Antidiabetic Activity of Hydromethanolic Extract of Crude *Dorstenia barnimiana* Root: Validation of In Vitro and In Vivo Antidiabetic and Antidyslipidemic Activity

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Background: Despite modern therapeutic armamentaria, DM remains a 21st-century public health menace. Novel phytomedicines are a rapidly expanding focus of research. The juice of *Dorstenia barnimiana* roots has long been used for the treatment of diabetes mellitus in traditional Ethiopian medicine, but its efficacy has not been supported by in vitro or in vivo scientific study. To investigate this, the present work was performed.

Methods: In this experimental study, simple random sampling was applied. Healthy male mice were used in normoglycemic and oral glucose-tolerance test (OGTT) models. Streptozotocin (IP, 150 mg/kg)–administered diabetic male mice were utilized. Animals were randomly divided into five groups of six each. Group I received 10 mL/kg distilled water, groups II–IV received 100 (DB100), 200 (DB200), and 400 (DB400) mg/kg crude extract, respectively, and group V received glibenclamide 5 mg/kg. A sham group (group VI) was added that received 10 mL/kg distilled water. All treatments were given orally. FBG, serum-lipid profiles, and body-weight changes were then measured. In vitro α -amylase inhibitory activity was also evaluated.

Results: The doses were atoxic up to 2,000 mg/kg. There was α -amylase inhibition activity of 67.52% at 500 μ g/mL with an IC_{50} of 4.595 μ g/mL. The OGTT revealed an antihyperglycemic effect of the crude extract. This was not attributed to a hypoglycemic side effect. In the diabetic mouse model, it shrank FBG levels remarkably. There were also significant reductions in serum TC, TGs, VLDL-C, and LDL-C. Nevertheless, HDL-C and body-weight levels returned.

Conclusion: The present study confirmed the safety and promising in vivo antidiabetic and antidyslipidemic activity of *D. barnimiana*, thus corroborating the traditional claim.

Keywords: antidiabetic, antidyslipidemic, *D. barnimiana*, streptozotocin, α -amylase

Introduction

Diabetes mellitus (DM) is a chronic metabolic disease attributed to hyperglycemia, hyperlipidemia, and hyperaminoacidemia, along with insulin insufficiency, insulin-receptor resistance, or both.¹ It affects nearly 10% of the global population. Despite the introduction of modern hypoglycemic agents, secondary complications remain a major medical issue. Also, most modern drugs have drawbacks, such as gastrointestinal disorders and weight gain.² Multiple pathogenic processes are engaged in the inauguration of DM, such as autoimmune destruction of β cells of the pancreas and others precipitating insulin resistance or both.³ Among the different types of DM, type 2 is more prevalent worldwide. DM is related to microvascular and macrovascular complications.⁴ Hyperglycemia is an important risk factor in the establishment and prognosis of complications.⁵ It is the third-leading cause of death, next to heart attacks and cancer.^{6,7}

Since ancient time, medicinal plants have been part and parcel of human attempts to fight diseases, including DM.⁸ According to a WHO estimation, around 80% of the population in the developing world relies on traditional phytomedicines for primary health-care needs.⁹ Near to 25% of contemporary medicines originate from traditional medicinal plants. For

example, metformin was detected in *Galega officinalis*.¹⁰ Based on ethnobotanical studies, there are >1,200 medicinal plants involved in DM treatment, due to their efficacy, few side effects, ease of use, and accessibility.¹¹

The WHO has suggested the appraisal of cultural medicinal plant treatments considered excellent potential for addressing for DM.¹² The leaves of *Croton macrostachys*,¹³ *Hypoestes forskalii*,¹⁴ *Thymus schimperi*,¹⁵ *Rubus erlangeri*,¹⁶ *Bersama abyssinica*,¹⁷ *Calpurnia aurea*,¹⁸ *Ajuga remota*,⁹ *Otostegia integrifolia*,¹⁹ roots of *Morus alba*,¹¹ *Polystachys delile*,²⁰ and *Ficus religiosa*, *Datura stramonium*²¹ and *Schinus molle* seed extracts,²² *Hagenia abyssinica* flowers,²³ and *Ficus bengalensis*,¹¹ and *Terminalia brownii* stem bark²⁴ are some of the medicinal plants used in traditional Ethiopian medicine for the treatment of DM. Single-plant preparations and formulated mixtures have been utilized for DM management. For example, dendrobium mixture and *fufang zhenzhu tiaozhi* formula are preparations used in traditional Chinese medicine for long-term clinical treatment of DM and its complications.^{25,26}

Dorstenia barnimiana is one herb with an indispensable role in the treatment of DM. It is a seasonal African plant used in Ethiopian folk medicine for multiple diseases. In many regions of Ethiopia, the root of the plant is used against common colds, and the leaves are used to remove retained placenta.²⁷ Its root powder is used for treatment of hepatitis, rabies, syphilis, unhealthy weight loss, dysentery, fever, hypertension, diarrhea, and DM.²⁸ The whole plant has been used for the treatment of schizophrenia via inhalation.²⁹ Crushed *D. barnimiana* leaves have been applied topically for wound healing.³⁰ Its root powder has shown antiplasmodial activity in a mouse model.³¹ Other medicinal plants in the same family and genus have confirmed anti-DM effects, eg, *D. psilurus* and *D. picta*.³² *Dorstenia* is a potential source of secondary metabolites, such as steroids, benzofuran derivatives, prenylated and geranylated coumarins, flavonoids, styrenes, and triterpenoids.^{33,34} Despite the fact that *D. barnimiana* juice from dry roots mashed and mixed with water has been used for the treatment of DM in traditional Ethiopian medicine, there has been neither in vivo nor in vitro scientific study on its antidiabetic activity to our knowledge. To investigate its use, the current study evaluated *D. barnimiana*'s antidiabetic and antidyslipidemic effects in vitro in animals.

Methods

Drugs and Chemicals

Drugs, chemicals, and instruments used in the experiment were absolute methanol (ReAgentchem), 0.9% normal saline (Cadila Pharmaceuticals), distilled water (EPharm), chloroform, ethyl acetate (Nice Laboratory Reagent), examination gloves, 0.5% trisodium citrate (Loba Chemie), 95% sulfuric acid (Thermo Fisher Scientific), Dragendorff's reagent (Super-Tek), glacial acetic acid (Loba Chemie), benzene (Nice Laboratory Reagent), ammonia solution (HiMedia), FeCl₃ (Super-Tek), 10% ammonium hydroxide (Rankem), porcine α -amylase (Molychem), 0.01 N sodium hydroxide (Blulux Laboratories), acetic anhydride and Mayer's reagent (May and Baker), 40% glucose solution (Reyoung Pharmaceuticals), glibenclamide (Glc; Cadila Pharmaceuticals), streptozotocin (Stz; Thermo Fisher Scientific), acarbose (Bayer), and analytical graded syringes were purchased from Addis Ababa pharmaceutical and chemical suppliers.

Apparatus and Supplies

Instruments, apparatus, and supplies used in the experiment a mini orbital shaker (Stuart Scientific), digital weighing balance, lyophilizer (Wagtech Jouan Nordic DK-3450), deep freezer, Whatman filter paper (number 1, Schleicher and Schuell Microscience), permanent marker, test tubes, volumetric flasks, Erlenmeyer flasks, measuring cylinder, mouse cages, gavages (oral feeding syringes), syringes (1 mL and 3 mL), desiccators, refrigerator, micro-hematocrit centrifuge, digital thermometer, surgical blades, one-touch basic glucometer and test strip (Hoffman-La Roche), UV-visible spectrophotometer (Agilent Technologies), grinder (KZ-III), rota vapor (Rotary Evaporator RE300), and BS-240 clinical chemistry analyzer (Mindray).

Experimental Animals

Male Swiss albino mice (body weight 27–35 g, age 8–12 weeks) were used in the present study, except for the acute oral toxicity study, where healthy nonpregnant and nulliparous female mice (body weight 27–35 g, age, 8–12 weeks) were used. The mice were purchased from the Ethiopian Health and Nutrition Research Institute and kept at the animal house of the Pharmacy Department, Debre Tabor University. Animals were housed in ventilated cages at 25 $\pm$ 2°C and a 12-

hour light–dark cycle, with free access to water and standard mouse pellets ad libitum. After indiscriminate grouping and before the commencement of the experiment, animals were acclimatized for 2 weeks. This period allowed them to adapt a strange environment and also promote reproducible experimental outcomes. All experimental protocols were followed as per the guiding principles for research involving animals and as approved by the Research and Ethics Review Committee of the College of Health Sciences, Debre Tabor University, Ethiopia.

Collection and Authentication of Plant Materials

Roots of *D. barnimiana* were collected at Fogera, Debub Gondar, Amhara in May 2021. Botanical identification and authentication were done by a taxonomist at the Department of Biology, College of Natural and Computational Sciences, University of Gondar, and a voucher specimen (Wor2/2021) was deposited there for future reference.

Preparation and Extraction of the Plant

After collection, roots were initially washed using running tap water to remove extraneous materials, air-dried under shade in the Debre Tabor University pharmacology laboratory, then chopped into small pieces and ground into coarse powder. The crude 80% methanolic crude extract of the root of *D. barnimiana* was prepared by cold maceration. After weighing 1,000 g coarsely powdered root using a sensitive digital weighing balance, it was soaked into three clean Erlenmeyer conical flasks, each containing 2 L 80% methanol. Flasks were then placed on a shaker at 120 rpm for 72 hours at room temperature for intermittent agitation. The entire mixture was first filtered through a funnel plunged with muslin cloth twice, then the filtrate was passed through Whatman filter paper (number 1). After filtration, the remaining residue or marc was remacerated twice for 6 days with a fresh solvent of 80% methanol to maximize the yield. The resulting solution after successive filtration was concentrated using a rotary evaporator set at 40°C to remove methanol. Finally, the concentrated aqueous filtrate was placed in the deep freezer overnight, then subjected to lyophilization at –50°C and vacuum pressure to remove water. The percentage yield of the powdered extract was calculated, and it was kept in an airtight container at –4°C until used.³⁵

Preliminary Phytochemical Analysis

Qualitative preliminary phytochemical profiling tests were carried out to determine the presence or absence of major classes of phytochemicals for an 80% methanol crude extract of the root of *D. barnimiana* using standard tests.^{36,37}

Acute Oral Toxicity Study

The median lethal dose (LD₅₀) of *D. barnimiana* extract was estimated orally in the female mice following the 2008 Organization for Economic Cooperation and Development guideline 425. Accordingly, five female mice aged 6–8 weeks were engaged. All mice were fasted (food but not water) for 4 hours before and 2 hours after administration of the extract. First, a sighting study was performed to determine the starting dose. For this, 2,000 mg/kg was given per mouse by oral gavage. The first animal received a limited dose of 2,000 mg/kg and monitored for death for the last 24 hours. Then, four other mice were sequentially treated. The animals were housed separately and observed externally for 4 hours with 30-minute intervals and then for 14 consecutive days, with an interval of 24 hours for general signs and symptoms of toxicity, like food and water intake, death, changes in skin and fur, eyes, and mucous membranes, and respiratory and behavior patterns.³⁸

Determination of α -Amylase Inhibition Activity

Commenced using the chromogenic 3,5-dinitrosalicylic acid (DNSA) method in congruent with previous studies.¹⁷ The hydromethanolic crude extract of *D. barnimiana* root and standard agent (acarbose) were first dissolved in PBS. Sample concentrations of 50–500 μ g/mL were prepared accordingly. α -Amylase solution (200 μ L) was mixed with an equal volume of each concentration of the extract and acarbose and incubated for 10 minutes. Afterward, an equal volume of the starch solution (1% in water, w/v) was added to each test tube. The reaction was ended after 3 minutes upon adding 200 μ L DNSA, then boiled for 10 minutes in a water bath at 85°C. Thereafter, it was allowed to cool to room temperature and 5 mL of DW was added. UV-visible spectrophotometry at 540 nm wavelength was used to measure absorbance capacity. The blank with full enzyme kinetics was created by mixing 200 μ L enzyme with an equal volume of buffer. A control blank solution, and sample blank (extract or acarbose) without enzyme were created in a similar fashion. The α -

amylase inhibitory activity is articulated as percentage inhibition and was calculated using the equation given below.²³ Finally, α -amylase inhibition was plotted against the extract and standard concentration:

$$\text{Inhibition (\%)} = \frac{(Ac - Acb) - (As - Asb)}{Ac - Acb} \times 100 \quad (1)$$

where Ac refers to absorbance of the control (enzyme and buffer), Acb absorbance of the control blank (buffer without enzyme), As absorbance of the sample (enzyme and inhibitor), and Asb absorbance of the sample blank (inhibitor without enzyme).

Grouping and Dosing

Simple random sampling was utilized for the organization of experimental groups and allocation of treatment types. Male mice were involved in all models in this study, due to insensitivity of female mice to Stz and insulin.^{39,40} In normoglycemic and oral glucose-loaded models, overnight-fasted normal mice were arbitrarily placed into five groups. Group I (negative control) received distilled water (DW10, 10 mL/kg, PO), groups II–IV 100 mg/kg (DB100), 200 mg/kg (DB200), and 400 mg/kg (DB400) hydromethanolic root extract of *D. barnimiana* PO, respectively, and group V (positive control) 5 mg/kg glibenclamide (Glc5).

In the Stz-induced diabetic model, overnight fasted-diabetic mice were randomly divided into six groups and treated:

Group I: Diabetic controls DW10

Group II: Diabetic mice DB100

Group III: Diabetic mice DB200

Group IV: Diabetic mice DB400

Group V: Diabetic mice Glc5

Group VI: Normal controls (sham group) DW10

Determination of Parametric Measures

A blood sample was withdrawn from the tail vein of each fasted mouse aseptically in all animal models. Blood-glucose (BG) levels were measured using the electronic glucometer and test strip. Average values were calculated after triplicate measurements of fasting BG (FBG), body weight, and sample weight.

Induction of Experimental DM

Stz solubilized in 0.1 M citrate buffer (pH 4.5) was applied. Stz (150 mg/kg) solution was administered IP to overnight-fasted mice. Thirty minutes after Stz administration, normal pellet feeding was allowed. Six hours later, in order to avert deaths associated with hypoglycemic, episodes animals were given access to 5% glucose solution for a day. Three days later, they were screened for DM, and mice with FBG >200 mg/dL were used for further study.^{41,42}

Determination of Hypoglycemic Effects in Normoglycemic Mice

Normal mice were fasted overnight, but had water ad libitum. Then, they were partitioned into the five. After treatment, the BG level (BGL) of each mouse was evaluated before treatment at baseline and then at 1, 2, 4, and 6 hours posttreatment.²⁴

Determination of Antihyperglycemic Effects in Oral Glucose-Loaded Mice

Thirty fasted mice were assigned indiscriminately into five groups and then treatment was administered accordingly. Glucose solution (40% w/v, 2 g/kg) was given to each mouse orally after 30 minutes' therapy. The BGL of each mouse was evaluated prior to oral glucose loading (baseline) and then at 30, 60, 90, and 120 minutes after glucose introduction.²⁴

Determination of Effects of the Extract on Blood Glucose, Lipid Profile, and Body Weight of Diabetic Mice

Stz-induced diabetic and normal male mice were partitioned arbitrarily into six groups. Treatment was given to mice once daily for 4 consecutive weeks. At 0, 7, 14, 21, and 28 days, FBG and body weight were evaluated. The weight of each mouse had also been measured before induction of DM. Before the end of the experiment, mice were euthanized humanely with pentobarbitone 150 mg/kg IP,⁴³ then blood samples collected via decapitation. The samples were used to

measure the level of total cholesterol (TC), triglycerides (TGs), and low-density lipoprotein (LDL), very low-density lipoprotein (VLDL), and high-density lipoprotein (HDL) cholesterol (HDL-C).³⁵

Statistical Analysis

IC₅₀ values were calculated from plots of inhibitor concentration versus percentage inhibition of standard curves. Data are given as means ± SEM and were analyzed using SPSS 24. Between- and within-group analyses were done using one-way ANOVA followed by post hoc Tukey multiple-comparison tests. *P* < 0.05 was considered statistically significant.

Results

Yield Determination

From 1,000 g root powder of *D. barnimiana*, 406.76 g hydromethanolic crude extract was obtained (see Table 1).

Phytochemistry

Preliminary qualitative tests were done as per standard test guidelines. These tests revealed the existence of terpenoids, saponins, tannins, flavonoids, glycosides, alkaloids, and phenolic compounds, whereas resins, anthraquinones, and steroids were not detected (See Table 2).

In Vitro α -Amylase Inhibitory Activity

The α -amylase inhibitory activity of the acarbose and crude extract are shown in Figure 1. Concentration-dependent inhibition was seen for various concentrations of the tested extract and the standard. Hydromethanolic crude root extract of *D. barnimiana* showed a maximum α -amylase enzyme inhibition activity of 67.52% at 500 μ g/mL, with IC₅₀ of 4.95 μ g/mL. The maximum α -amylase inhibition was observed at a high concentration of acarbose (94.65%), with IC₅ of 2.14 μ g/mL.

Table 1 Quantity and quality of hydromethanolic crude root extract of *D. barnimiana*

Solvent	Color and texture of extract	Outcome (g)	Outcome (w/w)
80% hydromethanol	Dark-brown powder	406.76	40.68%

Abbreviations: gm, gram; w/w, weight by weight.

Table 2 Qualitative phytochemical profiling of hydromethanolic crude root extract of *D. barnimiana*

Sample	Phytochemical	Test performed	Result
1	Alkaloids	Wagner's	+
2	Anthraquinones	Borntrager's	–
3	Flavonoids	Alkaline reagent (NaOH)	+
4	Glycosides	Keller–Killiani	+
5	Phenolic compounds	Ferric chloride	+
6	Plant steroids	Liebermann–Burchard	–
7	Resins	Precipitation	–
8	Saponins	Froth	+
9	Tannins	Braemer's	+
10	Terpenoids	Salkowski's	+

Notes: (+): exists, (–): does not exist.

Figure 1 α -Amylase inhibitory activity of crude root extract of *Dorstenia barnimiana* and acarbose.

Note: Values presented as means $\pm$ SEM (n=3).

Abbreviation: SEM, standard error of the mean.

Acute Toxicity

Mortality was not observed at 2,000 mg/kg during observation, indicating that the LD₅₀ for the extract was >2,000 mg/kg. The mice did not show any physical or behavioral abnormalities.

Effect on Basal Plasma Glucose

An investigation in normoglycemic mice made it possible to investigate hypoglycemic effects of the crude extract. Administration of the vehicle (DW 10 mL/kg) did not change plasma-glucose levels significantly during testing. Likewise, administration of hydromethanolic extracts at different test doses did not significantly ameliorate basal plasma-glucose levels compared to the normal control group, whereas the standard agent Glc5 markedly lowered plasma-glucose levels from 129.16 $\pm$ 0.39 mg/kg to 64.72 $\pm$ 1.95 mg/kg (see Table 3).

Table 3 Effect of hydromethanolic crude root extract of *D. barnimiana* on blood-glucose levels in normoglycemic mice

Group	Fasting blood glucose (mg/dL)				
	0	1	2	4	6
DW10	129.66 $\pm$ 0.84	128.49 $\pm$ 0.64	127.43 $\pm$ 0.87	125.11 $\pm$ 1.23	123.42 $\pm$ 1.55
DB100	127.70 $\pm$ 0.50	126.87 $\pm$ 0.41 ^{β*}	125.71 $\pm$ 0.44 ^{β*}	124.57 $\pm$ 0.53 ^{β*}	123.30 $\pm$ 0.58 ^{β*}
DB200	130.12 $\pm$ 0.84	128.82 $\pm$ 0.99 ^{β*}	127.82 $\pm$ 1.04 ^{β*}	127.14 $\pm$ 1.10 ^{β*}	125.99 $\pm$ 1.17 ^{β*}
DB400	128.37 $\pm$ 0.36	128.08 $\pm$ 0.39 ^{β*}	127.26 $\pm$ 0.44 ^{β*}	126.61 $\pm$ 0.51 ^{β*}	125.41 $\pm$ 0.35 ^{β*}
Glc5	129.16 $\pm$ 0.39	91.49 $\pm$ 3.21 ^{α*}	81.07 $\pm$ 3.01 ^{α*}	68.82 $\pm$ 3.59 ^{α*}	64.72 $\pm$ 1.95 ^{α*}

Notes: Values expressed as means $\pm$ SEM (n=6) and analyzed by one-way ANOVA followed by post hoc Tukey tests. ^{α} Compared to negative control; ^{β} compared to Glc5. * P <0.001.

Abbreviations: DB, *Dorstenia barnimiana*; DW, distilled water; Glc, glibenclamide.

Effect of Extract on Oral Glucose Tolerance in Mice

The chief source of energy in animals body is glucose, and the oral glucose-tolerance test (OGTT) enables us to evaluate the body's capacity to use glucose wisely.⁴⁴ A single dose of crude extract was administered to normoglycemic mice to verify its antihyperglycemic activity. OGTTs with 40% glucose solution resulted in hyperglycemia with a peak of 179.51 ± 2.29 mg/dL 30 minutes after administration in the negative-control group, but that peak was significantly averted by both the standard drug (Glc5) and the crude extract at all test doses ($P < 0.001$, see Table 4).

Antihyperglycemic Activity in Stz-Induced Diabetic Mice

BGLs were measured every week in normal and diabetic mice. After induction, diabetic mice showed significant differences in baseline BGLs compared to normal mice ($P < 0.001$), but no significant differences were found among the groups of diabetic mice. Even though diabetic mice were treated for about 2 weeks with DB100 daily and showed BGL reduction to 275.65 ± 2.62 mg/dL in the first and 268.22 ± 3.10 mg/dL in the second week, this reduction fell short of statistical significance compared to the negative controls, though it had shrunk remarkably by day 21 ($P < 0.01$) and day 28 ($P < 0.001$) compared to diabetic controls (see Table 5). Correspondingly, medium and high doses and Glc5 had reduced FBG at 1 week after commencing therapy.

Effect on Body Weight of Diabetic Mice

Before induction of DM and baseline (3 days after Stz injection), there was no statistically significant weight-loss difference between diabetic and normal mice (see Figure 2). Stz induction had caused a significant loss of weight in

Table 4 Effect of hydromethanolic crude root extract of *D. barnimiana* on blood-glucose levels in mice challenged with oral glucose

Group	Fasting glucose level (mg/dL)				
	0	30'	60'	90'	120'
DW10	130.56±0.61	179.51±2.29	168.98±2.67	158.20±2.76	140.20±2.95
DB100	130.59±1.46	158.04±3.01 ^{a*}	148.88±2.26 ^{a*}	147.35±2.11 ^{a*}	139.63±1.14 ^{a*}
DB200	133.61±1.65	152.92±2.51 ^{a*}	142.34±1.26 ^{a*}	140.03±0.83 ^{a*}	128.27±1.03 ^{a*}
DB400	130.80±1.87	143.33±2.24 ^{a*}	118.81±2.35 ^{a*}	97.82±1.33 ^{a*}	87.47±0.74 ^{a*}
Glc5	131.62±1.85	135.55±2.08 ^{a*}	102.28±4.22 ^{a*}	94.65±4.34 ^{a*}	84.49±4.70 ^{a*}

Notes: Values expressed as means ± SEM (n=6) and analyzed by one-way ANOVA followed by post hoc Tukey tests. ^aCompared to negative control; ^ccompared to DB200 mg/kg; ^bcompared to DB400 mg/kg; ^dcompared to Glc5. * $P < 0.05$; ^a $P < 0.01$; ^{a*} $P < 0.001$.

Abbreviations: DB, *Dorstenia barnimiana*; DW, distilled water; Glc, glibenclamide.

Table 5 Antihyperglycemic effect of hydromethanolic crude root extract of *D. barnimiana* in streptozocin-induced diabetic mice

Group	Blood glucose (mg/dL) at weekly intervals				
	Day 0	Day 7	Day 14	Day 21	Day 28
DW10	277.17±2.00 ^{c*}	277.83±2.85 ^{c*}	275.18±2.34 ^{c*}	271.97±2.55 ^{c*}	268.57±1.64 ^{c*}
DB100	282.83±3.21 ^{c*}	275.65±2.62 ^{c*}	268.22±3.10 ^{c*}	259.27±1.84 ^{a*}	255.41±0.78 ^{a*}
DB200	279.02±6.39 ^{c*}	256.16±3.38 ^{a*}	252.56±3.06 ^{a*}	251.01±2.86 ^{a*}	248.69±2.86 ^{a*}
DB400	276.50±2.51 ^{c*}	220.59±1.89 ^{a*}	216.38±2.13 ^{a*}	214.43±1.98 ^{a*}	214.07±2.03 ^{a*}
Glc5	276.45±3.34 ^{c*}	185.56±2.65 ^{a*}	178.82±2.47 ^{a*}	165.94±1.75 ^{a*}	163.75±1.69 ^{a*}
NC	129.93±0.75 ^{a*}	128.67±0.38 ^{a*}	127.57±0.24 ^{a*}	126.08±0.54 ^{a*}	125.56±0.62 ^{a*}

Notes: Values expressed as means ± SEM (n=6) and analyzed by one-way ANOVA followed by post hoc Tukey tests. ^aCompared to negative control; ^ccompared to DB200 mg/kg; ^bcompared to DB400 mg/kg; ^dcompared to Glc5; ^ecompared to normal control. ^a $P < 0.01$; ^{a*} $P < 0.001$.

Abbreviations: DB, *Dorstenia barnimiana*; DW, distilled water; Glc, glibenclamide; NC, normal control (pretreatment significantly brought down the volume of BGL).

Figure 2 Effect of *Dorstenia barnimiana* crude extract on body weight of diabetic mice in Stz-induced model.

Notes: Values expressed as means $\pm$ SEM (n=6 mice in each group) and analyzed by one-way ANOVA followed by post hoc Tukey tests. ^aCompared to negative control; ^εcompared to NC. *P<0.001.

Abbreviations: DB, *Dorstenia barnimiana*; DC, diabetic control; DM, diabetes mellitus; DW, distilled water; Glc, glibenclamide; NC, normal control.

diabetic controls by days 7 ($P<0.05$), 14 ($P<0.001$), 21 ($P<0.001$), and 28 ($P<0.001$) compared to normal controls. In the first week of therapy, neither the crude extract nor the standard agent was able to retard the weight loss. Significant body-weight increments were recorded for DB200 ($P<0.05$), DB400 ($P<0.01$), and Glc5 ($P<0.01$) 2 weeks after treatment had commenced compared to the diabetic control group. Similarly significant body-weight recovery was recorded for DB100 ($P<0.05$), DB200 ($P<0.001$), DB400 ($P<0.001$), and Glc5 ($P<0.001$) on day 21 compared to diabetic controls. All doses of the crude extract and Glc5 showed an significant ($P<0.001$) increment in body weight after 28 days of therapy (see Table 6).

Effect on Serum-Lipid Profile of Diabetic Mice

The DB100 group showed significant reductions in serum levels of TC ($P<0.001$), TGs ($P<0.01$), LDL-C ($P<0.01$), and VLDL-C ($P<0.05$), but failed to reach statistical significance for HDL-C compared to diabetic controls (see Table 7). Likewise, DB200-treated groups showed significant reductions in serum levels of TC ($P<0.001$), TGs ($P<0.001$), LDL-C ($P<0.001$), and VLDL-C ($P<0.01$), but did not show a significant increase in serum HDL-C compared to diabetic

Table 6 Effect of crude extract on body weight of diabetic mice

Group	Body weight (g)		
	Day 7	Day 14	Day 21
DW10	27.66 $\pm$ 0.69 ^{ε\$}	25.53 $\pm$ 0.64 ^{ε*}	25.33 $\pm$ 0.56 ^{ε*}
DB100	27.79 $\pm$ 0.78	27.91 $\pm$ 0.56	28.12 $\pm$ 0.53 ^{α\$β\$δ\$}
DB200	28.22 $\pm$ 0.72	28.77 $\pm$ 0.54 ^{α\$}	29.75 $\pm$ 0.55 ^{α*}
DB400	28.46 $\pm$ 0.41	29.93 $\pm$ 0.40 ^{α^}	30.74 $\pm$ 0.41 ^{α*}
Glc5	27.84 $\pm$ 0.39	30.05 $\pm$ 0.79 ^{α^}	30.89 $\pm$ 0.53 ^{α*}
NC	31.05 $\pm$ 1.27 ^{α\$}	31.53 $\pm$ 1.26 ^{α*}	32.56 $\pm$ 0.90 ^{α*}

Notes: Values expressed as means $\pm$ SEM (n=6) and analyzed by one-way ANOVA followed by post hoc Tukey tests. ^aCompared to negative control; ^δcompared to DB400 mg/kg; ^βcompared to Glc5; ^εcompared to NC. ^{\$}P<0.05; [^]P<0.01; ^{*}P<0.001.

Abbreviations: DB, *Dorstenia barnimiana*; DW, distilled water; Glc, glibenclamide; NC, normal control.

Table 7 Effect of crude extract on serum-lipid levels in diabetic mice

Group	Serum Lipid Level (mg/dL)				
	TC	TGs	LDL-C	VLDL-C	HDL-C
DW10	174.07±2.98 ^{a*}	119.02±3.03 ^{a*}	115.21±3.34 ^{a*}	23.80±1.49 ^{a*}	35.06±0.86 ^{a*}
DB100	148.16±3.92 ^{a*β*∅^δ*}	101.62±3.10 ^{a*β*∅^δ*}	90.57±0.38 ^{a*β*∅*}	20.32±0.41 ^{a*β*∅*}	37.27±0.54
DB200	130.17±1.18 ^{a*}	85.31±3.18 ^{a*}	69.80±0.56 ^{a*β*}	17.06±0.57 ^{a*β*}	43.31±1.02 ^{β^δ*}
DB400	127.04±0.69 ^{a*}	83.59±1.54 ^{a*}	65.74±1.54 ^{a*β*}	16.72 ±0.62 ^{a*}	44.58±0.57 ^{a*}
Glc5	122.12±4.20 ^{a*}	81.09±2.47 ^{a*}	60.92±0.67 ^{a*}	16.22±1.01 ^{a*}	44.98±2.29 ^{a*}
NC	107.43±3.25 ^{a*}	96.75±2.33 ^{a*}	40.73±1.05 ^{a*}	19.35±0.42 ^{a*}	47.35±0.23 ^{a*}

Notes: Values expressed as means ± SEM (n=6) and analyzed by one-way ANOVA followed by post hoc Tukey tests. ^aCompared to negative control; [∅]compared to DB200 mg/kg; ^δcompared to DB400 mg/kg; ^βcompared to Glc5; ^ccompared to NC. [§]P<0.05; [^]P<0.01; ^{*}P<0.001.

Abbreviations: DB, *Dorstenia barnimiana*; DW, distilled water; Glc, glibenclamide; NC, normal control.

controls. However, the group of mice treated with the highest dose and standard agent for 28 days showed a significant decrease in TC, TGs, LDL-C, and VLDL-C and increased HDL-C markedly compared to diabetic controls.

Discussion

DM is a global public health challenge, especially with current medications being incapable of averting diabetic complications. Therefore, contemporary research aimed at identifying phytomedicines is needed for new preventive and therapeutic approaches. In Addis Ababa, Ethiopia, *D. barnimiana* tubes are sold for the treatment of a variety of diseases, including DM.³⁴ In this study, before advancing to the animal study, the antidiabetic effect of *D. barnimiana* root was investigated in vitro by ascertaining its α -amylase inhibition activity. The pancreas is the principal organ engaged in detecting the body's energy and dietary states through the concentration of BG. When the BGL shifts, the pancreas responds via monitoring the release of insulin, α -amylase, and other pancreatic enzymes. Pancreatic α -glucosidase and α -amylase inhibition is an efficient approach to decrease levels of postprandial hyperglycemia through starch-breakdown control. This is a key approach in the management of type 2 diabetic patients and prediabetics.⁴⁵ α -Amylase inhibition assays were done to verify the α -amylase inhibition effect of *D. barnimiana* by employing acarbose as a standard reference. Acarbose is a modern medicine for managing DM by inhibiting the α -amylase and α -glucosidase activity that converts starch into glucose,⁴⁶ whereas the plant extract acts as a natural inhibitor. The results showed that the plant may have antidiabetic properties, since there was no appreciable discrepancy between IC₅₀ values of the standard drug (acarbose) and natural inhibitor (extract) at 2.14 μ g/mL vs 4.95 μ g/mL, respectively. Flavonoids from *Dorstenia* spp. (eg, *D. psilurus*), such as artelastin, have been found to be α -glucosidase inhibitors, thereby reducing the risk of hyperglycemia by expanding the carbohydrate-absorption period.⁴⁷ The roots of *D. drakena* have antisecretory effects,⁴⁸ and as such, *D. barnimiana* may also be beneficial in decreasing the secretion of pancreatic enzymes.

The in vitro finding establishes the extract's candidacy for further analysis of its antidiabetic activity in in vivo models. The effect of the extract on BGL was evaluated by initiating normoglycemic, oral glucose tolerance, and Stz-induced diabetic mice models. In the acute oral toxicity study at the limit dose of 2,000 mg/kg, *D. barnimiana* root extract remained atoxic during the study period. This safety profile is congruent with a report by Derebe et al.³¹ Normoglycemic models provide clues to the mechanisms of action of natural remedies.²¹ The results from the normoglycemic studies implied that the hydromethanolic crude extract of *D. barnimiana* had no hypoglycemic activity compared to normal controls, suggesting a lower risk of hypoglycemia for individuals in a euglycemic state. These findings are comparable to Alene et al.⁴⁹

A drug that is efficient in DM will have the capacity to control dysglycemia through different hepatic and extrahepatic mechanisms. The capacity of the extract to prevent hyperglycemia can be determined with the OGTT.⁵⁰ Interestingly, in this model, even the lowest dose of the crude extract resulted in a significant reduction in BGL after 30 minutes. This

shows that the extract has the ability to enhance glucose regulatory mechanisms, indicating its potential for minimizing meal-related hyperglycemia and complications of DM. It was observed that the extract exerted its action in a dose-dependent manner, and the highest dose (DB400) produced the maximum activity in all tests undertaken. The lowest dose (DB100) appeared to be ineffective in reducing the resurgence of BGL in glucose-loaded animals significantly at 2 hours. This may be due to the lower concentration of the active principles in the serum attained initially associated with the low dose administration or may be due to rapid elimination of the drug from the body.

Stz causes DM in experimental animals due to its higher inductive rate and selective toxicity.⁴⁹ After IP or IV administration, it enters pancreatic β cells through the GLUT2 transporter. Accumulation of Stz in β cells results in cell death via methylating DNA, which depletes the production of insulin.⁵¹ Insulin deficiency disrupts the metabolism of carbohydrates, fats, and proteins in liver, adipose, and muscle tissue thereby raising levels of glucose, cholesterol, nitrogen balance, muscle wasting, and weight loss.²¹ Chemically induced DM is able to mimic the physiological and biochemical derangements in the actual disease.⁵² Oral administration of the extract resulted in appreciable improvement in glucose homeostasis in diabetic mice. As shown in Table 6, the crude extract possessed antihyperglycemic activity when administered to Stz-induced diabetic mice at all test doses. In light of the results obtained from the Stz-induced model, it could be deduced that the roots of *D. barnimiana* have authentic antidiabetic activity and that their use in traditional medicine to control DM may be justified.

DM is also distinguished by dyslipidemia, which accounts for elevated levels of TC, TGs, and LDL-C and lower levels of HDL-C and is a major cause of cardiovascular problems.⁵³ Insulin triggers lipoprotein lipase, which can metabolize TGs and VLDL-C. In diabetic patients, lipoprotein lipase is less active due to insulin deficiency, and this can lead to development of diabetic dyslipidemia.³⁵ The serum-lipid profile of the diabetic control group revealed a significant increase in serum TGs, TC, VLDL-C, LDL-C and decrease of HDL-C compared to normal controls ($P<0.001$). Interestingly, treatment with the extract resulted in significant improvements in lipid profile: at 4 weeks, serum TC, TGs, VLDL-C, and LDL-C had reduced and were able to enhance HDL-C levels. These findings also correlate with those of Tsofack et al⁴⁷ and Jyothsna et al,⁴⁴ who (respectively) reported that *D. picta* and *Cinnamomum zeylanicum* reduced cholesterol and TGs in experimentally induced diabetic rats. This activity of the plant extract may be related to the presence of secondary metabolites implicated in the antihyperlipidemic activity of plant extracts.

Several plant constituents (such as flavonoids) detected in *D. barnimiana* are known to decrease TGs, TC, LDL-C, and VLDL-C.^{47,54} Absorption of free fatty acids and free cholesterol can be inhibited by intervening the enzymatic activity of pancreatic lipase and pancreatic cholesterol esterases, thereby preventing hyperlipidemia related to DM. The antihyperlipidemic effects of the extract could be attributed to inhibition of those enzymes. In addition to their hypolipidemic properties, flavonoids have been shown to decrease serum glycated hemoglobin levels.⁵⁵ Herein, even though all doses of the crude extract were able to modify serum-cholesterol levels, the highest dosage (DB400) exhibited the best antihyperlipidemic activity. The antihyperlipidemic activity of medicinal plants points to new directions for the development of antidiabetic agents with multiple therapeutic targets.⁵⁶

Weight loss is another cardinal feature of DM. It is due to insulin insufficiency, which results in depletion in protein synthesis and in turn reduced amino-acid uptake by tissue, resulting in lipolysis and proteolysis in adipose tissue.¹⁵ In this study, there was a genuine weight reduction in Stz-induced diabetic mice. However, after treatment with extract, the weight lost was redeemed. This prevention of weight loss is in line with other studies done on the same plant. At all tested doses, the significantly prevented weight loss in a dose-dependent manner compared to negative controls on the 4-day suppression and curative tests, but not the prophylaxis test.³¹ The capacity of *D. barnimiana* extracts to prevent significant weight loss may be attributed to the bioactive compounds that suppresses oxidative stress from free radicals generated via hyperglycemia.

In the present findings, the root extract showed an antihyperglycemic effect in Stz-induced diabetic mice without causing hypoglycemia in normoglycemic mice while Glc showed both hypoglycemic and antihyperglycemic effects in normoglycemic and Stz-induced diabetic mice. As a result, the mechanism of action of root extracts of *D. barnimiana* in the treatment of diabetic mice is unquestionably different from the standard drug Glc, which acts by increasing insulin secretion from the pancreas and should be used with caution in patients with cardiovascular disease or in elderly patients, in whom hypoglycemia would be especially dangerous.⁵⁷ However, wide-ranging pharmacological and biochemical

research is required to address the detailed mechanisms and active principles responsible for the antihyperglycemic effects observed.

Preliminary qualitative analysis was part of our work which revealed the presence of secondary metabolites like terpenoids, alkaloids, flavonoids, phenols, saponins, tannins, and cardiac glycosides, whereas resins, anthraquinones, and steroids were not detected, similar to other *Dorstenia* spp., eg, *D. mannii*⁵⁸ and *D. picta*.⁴⁷ However, terpenoids were not detected in *D. barnimiana* in a qualitative study conducted by Derebe et al. From the present plant-extraction yield determination, there was no marked difference from a previously reported yield (40.68% vs 44.3% w/w).³¹ This discrepancy in outcomes might be explained by ecological, edaphic, or climate disparities.

Secondary metabolites with antidiabetic activity include alkaloids, glycosides, galactomannan gum, polysaccharides, peptidoglycans, hypoglycans, guanidine, phytosterols, flavonoids, phenols, saponins, tannins, carbohydrates, polyphenol, glycopeptides, terpenoids, amino acids, and inorganic ions.^{59–61} Tannins are excellent free-radical scavengers.⁶² Tannins also inhibit the activity of digestive enzymes, such as trypsin and amylase.⁶³ Saponins also exhibit hypoglycemic effects via several means such as rehabilitation of insulin production and release, insulin-receptor sensitivity, reduction of hepatic gluconeogenesis, activation of glycogen production, ameliorative effects on gluconeogenesis, inhibition of α -glucosidase activity, and inhibition of mRNA expression of glycogen phosphorylase and glucose-6 phosphatases.⁶⁴ Likewise, flavonoids from *D. psilurus* can activate, thereby stimulating glucose uptake, inhibiting glucose production, and lowering hyperglycemia.⁴⁷ Prenylated flavonoids isolated from *D. mannii* such as 6,8-diprenyleriodictyol, dorsmanin C, and dorsmanin F, are potent scavengers of the stable free radical DPPH.⁶⁵ *D. barteri*, *D. convexa*, *D. foetida*, *D. psilurus*, and *D. ciliate* also have proven antioxidant activity, generally related to the presence of flavonoids and other phenolic compounds.⁶⁶ Therefore, the activity of this plant extract against DM might be due to the content of these phenolic compounds.

Conclusion

The present glycemic study found that the hydromethanolic crude root extract of *D. barnimiana* has potential antidiabetic and antihyperlipidemic effect, which confirms the plausibility of traditional claims. However, before recommending it for use, further studies should be conducted with relevant models to elucidate its mechanisms of action and confirm its efficacy and safety profile.

Recommendations

This study clearly backs the traditional claims of this extract's therapeutic properties. As such, further in-depth investigation is needed using other relevant models through histopathological examinations of the pancreas for more comprehensive evidence. Isolation and characterization of the active components to isolate and identify the active principle(s) responsible for its ethnopharmacological activity and to elucidate the exact mechanism action of this plant using advanced technologies like HPLC and GMSC is needed.

Limitations

- This was a preliminary study that aimed to validate only the antidiabetic potential of the crude extracts of plant materials.
- The glucose-lowering effects of *D. barnimiana* might be mediated, at least partially, through its antioxidant effect, which need further study.
- Consequently, it was necessary to perform subchronic and chronic toxicological investigation of this plant empirically, in order to avoid the risk of side effects related to phytotherapy.
- Species variation would limit a straightforward extrapolation of the findings of this study to humans.

Abbreviations

FBG, fasting blood glucose; HDL, high-density lipoprotein; LDL, low-density lipoprotein; OGTT, oral glucose-tolerance test; Stz, streptozotocin; TC, total cholesterol; TGs, triglycerides.

Data Sharing

All the data used to support the findings of this study are available from the corresponding author upon reasonable request.

Ethics

The experimental animals were handled and cared for in the experimental procedures according to the local ethical and internationally accepted laboratory animal use, care and welfare guidelines such as Basel declaration,⁶⁷ International Council for Laboratory Animal Science ethical guidelines,⁶⁸ and the EU directive⁶⁹ on the protection of animals used for scientific purposes. This study was approved by the ethical clearance which was requested and obtained from the research and ethical review committee of College of Health Sciences, Debre Tabor University with reference code number of SOP12/173/21.

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Author Contributions

All the authors have been involved in the design, conducting the research, data analysis, drafting or revising the article and write-up of the study. We have agreed on the journal to which the article was submitted, given final approval of the version to be published, and agreed to be accountable for all aspects of the work.

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