}_5$] $^+$ supported the presence of a glucose unit in 1. The fragment ion peaks at m/z 124 ($\text{C}_{9,10}\text{-C}_{5,6}$ fission), 316 ($\text{C}_{13,17}\text{-C}_{14,15}$ fission), and 143 ($\text{C}_9\text{H}_9\text{O}$, side chain) suggested the presence of cucurbitane-type carbon framework along with a C_9 -side chain. Further evidence was provided by NMR (nuclear magnetic resonance) data [Table 2]. Signals at δ_{H} 1.23 (6H, s, $\text{H}_3\text{-26}, \text{H}_3\text{-27}$) and δ_{C} 71.32 (C-25) indicated the presence of hydroxyl at C-25. A three-proton doublet at δ_{H} 0.84 ($J = 3.3 \text{ Hz}, \text{H-24}'$) and a signal at δ_{C} 19.05 (C-24') supported the presence of methyl substituent at C-24. A double-doublet at δ_{H} 2.34 ($J = 4.6, 10.2 \text{ Hz}$) for H-10 and a slightly shielded signal at δ_{C} 36.18 for C-9 have been typically reported for cucurbitanes.^[18,19] A double-doublet at δ_{H} 3.67 ($J = 5.1, 11.2 \text{ Hz}$) was ascribed to H-3 carbinol proton placed in α -orientation on the basis of coupling constant. A singlet at δ_{H} 3.11 (3H) was assigned to methoxy protons. The downfield signals at δ_{H} 5.32 (dd, $J = 4.8, 9.4 \text{ Hz}, \text{H-6}$), δ_{C} 141.06 (C-5), and 121.31 (C-6) and indicated a double bond between C5 and C6. The signals at δ_{H} 4.90 (d, $J = 6.3 \text{ Hz}$) and δ_{C} 100.78 along with other signals for sugar protons and carbons in the respective spectra indicated the presence of a glucose unit attached to the cucurbitane unit. The COSY (correlation spectroscopy) NMR spectrum displayed an important cross peak between anomeric proton H-1' and H-3. It also showed correlations between H-10 and $\text{H}_2\text{-1}$; H-6 and $\text{H}_2\text{-7}$; H-24 and H-24'; and H-20 and $\text{H}_3\text{-21}$. The $^1\text{HNMR}$ (hydrogen-1 nuclear magnetic resonance OR Proton nuclear magnetic resonance) spectrum also displayed four singlets at δ_{H} 0.87 ($\text{H}_3\text{-18}$), 0.91 ($\text{H}_3\text{-19}$), 0.99 ($\text{H}_3\text{-28}$), and 0.74 ($\text{H}_3\text{-29}$) along with a doublet at δ_{H} 0.98 ($J = 5.5 \text{ Hz}, \text{H}_3\text{-21}$) to account for the corresponding methyl protons. The $^{13}\text{CNMR}$ = Carbon

It is evident that both the fruit extracts produced a significant reduction in AUC comparable to metformin ($P < 0.01$). Thus, the extracts of both the fruits produced significant blood glucose-lowering effect in experimental diabetic animals.

Table 1: Effect of chloroform extract of fruits of *Coccinia indica* and *Momordica balsamina* on fasting blood glucose in diabetic rats

Group	Treatment (dose)	Fasting blood glucose FBG (mg/dl)		
		T0	T3	T7
Diabetic control	Vehicle (1mL/kg BW of 0.3% CMC in distilled water, p.o)	176.2 ± 9.14	209.4 ± 11.90	259.2 ± 7.19
Positive control	Metformin (15 mg/kg BW, p.o)	161.8 ± 6.01	150.6 ± 6.69	129.4 ± 5.12**
<i>C. indica</i>	Fruit extract (250 mg/kg BW, p.o)	172.4 ± 7.09	161 ± 4.77	158.6 ± 5.41*
<i>M. balsamina</i>	Fruit extract (250 mg/kg BW, p.o)	166.4 ± 6.69	154.4 ± 7.54	144 ± 7.17*

T0 = basal glucose level, T3 = FBG after 3 days of treatment, T7 = FBG after 7 days of treatment. Values are expressed as mean ± S.D.; n = 5 in each group. *P<0.05, **P<0.01, significant treatment groups vs diabetic control.

Table 2: NMR data of compounds 1-3 (DMSO-d₆)

Position	1		2		3	
	¹ H	¹³ C	¹ H	¹³ C	¹ H	¹³ C
1	1.79 m, 1.82 m	36.80	1.77 m, 1.80 m	34.51	1.79 m, 1.83 m	35.94
2	1.78 m, 1.95 m	27.76	1.78 m, 1.91 m	27.66	1.76 m, 1.96 m	28.72
3	3.67 dd (5.1, 11.2)	76.97	3.68 dd (3.8,9.9)	76.93	3.67 dd (4.1,10.8)	77.18
4	-	41.80	-	40.41	-	40.77
5	-	141.06	-	139.39	-	140.89
6	5.32 dd (4.8,9.4)	121.31	5.57 dd (5.1,10.2)	130.01	5.44 d (6.0)	121.72
7	1.51 m, 1.98 m	26.51	1.51 m, 1.92 m	26.11	4.89 d (6.0)	121.72
8	1.95 m	49.55	1.93 m	50.12	-	140.89
9	-	36.18	-	37.23	-	36.67
10	2.34 dd (4.6, 10.2)	45.11	2.37 m	44.71	2.35 m	45.56
11	1.47 m, 1.82 m	20.55	1.47 m, 1.82 m	21.01	1.47 m, 1.86 m	20.20
12	1.33 m, 1.63 m	29.23	1.33 m, 1.52 m	28.98	1.34 m, 1.65 m	29.50
13	-	51.82	-	48.52	-	42.31
14	-	58.15	-	57.81	-	57.11
15	1.03 m, 1.54 m	30.01	1.03 m, 1.54 m	31.01	1.03 m, 1.54 m	31.87
16	1.35 m, 1.95 m	24.15	1.29 m, 1.95 m	24.37	1.36 m, 1.20 m	24.33
17	1.63 m	56.14	1.62 m	54.76	1.67 m	56.62
18	0.87 s	18.52	0.86 s	22.72	0.86 s	19.07
19	0.91 s	28.59	0.96 s	28.96	0.94 s	29.10
20	1.85 m	35.47	1.90 m	32.62	1.86 m	33.77
21	0.98 d (5.5)	11.50	0.91 d (6.4)	24.95	0.91 d (6.3)	12.13
22	2.14 m, 2.55 m	33.31	5.31 m	130.35	4.80 d (5.6)	135.65
23	1.25 m, 1.42 m	29.03	5.35 m	114.01	4.80 d (5.6)	123.93
24	1.49 m	37.54	1.48 m	34.82	1.50 m	37.28
25	-	71.32	-	32.98	1.39 m	31.18
26	1.23 s	31.37	1.23 s	30.98	0.88 s	19.57
27	1.23 s	31.37	1.23 s	30.98	0.88 s	19.37
28	0.99 s	23.83	0.96 s	27.19	0.96 s	23.04
29	0.74 s	18.82	0.83 s	19.02	0.76 s	18.79
30	-	174.51	-	176.58	-	173.51
24'	0.84 d(3.3)	19.05	1.28 m	29.72	1.24 m	29.10
25'	-	-	0.79 t (7.0)	14.16	1.26 m	29.97
26'	-	-	-	-	1.27 m	31.18
27'	-	-	-	-	4.23 d (9.8)	63.27
COOMe	3.11 s	55.36	3.16 s	50.84	3.16 s	55.85
1'	4.90 d (6.3)	100.78	4.89 d (6.2)	103.47	5.04 bd (12.1)	101.18
2'	4.38 m	73.36	4.28 m	73.11	4.32 m	73.90
3'	4.41 m	76.62	4.06 m	75.51	4.35 m	75.69
4'	4.83 m	69.98	4.17 m	71.21	4.52 m	69.65
5'	4.85 m	76.71	4.48 m	76.73	4.65 m	76.02
6'	3.48 dd (4.2, 10.8)	61.06	3.16 dd (4.5,11.1)	61.51	3.87 dd (4.7, 10.2)	63.81

Coupling constants in Hertz are presented in parentheses.

-13 nuclear magnetic resonance spectrum of **1** displayed signals for 38 carbons that were assigned unambiguously on the basis of chemical shift values and DEPT (distortionless enhancement by polarization transfer) experiments. It exhibited nine methyl carbons, 10 methylene carbons, 12 methine carbons (including six carbinol signals), and seven quaternary carbons. Importantly, the ^{13}C NMR spectrum exhibited six signals between δ_{C} 70 and 80 for the carbinol carbons out of which the one at δ_{C} 71.32 disappeared in DEPT-135 experiment, suggesting the presence of a quaternary hydroxylated carbon (C-25). These findings and comparison of NMR spectra of **1** with those of known compounds^[22-24], led us to formulate its structure as 3 β , 25-dihydroxy-24-methyl-30-carboxy methyl cucurbita-5-en-3-O- β -d-glucopyranoside and was designated as coccinoside A.

Compound **2** was obtained as yellow amorphous powder from chloroform-methanol (9:1) eluants. Its ESI mass spectrum displayed a molecular ion peak at m/z 676 consistent with the molecular formula $\text{C}_{39}\text{H}_{66}\text{O}_9$. Its spectral data, as given in Table 2, was similar to **1** except for the presence of an additional double bond and an ethyl substituent in the side-chain. This led us to formulate its structure as 3 β , 25-dihydroxy-24-ethyl-30-carboxy methyl cucurbita-5, 22-dien-3-O- β -d-glucopyranoside and was designated as coccinoside B.

Compound **3** was obtained as light brown amorphous powder from chloroform-methanol (9:1) eluants. Its IR spectrum displayed absorption bands for hydroxyl group (3411 cm^{-1}), ester group (1724 cm^{-1}), and unsaturation (1636 cm^{-1}). On the basis of mass and ^{13}C NMR spectra, its molecular formula was established as $\text{C}_{41}\text{H}_{66}\text{O}_9$ consistent with that of a triterpenoidal glycosidic ester. A diagnostic ion peak at m/z 182 [$\text{C}_{12}\text{H}_{12}\text{O}$] $^+$ supported the presence of a hydroxybutyl substituted side chain in **3**. The fragments ion peaks at m/z 123 ($\text{C}_{9^{10}}\text{-C}_{5^{16}}$ fission), 316 ($\text{C}_{13^{17}}\text{-C}_{14^{15}}$ fission), and 163 (glycosidic linkage fission) indicated a cucurbitane carbon framework linked to glucose unit. Further evidence in support of proposed structure were derived from NMR data [Table 2]. The signals at δ_{H} 1.27 (2H, m, $\text{H}_2\text{-24}'$), 1.24 (4H, m, $\text{H}_2\text{-25}'$, $\text{H}_2\text{-26}'$), 4.23 (2H, d, $J=9.8\text{ Hz}$, $\text{H}_2\text{-27}'$), and δ_{C} 31.81 (C-24'), 29.87 (C-25', C-26'), 63.27 (C-27') indicated the presence of a hydroxybutyl unit C-24 of the side-chain. Two one-proton doublets at δ_{H} 5.44 ($J=6.0\text{ Hz}$) and 4.89 ($J=6.0\text{ Hz}$), and a two-proton doublet at δ_{H} 4.80 ($J=5.6\text{ Hz}$) were ascribed correspondingly to H-6, H-7, H-22, and H-23 vinylic protons. Relatively lower values for coupling constants for vinylic protons (6.0 and 5.6 Hz) indicated the *cis*-arrangement for H6/H8 and H22/H23. Four downfield signals at δ_{C} 140.89 (C-5, C-8), 121.72 (C-6, C-7), 135.65 (C-22), and 123.93 (C-23) supported the presence of double bonds between C5-C6, C7-C8 and C22-C23. A double-doublet at δ_{H} 3.67 ($J=4.1, 10.8\text{ Hz}$) was assigned to H-3 carbinol proton placed in α -orientation on the basis of coupling constant values. A sharp singlet at δ_{H} 3.16 was attributed to methoxy protons. The signals at δ_{H} 5.04 (bd, $J=12.1\text{ Hz}$, H-1') and δ_{C} 101.18 (C-1') along with other sugar protons and carbons in the respective spectra further supported the presence of a glucose attached to the cucurbitane unit. A signal at δ_{H} 2.35 (m, H-10) and a slightly shielded signals at δ_{C} 36.67 (C-9) and 45.11 (C-10) are characteristic to cucurbitanes as reported earlier.^[17,18] The ^1H NMR spectrum also displayed five singlets at δ_{H} 0.86 ($\text{H}_3\text{-18}$), 0.94 ($\text{H}_3\text{-19}$), 0.88 ($\text{H}_3\text{-26}$, $\text{H}_3\text{-27}$), 0.96 ($\text{H}_3\text{-28}$), 0.76 ($\text{H}_3\text{-29}$), and a doublet at δ_{H} 0.91 (3H, $J=6.3\text{ Hz}$, $\text{H}_3\text{-21}$) for the methyl protons. The ^{13}C NMR spectrum **3** exhibited signals for 41 carbons that were assigned unambiguously on the basis of chemical shift values and DEPT experiment [Table 2]. It displayed eight methyl signals, 11 methylene signals, 15 methine carbons, and seven quaternary carbon signals. The important signals appeared for ester carbon at δ_{C} 173.51 (C-30), hydroxymethyl carbons at δ_{C} 63.27 (C-27'), 63.81 (C-6'), and methoxy carbon at δ_{C} 55.85. The data were compared with previous reports on cucurbitanes and were found to be in good

agreement.^[24-26] Thus, the structure of **3** was determined as 3 β -hydroxy-24-(1-butanol-4-yl)-30-carboxy methyl cucurbita-5, 7, 22-trien-3-O- β -d-glucopyranoside, and was designated as coccinoside C.

Compounds **4-6** were obtained from *M. balsamina* from chloroform and chloroform-methanol (9:1) eluants. Their spectral data were compared with previous reports on cucurbitanes and were found to be in good agreement.^[23-27] This led us to formulate their structures as cucurbit-5, 7-dien-3 β -ol (**4**), cucurbita-5-en-3 β -ol-3-O- β -d-glucopyranoside (**5**), and cucurbit-5-en-3 β -ol-3-O- β -d-glucopyranosyl-(4' \rightarrow 1'')-O- β -d-glucopyranoside (**6**). This is the first report of these cucurbitacins from these plants.

CONCLUSION

Our study confirms the beneficial effects of *C. indica* and *M. balsamina* fruits in diabetes by lowering the elevated blood glucose level. The beneficial effect of these fruits could possibly be because of the constituents reported here. The study further supports their supplementation in the diet of newly detected diabetic patients for stricter glycemic control.

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Conflicts of interest

There are no conflicts of interest.

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