Phytochemical Screening and Antidiabetic, Antihyperlipidemic, and Antioxidant Effects of *Leptopus Cordifolius* Decne. In Diabetic Mice

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Plants are well known in traditional herbal medicines for their hypoglycemic and hypolipidemic activities and are often used due to their accessibility, affordability, and corollary effects. *Leptopus cordifolius* has been reported to control diabetes in folkloric medicine, but no known scientific research has been conducted to assess the plausibility of this assertion. Therefore, the current study is aimed to investigate the antidiabetic and hypolipidemic effects of *Leptopus cordifolius* leaves in alloxan-induced diabetic mice. The antidiabetic and antihyperlipidemic evaluation was conducted in Swiss albino mice at doses of 150–250 mg/kg for 15 days. The blood glucose, total cholesterol, triglyceride, LDL, HDL, creatinine, ALP, SGPT, and SGOT levels were estimated according to standard procedures. Phytochemicals of leaves were analyzed using GC-MS analysis. Enzymatic antioxidant activity of the plant was investigated spectrophotometrically by carrying out superoxide dismutase, peroxidase, and catalase assays. The membrane stabilization potential of *L. cordifolius* leaf extracts was carried out using an *in vitro* haemolytic assay. The results revealed a dose response effect with the methanolic extract of *L. cordifolius* which had significant antihyperglycemic effects at 150–250 mg/kg in alloxan treated mice, although less than the positive control (glibenclamide). Hyperlipidemic activity was significant at 250 mg/kg. The biochemical parameters, such as total cholesterol, triglyceride, LDL, HDL, creatinine, ALP, SGPT, and SGOT, were significantly improved (*p* < 0.01) by the methanolic extract of 250 mg/kg compared to the diabetic group. Treatment for 15 days showed significant elevation (*p* < 0.01) of antioxidant enzymes. GC-MS analysis provided tentative identifications of 52 compounds in the methanolic extract of *L. cordifolius* which had significant antihyperglycemic effects at 150–250 mg/kg in alloxan treated mice, although less than the positive control (glibenclamide). Hyperlipidemic activity was significant at 250 mg/kg. The biochemical parameters, such as total cholesterol, triglyceride, LDL, HDL, creatinine, ALP, SGPT, and SGOT, were significantly improved (*p* < 0.01) by the methanolic extract of 250 mg/kg compared to the diabetic group. Treatment for 15 days showed significant elevation (*p* < 0.01) of antioxidant enzymes. GC-MS analysis provided tentative identifications of 52 compounds in the methanolic extract of *L. cordifolius*, of which 12 compounds have reported antidiabetic activity. In conclusion, methanolic extract of *L. cordifolius* of 150 and 250 mg/kg body weight showed significant antidiabetic and antihyperlipidemic activities in alloxan-induced diabetic mice and, with further work, has the potential to be used to manage blood glucose and cholesterol levels.

**Keywords:** antidiabetic, alloxan, lipid profile, antioxidant, *leptopus cordifolius*
INTRODUCTION

Diabetes mellitus is one of the most severe and incurable metabolic disorders, characterized by an increase in blood glucose level due to an absolute or relative lack of insulin and failure of insulin to act on its target tissue (Dey et al., 2015). Chronic hyperglycemia in diabetes is associated with long-term damage, dysfunction, and failure of various organs (Bajpay, 1993). Diabetes mellitus is known to cause hyperlipidemia through various metabolic derangements. Among several metabolic derangements, insulin deficiency has been known to stimulate lipolysis in the adipose tissue and give rise to hyperlipidemia (Ahmad et al., 2014). The number of people suffering from diabetes is increasing due to urbanization, demographic growth, aging, and growing incidences of obesity and lack of exercise (Badran and Laher, 2012). The existing drug therapies of sulfonylureas, biguanides, α-glucosidase inhibitors, and glinides have challenges in terms of cost, safety, and efficacy (Omonije et al., 2019). In particular, these hypoglycemic agents may cause adverse effects like renal toxicity, hypoglycemia, hepatotoxicity, and gastrointestinal disorders (Carpio and Fonseca, 2014).

Medicinal plants in this regard have become popular because of their perceived fewer side effects and many biological activities (Peiris et al., 2015). The antidiabetic medicinal plants may provide valuable sources for finding safer hypoglycemic agents. Ethnobotanical statistics estimate that 1,200 rare plants worldwide may have antidiabetic potential (Arumugam et al., 2013).

Leptopopus cordifolius Decne (syn. Andrachne cordifolia) is a medicinal plant belonging to the family Phyllanthaceae, commonly found in the low hills of Northern hilly areas of Pakistan and the Himalayan region, including Azad Jammu and Kashmir. It is a small shrub, about 1 m tall, with ovate to elliptic leaves. It is distributed in Pakistan, Kashmir, India, and Nepal, on plains as well as on rocks, cliffs, wastelands, and on the bank of streams. The powder of the leaves of L. cordifolius is used for healing wounds and as a hypoglycemic agent (Ajaib, 2012; Munir and Qureshi, 2018). However, to the best of our knowledge, no scientific study documenting the antihyperglycemic and antihyperlipidemic behavior of L. cordifolius is available. Therefore, the current study was designed to investigate the antidiabetic ability of L. cordifolius in an alloxan-induced model of diabetic mice.

MATERIALS AND METHODS

Collection and Processing of Plant

The L. cordifolius taxonomically identified leaves were collected from District Buner, northwest of Pakistan, after confirmation of its identification and authentication. The sample specimen with voucher number (AWKB-002) was deposited in the Herbarium of Botany Department Abdul Wali Khan University Mardan for future reference. The leaves of the plant were washed and air-dried for two weeks under shade. Using an electric grinder, they were ground to a uniform coarse powder and held in an airtight container at room temperature until it was used for extraction.

Preparation of Extract

The powdered leaves of L. cordifolius were extracted by direct mollescence method using 70% methanol for 72 h. The extracts were weighed and the percentage yield was calculated by using the formula (David et al., 2017).
Experimental Animals
The Swiss albino mice of 25–35 g weight and 3–4 weeks of age of either sex were utilized in the experiment. In compliance with OECD guidelines (Guideline, 2001), experimental animals were acclimatized to laboratory conditions for one week with free access during the experimental time to commercial pellet laboratory diets and water ad libitum (Council, 2010). The experimental animal’s procedures were used with the approval of an ethical committee of the Department of Pharmacy, Abdul Wali Khan University Mardan (0015/2019). Animals under ether anesthesia were killed and sacrificed at the end of the experiment.

Diabetes Induction in Experimental Animals
Diabetes was provoked in overnight fasted mice by intraperitoneal injection of a single dose of alloxan monohydrate (150 mg/kg) (Szkudelski, 2001). The induction of diabetes was confirmed in mice 3 days after injection of alloxan by measuring the level of blood glucose. The animals with a blood glucose level greater than 200 mg/ml were considered as diabetic (Tzeng et al., 2014) and were designated for the study. The treatment with methanolic extracts, glibenclamide, and solvent fractions was initiated 72 h after alloxan injection to the mice.

Acute Toxicity Study
An acute toxicity test was conducted in conjunction with the limit test recommended by Guideline 425 of the Organization for Economic Cooperation and Development (Guideline, 2001). The tests for acute toxicity were carried out using a fixed-dose protocol. Gross behavioral changes such as tremors, traction, salivation, loss of appetite, lacrimation, hair erection, diarrhea, mortality, and other symptoms of toxicity for 15 days were observed in the mice (Alema et al., 2020).

Experimental Design
The mice were divided into nine groups consisting of six mice in each group. Group 1 served as normal control (saline, non-diabetic), group 2 served as alloxan (150 mg/kg)-treated control (diabetic control), group 3 received glibenclamide (10 mg/kg), group 4 and 5 received Leptopus Cordifolius methanolic extracts at the dose of 150 and 250 mg/kg respectively, whereas group 6, 7, 8, and 9 received n-hexane, chloroform, ethyl acetate, and aqueous fractions at the dose of 250 mg/kg orally. The treatment procedures lasted for 15 successive days. The extracts and saline solution were administered via oral gavage.

Determination of Anti-diabetic Activity
The anti-diabetic ability of extracts and solvent fractions were assessed by an in vivo oral glucose tolerance test (OGTT) using albino mice at a dose of 150 and 250 mg/kg body weight. The dose of 150 and 250 mg/kg body weight were selected based on the effectiveness of their traditional claim (Shukla et al., 2011).

Blood Glucose Determination
The blood glucose level of each mouse was determined by fractional tail amputation technique, taking blood from the tail vein by a one-touch electronic glucometer using glucose test strips (Akpan et al., 2012). The tails were then rubbed with ethanol to prevent infection.

Body Weight Analysis
The body weight of the experimental animals was recorded before commencing the treatment (day 0) and throughout the experiment (day 7 and 15), and fluctuations in weight were recorded.

Collection and Preparation of Blood Sample
Blood samples were collected from 24 h-fasted mice by cardiac puncture and retro-orbital plexus puncture method and were kept in sterile vials for 30 min for clotting. Serum samples were collected from clotted blood using a centrifuge operated at 3,000 rpm for 10 min (Chou et al., 2008; Parasuraman et al., 2015).

Collection of Organs
The mice were anesthetized under mild ether and dissected. The tissue sample was taken for assessment of biochemical parameters (Li et al., 2008).

Biochemical Analysis
The serum samples were then used for estimation of biochemical parameters such as alkaline phosphatase (ALP) (Sasaki, 1966), creatinine (Bowers, 1980), total cholesterol (TC) and triglycerides (TG) (Roeschlau et al., 1974), LDL (Friedewald et al., 1972), and HDL (Iwata et al., 1990). The Serum Glutamate Pyruvate transaminase (SGPT) and Serum Glutamic Oxaloacetic transaminase (SGOT) activities were determined according to the method of Reitman and Frankel (Reitman and Frankel, 1957). These biochemical assays were measured by using commercial diagnostic kits.

Histopathological Analysis
The liver, kidney, and pancreas of animals were expunged at the termination of the experiment and engrossed in 10% buffered formal-saline for histopathological investigations using Hematoxylin and Eosin (H&E) stains. The morphological changes in hepatic and pancreatic cells were examined under a microscope on albumenized glass slides, using 20× and 40× magnifications (Bancroft and Gamble, 2008).

Antioxidant Enzyme Assay
The spectrophotometric measurement of antioxidant enzymatic activity of Superoxide dismutase SOD (Shah et al., 2013),
TABLE 2 | Effects of L. cordifolius extracts on blood glucose level of alloxan-induced diabetic mice.

| S.No | Treatments               | Dose (mg/kg) | Blood glucose level (mg/dl) |
|------|--------------------------|--------------|-----------------------------|
|      |                          |              | Day 0          | Day 4          | Day 7          | Day 10         | Day 15         |
| 1    | Normal Saline            | 0.3 ml       | 100.4 ± 2.7     | 98.4 ± 3.2     | 97.4 ± 1.9     | 96.4 ± 4.2     | 94 ± 3.6       |
| 2    | Diabetic Control         | 0.3 ml       | 424.6 ± 2.7     | 438.8 ± 3.3    | 466.6 ± 2.7    | 474.8 ± 2.3    | 490 ± 2.7      |
| 3    | Glibenclamide            | 10           | 431.4 ± 3.3     | 402.8 ± 2.8    | 360.6 ± 3.04   | 272.2* ± 4.9   | 206.6* ± 5.4   |
| 4    | Methanolic Extract (1)   | 150          | 435.8 ± 4.4     | 412.6 ± 2.7    | 387.4* ± 3.6   | 346.6* ± 4.9   | 304.2* ± 2.8   |
| 5    | Methanolic Extract (2)   | 250          | 431.2 ± 3.9     | 408.2 ± 3.16   | 360.8 ± 3.1    | 320.8* ± 2.8   | 270.4* ± 4.5   |
| 6    | n-Hexane                 | 250          | 459.2 ± 3.4     | 458.2 ± 4.2    | 449.4 ± 3.6    | 454.6 ± 4.5    | 466.8 ± 2.16   |
| 7    | Chloroform               | 250          | 456.6 ± 3.6     | 453.8 ± 3.4    | 440.4 ± 3.04   | 429.6 ± 3.36   | 414.2 ± 4.3    |
| 8    | Ethyl acetate            | 250          | 419.8 ± 3.08    | 417.4 ± 0.04   | 421.2 ± 4.08   | 425.6 ± 2.4    | 440 ± 4.4      |
| 9    | Aqua                     | 250          | 414 ± 2.9       | 412.6 ± 3.6    | 412.8 ± 2.8    | 433.4 ± 4.03   | 456.2 ± 1.9    |

Values are expressed as mean ± SEM n = 6 in each group. Data analyzed by ONE WAY ANOVA followed by ‘s Dunnett’s multiple comparisons test. * P < 0.05; **P < 0.01; compared with diabetic control.

TABLE 3 | Effects of L. cordifolius extracts on body weight of mice.

| S.No | Treatments       | Dose (mg/kg) | Body weight (g) |
|------|------------------|--------------|-----------------|
|      |                  |              | Day 0          | Day 4          | Day 7          | Day 10         | Days 15        |
| 1    | Normal Saline    | 0.3 ml       | 22.2 ± 0.28     | 23.6 ± 0.20    | 24.6 ± 0.25    | 26.1 ± 0.47    | 26.8 ± 0.45    |
| 2    | Diabetic Control | 0.3 ml       | 29.2 ± 0.49     | 28.6 ± 0.38    | 26.3 ± 0.49    | 25.4 ± 0.38    | 24.5 ± 0.33    |
| 3    | Glibenclamide    | 10           | 30.3 ± 0.28     | 31.9 ± 0.38    | 32.8* ± 0.43   | 33.5* ± 0.31   | 34.9** ± 0.22  |
| 4    | Methanolic Extract (1) | 150   | 30.8 ± 0.27     | 31.7 ± 0.54    | 32.4* ± 0.36   | 32.6 ± 0.53    | 33.1* ± 0.36   |
| 5    | Methanolic Extract (2) | 250   | 29.3 ± 0.33     | 30.6 ± 0.93    | 32.4* ± 0.36   | 33.1* ± 0.41   | 34.6** ± 0.27  |
| 6    | n-Hexane         | 250          | 28.8 ± 0.27     | 28.2 ± 0.28    | 27.5 ± 0.24    | 27 ± 0.27      | 26.2 ± 0.31    |
| 7    | Chloroform       | 250          | 27.5 ± 0.28     | 27.3 ± 0.36    | 28.2 ± 0.16    | 28.4 ± 0.22    | 29.2 ± 0.20    |
| 8    | Ethyl acetate    | 250          | 29.5 ± 0.29     | 28.5 ± 0.31    | 27.6 ± 0.23    | 26.6 ± 0.22    | 25.7 ± 0.27    |
| 9    | Aqua             | 250          | 29.2 ± 0.20     | 28.6 ± 0.19    | 27.8 ± 0.20    | 26.7 ± 0.20    | 25.5 ± 0.27    |

Values are expressed as mean ± SEM n = 6 in each group. Data analyzed by ONE WAY ANOVA followed by Dunnnett’s multiple comparisons test. * P < 0.05; **P < 0.01; compared with diabetic control.

peroxidase POD (Khan et al., 2015), and Catalase CAT (Khan et al., 2012) was carried out at 560, 420, and 230 nm respectively.

Membrane Stabilization Assay
The membrane-stabilizing activity of the extracts was determined on human erythrocytes by collecting fresh blood samples from 10 volunteers and the absorbance was measured at 560 nm (Shinde et al., 1999; Omale and Okafor, 2008).

GC-MS Profiling
A 20 mg sample of leaf powder was dissolved in 1 ml of methanol, vortexed for 5 min, sonicated for 10 min, and then centrifuged for 5 min at 12,000 x g. The supernatant was transferred into a fresh vial and was subjected (without any dilution) to GC-MS analysis.

GC-MS analysis of the active methanolic extract of L. cordifolius was carried out by using the GC-MS instrument (Model GCMS-QP2010 Ultra, Shimadzu Co., Japan) equipped with a capillary column DB-1 (0.25 μm film × 0.25 mm i. d. ×30 m length). The instrument was operated in electron impact mode at ionization voltage (70 eV), injector temperature (230°C), and detector temperature (280°C). The carrier gas used was helium (99.9% purity) at a flow rate of 1 ml/min and about 1 μL of the sample was injected. The tentative identification of compounds from the spectral data was based on the available mass spectral records (NIST and WILEY libraries) (Swamy et al., 2015).

Statistical Analysis
All the values were expressed as mean ± Standard Error of the Mean (SEM). The results were analyzed for statistical significance using one-way ANOVA followed by Dunnett’s test with the aid of IBM Statistical Package for Social Scientist (SPSS-20). The level of significance was set at p < 0.05.

RESULTS
Extraction Yield
The extraction yield of methanolic extract and different partitions of L. cordifolius showed that the extraction with methanol had the highest yield compared to the solvent extraction (Table 1).

Acute Toxicity Study
The extract was found benign up to a dose of 2000 mg/kg body weight from the safety data obtained. The behavior of the animals during the study was closely monitored and no lethality, behavioral changes, or mortality was noticed, which is regarded as a therapeutic benefit.
### TABLE 4 | Effects of *L. cordifolius* extracts on lipid profile parameters.

| S.No | Treatments                | Dose (mg/kg) | TC (mg/dl) | TG (mg/dl) | LDL (mg/dl) | HDL (mg/dl) |
|------|---------------------------|--------------|------------|------------|-------------|-------------|
| 1    | Normal Saline             | 0.3 ml       | 181.3 ± 1.45 | 163.3 ± 4.7 | 173 ± 1.1   | 35 ± 0.5    |
| 2    | Diabetic Control          | 0.3 ml       | 183.4 ± 1.45 | 167.6 ± 1.76 | 161.9 ± 12.1 | 36.6 ± 0.4  |
| 3    | Glibenclamide             | 10           | 136.4** ± 0.9 | 145.8** ± 0.7 | 125.4** ± 0.2 | 43.4** ± 0.5 |
| 4    | Methanolic Extract(1)     | 150          | 165.1 ± 3.52 | 154.8 ± 1.9 | 142.3 ± 0.3 | 40.9 ± 1.6  |
| 5    | Methanolic Extract(2)     | 250          | 154.7** ± 3.76 | 148** ± 1.7 | 129.6** ± 0.7 | 41.2** ± 1.2 |
| 6    | n-Hexane                  | 250          | 178.4 ± 1.90 | 164.3 ± 0.6 | 147.4 ± 0.4 | 35.6 ± 0.7  |
| 7    | Chloroform                | 250          | 171.5 ± 5.7  | 165.5 ± 3.7 | 143.7 ± 0.4 | 37.3 ± 0.6  |
| 8    | Ethyl acetate             | 250          | 178.4 ± 1.81 | 165.3 ± 0.7 | 147.3 ± 0.6 | 35.8 ± 0.9  |
| 9    | Aqua                      | 250          | 178.8 ± 1.11 | 166.3 ± 1.03 | 148.4 ± 0.7 | 35.2 ± 0.4  |

Values are expressed as mean ± SD; *n* = 6 in each group. Data analyzed by ONE WAY ANOVA followed by Dunnett’s multiple comparisons test. *P < 0.05; **P < 0.01; compared with diabetic control.

### TABLE 5 | Effects of extracts of *L. cordifolius* on serum biochemical parameters.

| S.No | Treatments                | Dose (mg/kg) | SGPT (U/l) | SGOT (U/l) | ALP (U/l) | Creatinine (mg/dl) |
|------|---------------------------|--------------|------------|------------|-----------|-------------------|
| 1    | Normal Saline             | 0.03 ml      | 19.3 ± 2.08 | 23.4 ± 0.65 | 173.6 ± 6.77 | 1.05 ± 0.31       |
| 2    | Diabetic Control          | 0.03 ml      | 48.4 ± 1.45 | 43.6 ± 1.45 | 331.3 ± 1.76 | 2.3 ± 0.08        |
| 3    | Glibenclamide             | 0.03 ml      | 32.4** ± 0.36 | 25.2** ± 0.60 | 169.8** ± 1.44 | 0.98** ± 0.05   |
| 4    | Methanolic Extract (1)    | 150          | 29.3 ± 2.86  | 30.5 ± 0.57 | 280 ± 1.41  | 1.87 ± 0.03       |
| 5    | Methanolic Extract (2)    | 250          | 27.8** ± 2.60 | 26.4** ± 2.15 | 231.1** ± 2.25 | 1.26** ± 0.22   |
| 6    | n-Hexane                  | 250          | 37.3 ± 1.44  | 32.2 ± 0.17 | 280.9 ± 1.15 | 1.88 ± 0.02      |
| 7    | Chloroform                | 250          | 33.5 ± 0.92  | 29.4 ± 1.15 | 270 ± 2.96  | 1.84 ± 0.04      |
| 8    | Ethyl acetate             | 250          | 38.2 ± 1.43  | 34.2 ± 0.78 | 286.6 ± 1.55 | 1.90 ± 0.02      |
| 9    | Aqua                      | 250          | 35.9 ± 1.18  | 33.1 ± 0.31 | 296.1 ± 8.93 | 1.93 ± 0.02      |

Values are expressed as mean ± SD; *n* = 6 in each group. **P < 0.01 as compared with diabetic control at the same time (one-way ANOVA followed by Dunnett’s multiple comparison test). TC, total cholesterol; TG, triglycerides; LDL, low density lipids and HDL, high density lipids.

### TABLE 6 | Determination of antioxidant enzymes activity.

| S. No | Treatments                | Dose (mg/kg) | SOD (U/mg) | POD (U/mg) | Catalase (U/mg) |
|-------|---------------------------|--------------|------------|------------|-----------------|
| 1     | Normal Saline             | 0.03 ml      | 4.87 ± 0.24 | 6.27 ± 0.23 | 9.38 ± 0.11     |
| 2     | Diabetic Control          | 0.03 ml      | 3.72 ± 0.04 | 5.97 ± 0.02 | 8.63 ± 0.03     |
| 3     | Glibenclamide             | 0.03 ml      | 2.02** ± 0.03 | 4.80** ± 0.03 | 4.57** ± 0.40  |
| 4     | Methanolic Extract (1)    | 150          | 2.98* ± 0.03 | 3.90* ± 0.22 | 5.04* ± 0.26    |
| 5     | Methanolic Extract (2)    | 250          | 2.66** ± 0.10 | 4.62** ± 0.09 | 4.94** ± 0.60   |
| 6     | n-Hexane                  | 250          | 3.41 ± 0.12  | 2.86 ± 0.03 | 5.62 ± 0.32     |
| 7     | Chloroform                | 250          | 3.71 ± 0.21  | 3.32** ± 0.27 | 5.41 ± 0.06    |
| 8     | Ethyl acetate             | 250          | 3.12 ± 0.18  | 2.66 ± 0.14 | 5.91 ± 0.26     |
| 9     | Aqua                      | 250          | 2.95 ± 0.23  | 2.36 ± 0.17 | 5.85 ± 0.03     |

Values are expressed as mean ± SD; *n* = 6 in each group. **P < 0.01 as compared with diabetic control at the same time (one-way ANOVA followed by Dunnett’s multiple comparison test). SGPT, Serum Glutamate Pyruvate Transaminase; SGOT, Serum Glutamic Oxaloacetic Transaminase; ALP, alkaline phosphatase.

### TABLE 7 | Thrombolytic activity of *L. cordifolius* extract/fractions.

| S.No | Extract/Fractions | Concentration (mg/ml) | Thrombolytic activity | % Inhibition |
|------|-------------------|-----------------------|-----------------------|-------------|
| 1    | Methanolic        | 1                     | 0.1678 ± 0.06         | 76          |
| 2    | n-Hexane          | 1                     | 0.6016 ± 0.03         | 14.6        |
| 3    | Chloroform        | 1                     | 0.4411 ± 0.05         | 37.4        |
| 4    | Ethyl acetate     | 1                     | 0.5016 ± 0.04         | 28.8        |
| 5    | Aqueous           | 1                     | 0.5737 ± 0.19         | 18.6        |
| 6    | Acetyl salicylic  | 0.10                  | 0.706 ± 0.06          | 88          |

Values are expressed as mean ± SEM. Data analyzed by ONE WAY ANOVA followed by Dunnett’s multiple comparisons test. *P < 0.05; **P < 0.01, compared with diabetic control. Data are expressed as mean ± SD (*n* = 3).
Blood Glucose Level

The administration of the methanolic extract of *L. cordifolius* at 150 and 250 mg/kg significantly reduced \( p < 0.01 \) the blood glucose level in diabetic mice at the end of the experiment in a dose dependent fashion, but less than glibenclamide-treated mice. Other extracts were not effective. (Table 2).

Changes in Body Weight and Fasting Blood Glucose Level

The methanolic extracts at the dose of 150 and 250 significantly improved the body weight \( p < 0.05; p < 0.01 \) in alloxanised diabetic mice and produced maximum fall \( p < 0.01 \) in the FBG levels of diabetic mice after 15 days of treatment (Table 3). Other extracts were not effective.
The lipid profile in diabetic mice was severely disturbed in diabetic mice by alloxan administration when compared with the control group. The methanolic extract at the dose of 150 and 250 mg/kg of *L. cordifolius* significantly reduced the levels of lipid parameters to near normal (Table 4). Other extracts were not effective.

Serum Profile
A significant elevation in SGPT, SGOT, ALP, and creatinine was observed in alloxan-induced diabetic mice when compared to normal mice. Oral administration of *L. cordifolius* leaves, methanolic extract 150 and 250 mg/kg, and glibenclamide treatment significantly improved the above parameters (Table 5).

Antioxidant Enzymes
Diabetes mellitus significantly reduced antioxidant enzymes, like catalases (CAT), peroxidases (POD), and superoxide dismutase (SOD) levels, and elevated the action of reactive oxygen species. The methanolic extract at the dose of 150 and 250 mg/kg significantly increased the activities of antioxidant enzymes. However, other extracts treated group showed no significant difference in comparison to the glibenclamide-treated group (Table 6).

Membrane Stabilizing Activity
In the hypotonic-solution-induced condition, methanolic extract showed the highest protection of erythrocyte membrane (76%), which was comparable to haemolysis inhibited by standard acetyl salicylic acid (Table 7).

GC-MS Analysis
The results of GC-MS analysis revealed the tentative identity of 52 compounds present in methanolic extract *L. cordifolius*. The active principles with their retention time, probability, molecular weight, and concentration (peak area %) are presented in Table 8 and Figure 1.
fractions of *L. cordifolius* in alloxan-induced diabetic mice. Alloxan is a cytotoxic glycoside and acts as a diabetogenic by the annihilation of the islets of Langerhans and causes an enormous decline in insulin discharge, thus prompting hyperglycemia (Grover et al., 2000). Glibenclamide is used as a standard antidiabetic drug in alloxan-induced mild diabetes to compare the antidiabetic effects of a variety of bioactive compounds (Ramkumar et al., 2011). The extracts were found to be safe up to the dose of 2000 mg/kg in normal healthy mice from the safety data obtained (Yehya et al., 2019).

Alloxan induces diabetes through ROS that leads to rapid destruction of pancreatic beta cells, causing hyperglycemia (Stanely et al., 2000). Hyperglycemia in turn increases the generation of free radicals by glucose auto-oxidation (Bajaj and Khan, 2012). In the current research study, oral administration of *L. cordifolius* methanolic extract of 150 and 250 mg/kg body weight to diabetic mice for 15 days significantly reduced the fasting blood glucose and glycercides were reduced only at 250 mg/kg, compared with diabetic control mice. In support of our study, the *L. cordifolius* methanolic extract (250 mg/kg body weight) cause their anti-hyperglycemic activity by augmenting the production of insulin or regenerating pancreatic cells (Patel et al., 2012). Although the effective doses of 150 and 250 mg/kg reported here are comparable with other antidiabetic plants (Ghazanfar et al., 2014; Alema et al., 2020), they are at the upper end of the dose range and further research needs to be done to concentrate the active principles and thereby reduce the effective dose. Unfortunately, the results with different solvents did not achieve this result as they were less effective than methanol. The observed decline in body weight of diabetic mice could be attributed to the surging body mass index and the progression of skeletal size and epididymis fats (Wu and Huan, 2008). However, treatments with orally administered methanolic, and chloroform extracts of 250 mg/kg significantly improved the body weight compared with diabetic control, which signifies its protective effect in controlling muscle wasting i.e. reversal of gluconeogenesis (Mistry et al., 2017). In diabetes mellitus, hyperlipidemia occurs as a result of the excess mobilization of fats from the adipose tissue due to the underutilization of glucose (Akpan et al., 2012). Hyperglycemia is accompanied by a rise in TC, TG, and LDL, and a fall in HDL levels (Gao et al., 2009). The methanolic extract *L. cordifolius* and glibenclamide might act on reserved fats and inhibit the release of free fatty acids, decreasing the total cholesterol and triglyceride levels, and increasing the HDL level.

It is acknowledged that diabetes is linked to irreversible hepatic damage leading to the proliferation of different enzymes including SGOT, SGPT, ALP, and creatinine in the blood (Pari and Latha, 2002; Preethi and Kuttan, 2009). *L. cordifolius* methanolic extract significantly reduced the serum creatinine, thus treating and preventing the progression of renal damage in diabetic mice. Reports have shown that the reversal of such changes in the above parameters is allied with the presence of bioactive constituents in the extract (Lal et al., 2009; Sharma et al., 2019). In diabetes mellitus, high glucose can inactivate antioxidant enzymes SOD, POD, and, CAT by glycating these proteins, thus producing induced oxidative stress, which in turn causes lipid peroxidation. The *L. cordifolius* methanolic extract increased the activities of SOD, POD, and CAT, thus maintaining the antioxidant status in the erythrocytes of diabetic mice. Several studies revealed that injury to membranes of RBCs due to hyperglycemia makes cells more susceptible to secondary damage through lipid peroxidation, haemolysis, and oxidation of haemoglobin (Halliwell, 1999). The inhibition of the erythrocyte lysis property of *L. cordifolius* could be the possible mechanism for its anti-diabetic activity. It may also be due to the presence of plant biochemical compounds in the extract which exert a profound stabilizing effect on lysosomal membranes and their cation-binding ability (Oyedapo et al., 2004).

Various secondary metabolites and phytochemicals isolated from diverse plant species have been found to have potent antihyperglycemic and glucose suppressive effects (Sharma et al., 2010). The secondary metabolites which were tentatively identified in *L. cordifolius* crude methanolic extract may be accountable for its observed glucose suppressive and antihyperglycemic activity, due to its ability to stimulate glycogenesis in the liver, release insulin from pancreatic β-cells, or inhibit glucose absorption in the gut (Sezik et al., 2005).

Several studies revealed that injury to membranes of RBCs makes cells more susceptible to secondary damage through lipid peroxidation, haemolysis, and oxidation of haemoglobin (Halliwell and Whitman, 2004). The inhibition of the erythrocyte lysis property of *L. cordifolius* could also be the possible mechanism for its anti-diabetic activity (Chaitanya et al., 2011). GC-MS analysis of methanolic extract of *L. cordifolius* tentatively identified the presence of various bioactive compounds with different retention times. The tentatively identified compounds are known to possess several biological and pharmacological activities (Grover et al., 2002; Kumar et al., 2010; Hema et al., 2011). Several studies have shown that β-Sitosterol, stigmasterol, betulin, ergost-8(14)-en-3-ol, n-hexadecanoic acid, and palmitic acid have been found to have hypoglycemic effects by reducing the absorptions of cholesterol from the gut as shown in the Table 9 (Ikeda et al., 1988; Rajasekaran et al., 2004).

**CONCLUSION**

The current research study demonstrated the first pharmacological insight into the antioxidant, antidiabetic, and antihyperlipidemic potential of the *L. cordifolius*. Based on the results exhibited by this study, it is concluded that *L. cordifolius* methanolic extract had significant antihyperglycemic activity in alloxan-induced diabetic mice compared with that of glibenclamide, as evident from the restoration of blood glucose levels, thus validating its traditional utilization. Future research needs to focus on bioassay guided isolation and confirmed identification of the active principles using standard or spectral workup. This could inform better extraction and formulation procedures to lower the dose required, as well as lead to a validated phytochemical analysis of a standardized product.
Further study of the mode of action of active principles is warranted.

**DATA AVAILABILITY STATEMENT**

The original contributions presented in the study are included in the article/Supplementary Material, further inquiries can be directed to the corresponding author.

**ETHICS STATEMENT**

The animal study was reviewed and approved by Veterinary Research Institute, Peshawar, Pakistan.

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**AUTHOR CONTRIBUTIONS**

SR and GJ designed, performed the experiment, and wrote the original manuscript. Statistical analysis, results, and discussion were accomplished by SR and FJ. HR assisted during data collection, visualization, characterizations, editing, and reviewing of the manuscript. All the authors read and approved the final manuscript.

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**Conflict of Interest:** The authors declare that the research was conducted in the absence of any commercial or financial relationships that could be construed as a potential conflict of interest.

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