Research Article

Isolation of Thymol from *Trachyspermum ammi* Fruits for Treatment of Diabetes and Diabetic Neuropathy in STZ-Induced Rats

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Terpenoids and phenols from *Trachyspermum ammi* (*T. ammi*) have reported some pharmacological actions. The objective of the work was to isolate the active constituent, its identification by spectroscopic techniques, and evaluation of the antidiabetic and neuroprotective activity from *T. ammi* on STZ Wistar rats. The dried fruits of *T. ammi* were kept in a hydrodistillation apparatus to collect essential oil. The isolated fraction went through TLC, UV, FTIR, HPLC, HRMS, C\(^{13}\), and \(^{1}\)H NMR for characterization. Two dosage concentrations from the isolated compound were prepared as 10 and 20 mg/kg for treatment groups. The groups were tested for thermal and mechanical hyperalgesia, writhing, grip strength, spontaneous locomotor test, neuromuscular coordination tests, and histopathological and lipid profile analysis. Diabetes was induced by streptozotocin (45 mg/kg i.p.) and 12 weeks of treatment-induced diabetic neuropathy in Wistar rats. Biomarkers were evaluated to understand the neuropathic protection of thymol on STZ-treated Wistar rats. The biomarker studies (SOD, NO, LPO, Na\(^{+}\)K\(^{-}\)ATPase, and TNF-\(\alpha\)) further confirmed thymol’s diabetic neuropathy protective action. This study suggests that isolated compound thymol was antidiabetic and neuroprotective as it has shown controlled glucose levels defensive nerve damage in STZ Wistar rats. \(P < 0.05\) level of significance was observed in the levels of endogenous biomarkers, fasting blood glucose levels, actophotometer response, and response latency in treated groups compared to the diabetic group, whereas \(P < 0.001\) level of significance during lipid profile levels, thermal algesia, and neuromuscular comparison tests was noted in treated groups compared to the diabetic group.

1. Introduction

Patients suffering from diabetes mellitus (DM) observe a severe condition of peripheral nerve dysfunction called diabetic neuropathy (DN) [1, 2]. In India, the prevalence is higher (4.3%) compared to the western countries where 1%–2% population of DM faces these conditions. This could be due to the probability that Asian Indians are prone to the condition of insulin resistance [3–5]. Around 2/3rd of the diabetic population suffers from clinical or subclinical neu-
apotrophic conditions, where approximately 10% of the people with diabetes face persistent pain. Diabetic neuropathy has a common feature of spontaneous pain, intractable or stimulus-induced pain [6]. There are many categories of DN where diabetic polyneuropathy (DPN) is a condition that prevails for a lifetime and is a major cause of nerve injury, foot ulceration, gait disturbance, and amputation [7–11]. The central control observed for DN is preventive management, as the check on the glycemic control helps prevent neuropathic complications [12].

The negative symptom observed in patients suffering from DN is a decrease in sensation-induced numbness, whereas the positive symptoms observed are aching, prickling, and burning sensations [13]. Mechanisms suggest that the tiny and unmyelinated nerve fibres are responsible for conveying sensations like temperature, touch, and pain, while the long white fibres get sensations of joint position and vibratory senses. Most of the patients report mild discomfort but around 25% report painful neuropathic conditions in diabetes. Pain in diabetic neuropathy often worsens at night [1, 14–16].

Diabetes has gradually become the most prominent problem in the global healthcare issue of the 21st century. The population of diabetics is predicted to double between the years 2000 to 2030 by reaching to level of 366 million people [17, 18]. This metabolic disorder shows prominence of hyperglycemia and includes defects of insulin productions or insulin secretion or both. Cases of chronic hyperglycemia in diabetes lead to dysfunctions and damages in the kidney, nerves, eyes, heart, and blood vessels. Prominent symptoms of hyperglycemia include weight loss, polydipsia, polyuria, blurred vision, and sometimes polyphagia. Susceptibility to infections and impairment of growth are chronic effects of prolonged hyperglycemia. In contrast, the acute fatal consequences of diabetes are nonketotic hyperosmolar syndrome and hyperglycemia with ketoadiposis [19, 20].

Prime causes of Type 2DM are genetics and lifestyle factors [21]. Advancing age is also a risk factor of T2DM, but the increasing incidences of obesity in childhood have resulted in prominent cases of T2DM in adolescents, children, and teenagers which is a serious concern and an emerging public health epidemic [22].

*T. ammi* is commonly known as “Ajwain” and is found in major parts of India, dominantly in Rajasthan and Gujarat regions. The plant belongs to the family Umbelliferae and also is called Omum, Ajowan, in Sanskrit [23, 24], Ajwain in German, Kammun or Al-Yunan in Arabic, Hounastan in Armenian, and Xi Ye Cao Guo Qin in Dutch. This plant is native to Indian region, Pakistan, Iraq, and Afghanistan but also grows in regions near Mediterranean Sea, Egypt, and southwest Asia. The herb possesses many medicinal values and is used traditionally for curing conditions like atonic dyspepsia, abdominal tumors, lack of appetite, flatulence, bronchial problems, abdominal pains, diarrhea, piles, asthma, and amenorrhea. Many pieces of research have proved the plant to be antinociceptive, hypolipidemic, antimicrobial, abortifacient, nematicidal, antioxi-
dant, antifungal, antispasmodic, anti-hypertensive, antilithiasis, diuretic, antitussive, antifilarial, abortifacient, hypolipidemic, and cytotoxic and having bronchodilating actions [25].

Currently, a huge inclination to the herbal sources is noted in the case of many treatments. Since the ancient era, humans have depended on natural sources as remedies for diseased conditions. In India, Ayurvedic remedies are used as treatments for ages. Due to reported adverse reactions in synthetic medications and temporary relief, a huge population has drifted to natural and herbal sources for curing ailments. There has been a lot of reaches where herbs and their extracts have been proved useful in the treatment of diseases [26–29].

Based on the literature review, we found no study on diabetes and diabetic neuropathy of thymol isolated from dried fruits of *T. ammi*. Therefore, we planned to isolate thymol (an active constituent) from the dried fruits of *Trachyspermum ammi* and evaluate its effects on diabetes and diabetic neuropathy by experimentally inducing it on animals.

2. Material and Methods

2.1. Drugs and Chemicals. Streptozotocin was acquired from Sigma-Aldrich India, while Sanoﬁ India Ltd. provided gliben-
clamide for a research work. The chemicals were of analytical grade obtained from the laboratory facility of the institute.

2.2. Instruments. UV-1700 PharmaSpec on SHIMADZU, FTIR–Spectrum Two on PerkinElmer, and HPLC on LC-2010CHT by Shimadzu were used from PSIT, Kanpur laboratory facility for performing UV and FTIR spectroscopy, while C13 NMR and 1H NMR were performed by Indian Institute of Technology, Kanpur laboratory facility, where DMSO–D6 was used as solvent. The HRMS was done by SAIF CDRI, Lucknow, where ethanol was used as a solvent. A glucometer (Accu-chek®) was used to estimate blood glucose levels and a semiauto analyzer (Remi Industries Ltd., India) was used for biochemical analysis.

2.3. Test Animal. Adult rats weighing (180–220 g) of both genders were obtained from the animal house of Institute of Pharmacy, PSIT, Kanpur. Animals were kept in cages (polyacrylic) which were spacious and large. An ambient room temperature was maintained with 12 h light/12 h dark cycle. Purified water and standard pellet diet ad libitum were made available to experimental rats [30]. This study was approved by IAEC, and the experiment was performed in accordance to CPCSEA (Ref No. 1273/PO/Re/S/09/ CPCSEA). The experimentation design chosen is mentioned in Table 1, and the process is explained in Figure 1.

2.4. Collection and Identiﬁcation of Plant Material. The fruit of *T. ammi* was obtained from the local market in Kanpur, Uttar Pradesh, India (Figure 2). They were collected in November 2018. The powdered fruit underwent hydrodistillation and was purified to obtain an isolated compound (Figure 3).

2.5. Identiﬁcation of Compound by TLC. The TLC of the crystalline material obtained after hydrodistillation and purification showed an *Rf* value of 0.52. The system of TLC used had silica gel (stationary phase) and benzene–chloroform (3:1 *V/V*) as the mobile phase.
2.6. Identification of Compound by $^1$H NMR, C$^{13}$ NMR, HPLC, HRMS, UV, and FTIR Analysis. The crystalline material obtained from the Crude Essential oil collected was taken for UV, HPLC, and FTIR spectroscopy studies using ethanol as a solvent. These studies were carried out in the Department Pharmacy of PSIT (Kanpur), and the results obtained are linked. The HRMS for the estimation of compound molecular weight was performed by SAIF, CDRI, Lucknow, using ethanol as a solvent. The $^1$H NMR and C$^{13}$ NMR were performed by the Indian Institute of Technology, Kanpur, where DMSO–D$_6$ was used as a solvent. The melting point was registered as 50°C using a melting point apparatus. On analysing the TLC results and all spectroscopies, the interpretation of the compound was found to be thymol (Figure 4).
2.7. Acute Toxicity Study. The ethanolic extract of *Trachyspermum ammi* was checked up to a dose of 3200 mg/kg orally, and it was hence confirmed that the lethal dose was beyond 3200 mg/kg (b.wt) [31]. The acute toxicity experiments were done in accordance with guideline no. 425 of the OECD. The treated rats were given isolated compound doses of 5, 50, 300, and 2000 mg/kg, respectively. The dose chosen for the activity was 10 and 20 mg/kg (b. wt.) of compound isolated. Rats were then checked for their behaviour like irritability, alertness, restlessness, and fearfulness. They were also checked for neurological abnormalities like pain response, touch response, gait and spontaneous activity, and autonomic reflex-like urination as well as for defecation for 24 hours. 14 days from the administration of dose, the animals were observed for any case of mortality, and on the 14th day, they were sacrificed to isolate organs to note any visible morphological changes or signs of toxicity (Figure 5) [32–34].

2.8. Induction of Diabetes Mellitus. An intraperitoneal route was used for administration of streptozotocin (Sigma-Aldrich) at a dose of 45 mg/kg post overnight fasting of at least 12 hours to induce hyperglycemia. The hyperglycemia was confirmed post 72 hours of STZ injection and was marked by increased blood glucose levels. After the induction of diabetes, the animals were randomized as per their body weight and glucose levels thus having 4 groups according to protocol. A control group was established where diabetes was not induced. Estimation of fasting blood glucose on 7th, 14th, 21st, and 28th days was done via a glucometer from Accu-chek® [35].

2.9. Induction of Diabetic Neuropathy. Diabetes was induced, and 12 weeks after the experimentation, its effects were evaluated on the nerves to find out the possibilities of generation of diabetic neuropathy. The neuropathic development was assessed with the help of various parameters like locomotor tests and examination of the nerve tissue morphologically and microscopically on weeks 2, 4, 8, and 12 in all animals [37].

2.9.1. Estimation of Fasting Blood Glucose. In each group, rats were fasted one day prior to testing their glucose levels on days 7, 14, 21, and 28 of experimentation. Glucose levels were obtained by collecting small amount of blood from tail and tested via a diagnostic kit [35].

2.9.2. Body Weight, Food, and Water Consumed. The body weight for each animal, consumption of food, and water were monitored during the experiment, recorded, and compared [36].

2.9.3. Estimation of Lipid Profile. Serum was separated by the help of centrifugation process by using centrifuge at a speed of 15000 rpm for 10 min—Remi Industries Ltd. (Mumbai, India). The isolated serum was analysed for TC, LDL, HDL, and triglyceride analysis using a semiautoanalyser (Span Diagnostics Ltd., India). The lipid profile estimations were performed at the end of experimentation, and the blood was collected from the retroorbital method.

2.10. Behavioural Studies

2.10.1. Thermal Hyperalgesia. Animals were kept on the analgesimeter (hot plate device) manufactured by Columbus instruments. 55 ± 1°C was set as the device temperature for the experiment. Initially, the symptoms like jumping or licking of paws were recorded and the reaction time in seconds was noted. Cut-off time for the investigation was 10 seconds to prevent any damages to rat paws. Eddy’s hot plate method was used for testing the hyperalgesia on weeks 2, 4, 8, and 12 in all animals [37].

2.10.2. Writhing Responses. Assessment of neuropathic pain was done by recording writhing responses in animals. 1% *v/v* acetic acid (in distilled water) was administered to rats in a volume of 0.1 ml/10 g body weight to stimulate writhing responses. All episodes of stretching of back, elongation of body, and limb extensions were noted and counted in all animals [38].

2.10.3. Cold Hyperalgesia. The acetone drop test was conducted to evaluate the cold sensitivity of the rats. Animals from every group were kept in separate mesh cages for acclimatization. Fresh drops of 50 μl acetone were then applied on the midplantar surface of paws gently. The stimulation of cold was created around 2-5 sec of application, and then the responses like mild paw withdrawal, shaking, rubbing, or licking of paw were noted. These reactions were noted as nociceptive responses, no responses (antinociceptive effects), or delayed responses (delayed nociception) in terms of minutes on both paws at intervals of 5 minutes; hence, the mean response was noted [39].

2.10.4. Evaluation of Mechanical Hyperalgesia. A pinprick test was performed on rats (hind paw) where the paw was pricked with a 900 C bent gauge needle but no piercing was done. Reflexes were recorded for paw withdrawal post-gentle prick, and responses were recorded in seconds. The cut-off time was 20 sec. Results were noted on weeks 2, 4, 8, and 12 of experimentation [40].

2.10.5. Grip Strength. The tests were performed to determine the neuromuscular strength of animals by dangling them on forelimbs with metal wire held tight on the poles. Duration of falling was recorded before falling on the surface hence estimating their muscle strength. Time was noted, and the interpretations were made on the basis of time of fall (as a week or damaged muscles will lead to less time of fall) [41].
2.10.6. Spontaneous Locomotor Testing. This test was performed by introspecting the animal’s behaviour and activity in an actophotometer. Animals were kept in a 30 × 30 × 30 cm actophotometer with photocells and a digital counter to record the interruptions in the beam during activity recording [42].

2.10.7. Estimation of Neuromuscular Coordination. A motor coordination test was performed using a rotarod apparatus where rotarod was operating on 25 rpm speed. Cut-off time was 5 minutes, and fall-off time was recorded for all animals during the test. The low fall of time reciprocates to damaged nerve and muscle [43].

2.11. Histopathological Study. Tissues were isolated from the liver and sciatic nerves of rats at the termination of experiment. On week 12 of the study, the sciatic nerve from the lower limb’s thigh region and the liver from the study animal were isolated, washed, and then stored (Figure 6). The nerve and liver were cleansed and stored in 4% formalin solution with pH 6.9. They were kept at freezing temperatures and then, the histopathological analysis was carried out. The samples were precisely cut and isolated and then stained with eosin and hematoxylin [44, 45].

2.12. Biochemical Estimations

2.12.1. Preparation of Homogenate of Sciatic Nerve. At the termination of the procedure, animals were exposed to a high dose of anesthesia to isolate the sciatic nerve. Homogenate of tissue was prepared using a buffer solution (0.1 M
Tris-HCl) with a pH of 7.4. The supernatant fluid of the homogenate was used for the estimation of membrane-bound inorganic phosphate, SOD (superoxide dismutase), lipid peroxidation (MDA content-LPO), TNF-α, and NO content (nitric oxide) [46, 47].

2.12.2. Evaluation of TNF-α. Quantification of TNF-α was done using a Thermo Scientific immunoassay kit for estimation of TNF-α levels in rat solid-phase ELISA for a duration of 4.5 h. A microplate was precoated with a monoclonal antibody, and 50 μl buffer was added to each well plate.

Samples from homogenate obtained from sciatic nerve of each group were incubated at RT for 1 hour. Immobilized antibodies will bind to TNF-α present in the well.

Washing of sample was done followed by pouring of biotinylated antibody reagents into all wells, respectively, and was then incubated at RT for 1 hour. It was further followed by washing. Then, 100 μl of streptavidin-HPR reagent was added in each well. Antibody-enzyme was then added to remove any traces of unbound antibody. 100 μl of 3,3′,5,5′
The color intensity was noted at 550 nm [46, 47].

3.1. Isolation. During the characterization of the isolated compound, it was analysed that UV \( \lambda \) max = 272.40; HPLC: retention time 9.94; HRMS-151 – 201, Molecular Ion Peak, 108 – Fragment Ion Peak; IR (KBr, cm -1): 1610 (C=C str., Aromatic ring), 1422 (O-H ben.), 1236.53 (C-O str.), 1227.02 (O-H str, Phenol), 2955.48 (C-H str), 1422 (O-H ben.), 1236.53 (C-O str.).

2.13. Statistical Analysis. Experiments were done by grouping animals as 6 in each group. Quantitative data was tabulated via ANOVA (one-way) method, and post hoc analysis of data was done via Tukey-Kramer’s test.

3. Results

3.1. Isolation. During the characterization of the isolated compound, it was analysed that UV \( \lambda \) max = 272.40; HPLC: retention time 9.94; HRMS-151– Molecular Ion Peak, 108 – Fragment Ion Peak; IR (KBr, cm -1): 1610 (C=C str., Aromatic ring), 1422 (O-H ben.), 1236.53 (C-O str.), 1227.02 (O-H str, Phenol), 2955.48 (C-H str). One-way ANOVA; Tukey-Kramer’s multiple comparisons test: "P < 0.05 in comparison with normal control; and "P < 0.05 in comparison with the diabetic control.

Values are expressed as the mean ± SEM; n = 6. One-way ANOVA; followed by Tukey-Kramer’s multiple comparisons test: "P < 0.05 in comparison with normal control and "P < 0.05 in comparison with the diabetic control.

3.2. Acute Toxicity. 14 days from the dose of administration, the animals were observed for any case of mortality. On the 14th day, they were sacrificed to isolate organs to note any visible morphological changes or signs of toxicity. The rats from all groups were checked for neurological abnormalities and autonomic reflexes. No exceptions were reported, and all results were normal. The organs isolated were observed for any morphological changes, and no changes were found. The results are recorded in Figure 5.

3.3. Fasting Blood Glucose. Glucose levels rise uncontrollably in diabetic groups which were noticed as 308.27 ± 0.37 in Group II animals (Table 2). The levels of glucose were measured in all groups where it was found that Groups IV and V have shown 129.01 ± 3.45 and 117.12 ± 1.35 mg/dl concentrations, respectively, which shows improvement in glucose levels when glucose levels were measured in all groups where it was found that Groups IV and V have shown 129.01 ± 3.45 and 117.12 ± 1.35 mg/dl concentrations, respectively.

Table 2: Fasting blood glucose testing during the antidiabetic study for 28 days.

| S. No. | Treatment type | Day 7 | Day 14 | Day 21 | Day 28 |
|--------|----------------|-------|--------|--------|--------|
| 1      | Normal control | 95.4 ± 0.18 | 96.35 ± 1.76 | 97.24 ± 2.04 | 97.03 ± 2.19 |
| 2      | Diabetic control | 157.21 ± 0.32a | 261.18 ± 2.24a | 297.12 ± 1.37a | 308.27 ± 0.37a |
| 3      | Standard | 154.01 ± 0.14* | 153.13 ± 1.91* | 141.32 ± 1.41* | 121.18 ± 3.23* |
| 4      | Treated 1 | 152.14 ± 1.62* | 159.37 ± 0.37* | 149.38 ± 3.27* | 129.01 ± 3.45* |
| 5      | Treated 2 | 1521.53 ± 1.61* | 152.16 ± 2.81* | 135.14 ± 2.47* | 117.12 ± 1.35* |

Values are expressed as the mean ± SEM; n = 6. One-way ANOVA; followed by Tukey-Kramer’s multiple comparisons test: "P < 0.05 in comparison with normal control; and "P < 0.05 in comparison with the diabetic control.

Table 3: Record of the body weight postinduction of diabetes in Wistar rats on weeks 2, 4, 8, and 12.

| S. No. | Treatment type | 2nd week | 4th week | 8th week | 12th week |
|--------|----------------|-----------|-----------|----------|----------|
| 1      | Normal control | 152.3 ± 2.03 | 168.27 ± 2.21 | 190.25 ± 5.13 | 231.45 ± 0.26 |
| 2      | Diabetic control | 154.10 ± 2.40a | 149.15 ± 1.32a | 144.27 ± 1.53a | 132.36 ± 3.26a |
| 3      | Standard | 153.13 ± 1.26 | 150.13 ± 0.37 | 159.23 ± 1.32*** | 199.2 ± 1.12*** |
| 4      | Treated 1 | 155.24 ± 1.32 | 163.4 ± 2.24*** | 168.25 ± 0.25*** | 201.34 ± 1.38*** |
| 5      | Treated 2 | 159.24 ± 1.35 | 166.14 ± 2.14*** | 182.17 ± 1.34*** | 211.23 ± 1.42*** |

Values are expressed as the mean ± SEM; n = 6. One-way ANOVA; Tukey-Kramer’s multiple comparisons test: "P < 0.001 in comparison with normal control; and ***"P < 0.001 in comparison with the diabetic control.

Table 4: Assessment of food and water consumption post treatment in Wistar rats.

| S. No. | Treatment type | Water intake (ml/day) | Food intake (g/day) |
|--------|----------------|----------------------|---------------------|
| 1      | Normal control | 13.46 ± 0.82 | 18.21 ± 1.34 |
| 2      | Diabetic control | 36.25 ± 6.34a | 27.26 ± 0.31a |
| 3      | Standard | 21.17 ± 2.35*** | 21.14 ± 2.62*** |
| 4      | Treated 1 | 17.42 ± 1.35*** | 22.47 ± 1.37*** |
| 5      | Treated 2 | 16.13 ± 0.59*** | 21.02 ± 0.15*** |

Values are expressed as the mean ± SEM; n = 6. One-way ANOVA; Tukey-Kramer’s multiple comparisons test: "P < 0.001 in comparison with normal control; and ***"P < 0.001 in comparison with the diabetic control.

- Tetramethylbenzidine (TMB)—a substrate solution—was poured which changed the color from blue to yellow. The color intensity was noted at 550 nm [46, 47].
Table 5: Effects on lipid profile of groups in Wistar rats.

| S. No. | Treatment type   | TC   | Triglyceride | LDL  | HDL  | VLDL |
|--------|-----------------|------|--------------|------|------|------|
| 1      | Normal control  | 74.29 ± 0.35 | 65.16 ± 0.28 | 36.29 ± 0.19 | 27.04 ± 4.15 | 13.28 ± 0.27 |
| 2      | Diabetic control| 143.29 ± 0.25² | 157.21 ± 0.28² | 94.19 ± 0.14² | 15.19 ± 0.42² | 46.38 ± 0.25² |
| 3      | Standard        | 77.23 ± 2.36∗∗∗ | 65.48 ± 1.49∗∗∗ | 37.16 ± 0.25∗∗∗ | 29.8 ± 2.46∗∗∗ | 15.12 ± 1.38∗∗∗ |
| 4      | Treated 1       | 85.02 ± 1.91∗∗∗ | 72.26 ± 2.43∗∗∗ | 45.26 ± 2.36∗∗∗ | 20.35 ± 2.26∗∗∗ | 21.13 ± 1.27∗∗∗ |
| 5      | Treated 2       | 74.59 ± 0.24∗∗∗ | 65.15 ± 1.28∗∗∗ | 37.24 ± 2.29∗∗∗ | 28.03 ± 0.29∗∗∗ | 14.01 ± 0.13∗∗∗ |

Values are expressed as the mean ± SEM; n = 6. One-way ANOVA; Tukey-Kramer’s multiple comparisons test: *P < 0.001 in comparison with normal control; and **P < 0.001 in comparison with the diabetic control.

Table 6: Thermal hyperalgesia: Eddy’s hot plate test on animals in all groups to assess diabetic neuropathy in Wistar rats on weeks 2, 4, 8, and 12.

| S. No. | Treatment type | Reaction latency (2nd week) | Reaction latency (4th week) | Reaction latency (8th week) | Reaction latency (12th week) |
|--------|---------------|-----------------------------|-----------------------------|-----------------------------|-------------------------------|
| 1      | Normal control| 4.12 ± 1.43                 | 4.19 ± 1.28                 | 4.01 ± 1.03                 | 4.01 ± 1.45                   |
| 2      | Diabetic control| 6.77 ± 1.28⁴               | 5.19 ± 1.48⁴               | 8.23 ± 2.92⁴               | 9.85 ± 1.86⁴                 |
| 3      | Standard      | 6.25 ± 3.14                 | 5.30 ± 4.19                 | 5.89 ± 1.76∗∗∗             | 5.91 ± 1.74∗∗∗               |
| 4      | Treated 1     | 6.92 ± 0.29                 | 6.55 ± 1.32∗∗∗             | 5.18 ± 0.21∗∗∗             | 6.08 ± 1.41∗∗∗              |
| 5      | Treated 2     | 6.01 ± 0.48                 | 4.41 ± 1.34                 | 5.11 ± 1.65∗∗∗             | 5.93 ± 0.24∗∗∗              |

Values are expressed as the mean ± SEM; n = 6. One-way ANOVA; followed by Tukey-Kramer’s multiple comparisons test: *P < 0.001 in comparison with normal control; **P < 0.001 in comparison with the diabetic control.

Table 7: Recoding the writhing responses to assess the neuropathic pain in all groups of Wistar rats on weeks 2, 4, 8, and 12.

| S. No. | Treatment type | Number of writhing responses in ten minutes (counts) | (2nd week) | (4th week) | (8th week) | (12th week) |
|--------|---------------|----------------------------------------------------|------------|------------|------------|-------------|
| 1      | Normal control| 10.01 ± 0.12                                      | 11.16 ± 0.22 | 10.24 ± 0.22 | 8.13 ± 1.32 |             |
| 2      | Diabetic control| 11.14 ± 1.44⁴                                      | 10.22 ± 6.13⁴ | 12.58 ± 4.80⁴ | 13.35 ± 0.35⁹ |             |
| 3      | Standard      | 12.35 ± 2.24                                      | 9.15 ± 2.34  | 9.64 ± 2.32∗∗∗ | 8.94 ± 1.35∗∗∗ |             |
| 4      | Treated 1     | 14.94 ± 4.25                                      | 9.24 ± 1.34  | 9.94 ± 1.31∗∗∗ | 8.28 ± 1.44∗∗   |             |
| 5      | Treated 2     | 13.32 ± 1.32                                      | 9.14 ± 2.11  | 9.61 ± 0.21∗∗∗ | 8.24 ± 0.17∗∗∗ |             |

Values are expressed as the mean ± SEM; n = 6. One-way ANOVA; Tukey-Kramer’s multiple comparisons test: *P < 0.001 in comparison with normal control; **P < 0.001 in comparison with diabetic control.

Table 8: Cold hyperalgesia: acetone drop test in all groups in Wistar rats on weeks 2, 4, 8, and 12.

| S. No. | Treatment type | Cold hyperalgesia (acetone drop test) in sec | (2nd week) | (4th week) | (8th week) | (12th week) |
|--------|---------------|---------------------------------------------|------------|------------|------------|-------------|
| 1      | Normal control| 4.62 ± 1.35                                  | 4.27 ± 0.25 | 4.07 ± 2.03 | 4.12 ± 1.38 |             |
| 2      | Diabetic control| 8.16 ± 2.18⁴                                 | 4.32 ± 1.91⁴ | 8.26 ± 0.54⁴ | 9.17 ± 1.34⁶ |             |
| 3      | Standard      | 6.19 ± 0.37                                  | 5.19 ± 1.36 | 6.17 ± 1.11∗∗∗ | 6.11 ± 2.12∗∗∗ |             |
| 4      | Treated 1     | 7.16 ± 1.15                                  | 5.85 ± 0.26∗∗∗ | 6.26 ± 2.46∗∗∗ | 5.93 ± 1.38∗∗ |             |
| 5      | Treated 2     | 6.31 ± 0.35                                  | 5.16 ± 1.82 | 6.31 ± 1.81∗∗ | 5.15 ± 1.27∗∗ |             |

Values are expressed as the mean ± SEM; n = 6. One-way ANOVA; followed by Tukey-Kramer’s multiple comparisons test: *P < 0.001 in comparison with normal control; **P < 0.001 in comparison with the diabetic control.
Table 9: Mechanical hyperalgesia: pinprick test in all groups in Wistar rats on weeks 2, 4, 8, and 12.

| S. No. | Treatment type | Response latency in sec (2nd week) | Mechanical hyperalgesia (pinprick test) sec | Response latency in sec (4th weeks) | Response latency in sec (8th weeks) | Response late response latency in sec (12th weeks) |
|--------|----------------|----------------------------------|-------------------------------------------|----------------------------------|----------------------------------|-----------------------------------------------|
| 1      | Normal control | 2.26 ± 1.36                      |                                            | 2.19 ± 1.48                      | 1.97 ± 1.39                      | 2.18 ± 1.92                                    |
| 2      | Diabetic control | 5.16 ± 1.36                      |                                            | 4.96 ± 1.04                      | 6.71 ± 0.48                      | 9.71 ± 1.04                                    |
| 3      | Standard       | 4.22 ± 0.23***                   |                                            | 3.12 ± 2.05***                   | 4.43 ± 2.82***                   | 4.74 ± 2.72***                                 |
| 4      | Treated 1      | 5.01 ± 0.11                      |                                            | 3.15 ± 0.45***                   | 4.21 ± 2.91***                   | 4.89 ± 1.25***                                 |
| 5      | Treated 2      | 4.14 ± 1.26***                   |                                            | 3.38 ± 1.26                      | 4.96 ± 2.91***                   | 4.51 ± 0.21***                                 |

Values are expressed as the mean ± SEM; n = 6. One-way ANOVA; Tukey-Kramer’s multiple comparisons test: *P < 0.001 in comparison with normal control; **P < 0.001 in comparison with the diabetic control.

Table 10: Grip strength to record the response latency in sec in all groups in Wistar rats on weeks 2, 4, 8, and 12.

| S. No. | Treatment type | Response latency in sec (2nd week) | Grip strength for each animal in a group ± SEM (n = 6) | Response latency in sec (4th weeks) | Response latency in sec (8th weeks) | Response latency in sec (12th weeks) |
|--------|----------------|----------------------------------|------------------------------------------------------|----------------------------------|----------------------------------|-----------------------------------------------|
| 1      | Normal control | 31.11 ± 0.13                     | 30.97 ± 1.41                                         | 31.21 ± 1.91                     | 34.31 ± 0.19                     |
| 2      | Diabetic control | 13.03 ± 0.38*                     | 8.93 ± 1.51*                                         | 6.49 ± 1.41*                     | 4.01 ± 0.25*                     |
| 3      | Standard       | 15.14 ± 1.26*                     | 24.31 ± 0.87*                                         | 25.56 ± 1.32*                    | 28.21 ± 2.13*                    |
| 4      | Treated 1      | 15.77 ± 1.06*                     | 21.11 ± 3.71*                                         | 23.27 ± 0.39*                    | 24.35 ± 1.16*                    |
| 5      | Treated 2      | 19.03 ± 1.36*                     | 25.25 ± 0.27*                                         | 26.18 ± 0.23*                    | 32.16 ± 1.62*                    |

Values are expressed as the mean ± SEM; n = 6. One-way ANOVA; followed by Tukey-Kramer’s multiple comparisons test: *P < 0.05 in comparison with normal control; **P < 0.05 in comparison with the diabetic control.

Table 11: Spontaneous locomotor (exploratory) test: actophotometer on all groups in Wistar rats on weeks 2, 4, 8, and 12.

| S. No. | Treatment type | Counts (2nd week) | Actophotometer: counts per 5 minutes | Counts (4th weeks) | Counts (8th weeks) | Counts (12th weeks) |
|--------|----------------|------------------|-------------------------------------|--------------------|--------------------|--------------------|
| 1      | Normal control | 121.36 ± 1.34    | 126.11 ± 2.52                       | 122.52 ± 3.65      | 127.43 ± 2.01      |
| 2      | Diabetic control | 35.14 ± 3.26*    | 28.18 ± 0.14*                       | 19.04 ± 1.81*      | 15.01 ± 3.24*      |
| 3      | Standard       | 41.52 ± 2.42*    | 70.27 ± 2.44*                       | 84.24 ± 2.22*       | 93.25 ± 0.42*      |
| 4      | Treated 1      | 42.37 ± 3.25*    | 68.26 ± 2.32*                       | 80.14 ± 3.62*       | 90.33 ± 2.66*      |
| 5      | Treated 2      | 44.46 ± 3.23*    | 72.24 ± 2.72*                       | 89.25 ± 2.43*       | 100.24 ± 2.55*     |

Values are expressed as the mean ± SEM; n = 6. One-way ANOVA; followed by Tukey-Kramer’s multiple comparisons test: *P < 0.05 in comparison with normal control; *P<0.05 in comparison with the diabetic control.

Table 12: Neuromuscular coordination test (motor coordination): rotarod test on all groups in Wistar rats on weeks 0, 2, 4, and 6.

| S. No. | Treatment type | Counts (2nd week) | Rotarod test (counts) | Counts (4th weeks) | Counts (8th weeks) | Counts (12th weeks) |
|--------|----------------|------------------|-----------------------|--------------------|--------------------|--------------------|
| 1      | Normal control | 123.14 ± 2.01    | 125.1 ± 0.25          | 127.41 ± 1.02      | 129.37 ± 4.14      |
| 2      | Diabetic control | 43.11 ± 1.10*    | 21.41 ± 3.14*         | 16.24 ± 0.61*      | 10.39 ± 2.05*      |
| 3      | Standard       | 50.26 ± 4.39     | 84.01 ± 5.23***       | 97.52 ± 2.19***    | 109.13 ± 0.28***   |
| 4      | Treated 1      | 49.40 ± 2.91***  | 78.27 ± 2.71***       | 89.24 ± 3.27***    | 100.25 ± 4.82***   |
| 5      | Treated 2      | 52.15 ± 4.16     | 96.13 ± 3.37***       | 99.24 ± 4.46***    | 102.18 ± 0.31***   |

Values are expressed as the mean ± SEM; n = 6. One-way ANOVA; Tukey-Kramer’s multiple comparisons test: *P < 0.001 in comparison with normal control; **P < 0.001 in comparison with the diabetic control.
which shows improvement in glucose levels compared with the diabetic control group on day 28 on the experiment. Group III has shown demonstrated improvement of glucose levels with $121.18 \pm 3.23$ mg/dl concentration on day 28.

3.4. Body Weight, Food, and Water Consumption. Body-weight of Group IV and Group V was $201.34 \pm 1.38$ g and $211.23 \pm 1.42$ g at conclusion of the experiment, which is higher when compared against diabetic control ($132.36 \pm 3.26$) and also efficient when compared with Group III ($199.2 \pm 1.12$ g), while food intake was $22.47$ and $21.02$ g/day which is less than the diabetic control group ($27.26 \pm 0.31$ g/day). The glibenclamide group (Group III) has shown efficient results with $21.14 \pm 2.62$ g/day food intake.

On recording the water intake, it was found that Group IV and Group V consumed $17.42 \pm 1.35$ and $16.13 \pm 0.59$ ml/day, which is less than the diabetes group, while Group III has shown $21.17 \pm 2.35$ ml/day of average water intake (Tables 3 and 4).

3.5. Lipid Profile. Groups IV and V have shown $85.02 \pm 1.91$ mg/dl and $74.59 \pm 0.24$ mg/dl levels of TC; $72.26 \pm 2.43$ mg/dl and $65.15 \pm 1.28$ mg/dl levels of triglyceride; $45.26 \pm 2.36$ mg/dl and $37.24 \pm 2.29$ mg/dl of LDL levels; $21.13 \pm 1.27$ mg/dl and $14.01 \pm 0.13$ mg/dl of VLDL levels; and $20.35 \pm$
Figure 8: Histopathological analysis of the liver tissues which were obtained post 12 weeks of experiment (experimentally induced diabetic neuropathy) from all animal groups. Images represent the control group, diabetic control, standard, and treated groups. Samples were stained and show results as follows: (a) hepatic cells were observed as normal, and narrow sinusoids were clearly visible. (b) Fatty necrosis and ballooning degradation were noted in this image which represents disruptive cells in the liver. (c) Narrow sinusoids were observed in the slides, and comparatively healthy tissue was noted. (d) Normal hepatic cells with narrow sinusoids. (e) Normal hepatic cells with narrow sinusoids were observed. CV: central vein; NS: narrow sinusoids; FN: fatty necrosis; BD: ballooning degradation.

Table 13: Effect of thymol on endogenous biomarkers in rats.

| S. No. | Treatment type | Group type | SOD (U/mg of protein) | NO (μg/ml) | LPO (nM/mg of protein) | Na⁺K⁺ATPase (μmol/mg of protein) | TNF-α (pg/ml) |
|--------|----------------|------------|-----------------------|------------|-----------------------|---------------------------------|--------------|
| 1      | Normal control |            | 24.03 ± 1.42          | 104.03 ± 1.36 | 2.34 ± 1.13          | 10.11 ± 1.23                   | 51.35 ± 1.13 |
| 2      | Diabetic control|            | 5.32 ± 1.13          | 303.11 ± 2.11   | 9.11 ± 1.13        | 2.34 ± 0.17                   | 160.01 ± 1.21 |
| 3      | Standard       |            | 16.12 ± 0.12         | 224.32 ± 2.12   | 6.13 ± 1.41        | 6.21 ± 0.21                   | 119.13 ± 1.12 |
| 4      | Treated 1      |            | 21.11 ± 1.23        | 158.09 ± 2.12   | 4.13 ± 1.23        | 7.31 ± 2.03                   | 90.21 ± 1.13  |
| 5      | Treated 2      |            | 23.24 ± 1.22        | 129.22 ± 2.01   | 2.71 ± 1.31        | 9.34 ± 1.34                   | 67.25 ± 0.21  |

Values are expressed as the mean ± SEM; n = 6. One-way ANOVA; Tukey-Kramer’s multiple comparisons test: *P < 0.05 compared to normal control; **P < 0.05 in comparison with the diabetic control.
2.26 mg/dl and 28.03 ± 0.29 mg/dl of HDL levels which are improved values compared with the diabetic group. The standard group has shown improvements in lipid profile with TC level as 77.23 ± 2.36 mg/dl, triglyceride as 65.48 ± 1.49 mg/dl, LDL levels as 37.16 ± 0.25 mg/dl, HDL levels as 29.8 ± 2.46 mg/dl, and VLDL levels as 15.12 ± 1.38 mg/dl which have shown improvements in comparison to the diabetic group [Table 5].

3.6. Behavioural Studies

3.6.1. Estimation of Thermal Hyperalgesia. In Eddy’s hot plate experiment, the response time recorded was 6.08 ± 1.41 s and 5.93 ± 0.24 s for Group IV and Group V, respectively, as recorded on the 12th week. The results justify the healing of the nerves compared with Group II, which has recorded a higher (9.85 ± 1.86 s) response time due to potential damage to the nerves due to diabetic neuropathy. Group III has shown 5.91 ± 1.74 s as response time, which improves results (Table 6).

3.6.2. Writhing Responses. Writhing responses have recorded values of 8.28 ± 1.44 and 8.24 ± 0.17 (number of reactions) on week 12, which have shown improved latency when compared against the diabetic control group (Group 2), where reported latencies on week 12 are high due to potential damage to the nerves. Group III responses have also improved (8.94 ± 1.35) [Table 7].

3.6.3. Cold Hyperalgesia. When the acetone drop test results were compared from Group IV and Group V against diabetic control on 12th week, the values obtained were 5.93 ±
Here, an improvement in the latencies of the treated groups was noted compared to the diabetic group. Group III results were recorded as $6.11 \pm 2.12$ which had improvement compared to the diabetic control group (Table 8).

### 3.6.4. Mechanical Hyperalgesia: Pinprick Test

On analysing the results of Group IV and V with Group II, readings of $4.89 \pm 1.25$ s and $4.51 \pm 0.21$ s have been noted on week 12 of experimentation. Though the changes have been quite significant throughout the experiment, the results on week 12 have shown improved results in the treated groups. The standard drug group has recorded a response of $4.74 \pm 2.72$ s which shows improvement (Table 9).

### 3.6.5. Grip Strength

The animals in the diabetic group have lost their grip strength due to prolonged diabetes; hence, the diabetic group on week 12 shows poor results. In comparison, significant values of $24.35 \pm 1.16$ and $32.16 \pm 1.62$ (sec) were recorded for Groups V and V, respectively. Group III has recorded $28.21 \pm 2.13$ (sec) as grip strength. All treatment groups show improvement in the mechanical strength of animals (Table 10).

### 3.6.6. Spontaneous Locomotor (Exploratory) Test

Significant readings of $93.25 \pm 0.42$, $90.33 \pm 2.66$, and $100.24 \pm 2.55$ (counts) were obtained on week 12 of experimentation in Groups III, IV, and V. The readings are very similar to normal groups and significantly different from the diabetic control group; hence, they show an improvement in the spontaneous locomotor activity due to healing effect of drug (Table 11).

### 3.6.7. Neuromuscular Coordination Testing

Motor coordination readings (rotarod test) for Groups III, IV, and V were recorded to be $109.13 \pm 0.28$, $100.25 \pm 4.82$, and $102.18 \pm 0.31$ which are significantly different compared to Group II on the 12th week of the experiment (Table 12).

### 3.7. The Histopathological Analysis

The histopathological analysis has shown normal results in treated groups, as shown in Figures 7 and 8. Little or no degradation was noted in sciatic nerve tissues similarly; no necrosis was noted in liver tissues of Groups IV and V. All results have shown significance against the diabetic control rats.

### 3.8. Biochemical Parameters

#### 3.8.1. Superoxide Dismutase Effect

The results have shown significant, dose-dependent results with isolated doses of thymol. A significant reduction in the value of superoxide dismutase was found ($5.32 \pm 1.13$ U/mg of protein) in the homogenate of the sciatic nerve obtained from diabetic rats (Group II) compared with the normal control group (Group I), i.e., $24.03 \pm 1.42$ U/mg of protein. Rats from treated groups have shown rise in SOD levels ($21.11 \pm 1.23$ and $23.24 \pm 1.22$ U/mg of protein) as shown in Table 13.

#### 3.8.2. The Effect of Thymol-Treated Groups on Nitrosative Stress

In the sciatic nerve, the neural nitrite levels from the diabetic groups were significantly increased to $108.77 \pm 0.82$, $100.25 \pm 2.66$, and $102.18 \pm 0.31$ which are significantly different compared to Group II on the 12th week of the experiment (Table 12).
2.11 μg/ml ($P < 0.05$) in comparison to the normal group recording values of 104.03 ± 1.36 ($P < 0.05$). Treated groups have shown values of 158.09 ± 2.12 and 129.22 ± 2.012 which are lowered significantly compared to the diabetic group rats (Table 13).

3.8.3. Lipid Peroxide Profile Evaluation. Neutral Lipid Peroxidase (LPO) levels were recorded in all groups at the end of 12 weeks of study. The diabetic group has shown a significant increase ($P < 0.05$) in LPO levels. The value recorded for Group II was 9.11 ± 1.13 nM/mg of protein compared with the levels from the normal group (2.34 ± 1.13 nM/mg of protein). The treated groups recorded values as 4.13 ± 1.23 and 2.71 ± 1.31 nM/mg of protein ($P < 0.05$) (Table 13).

3.8.4. Effect in Membrane-Bound Inorganic Phosphate. Na$^+$ K$^+$ ATPase level (membrane-bound inorganic phosphate) after 12 weeks of study was found to be 2.34 ± 0.17 μmol/mg of protein with $P < 0.05$ in rats from the diabetic group which is low in comparison to normal group results (10.11 ± 2.23). The treated group rats have shown values of 7.31 ± 2.03 and 9.34 ± 1.34 μmol/mg of protein which is significantly different from the diabetic group (Table 13).

3.8.5. Effect in the TNF-α. A significant increase in TNF-α from sciatic nerves was noted in STZ diabetic groups (160.01 ± 1.21 pg/ml) when compared to the control group (51.35 ± 1.13 pg/ml). Treated groups have shown improved results with values of 90.21 ± 1.13 and 67.25 ± 0.21 pg/ml compared to the diseased groups (Table 13). Improvement in motor coordination was noted in treated group SOD$_2$ Na$^+$ K$^+$ ATPase values of treated groups receiving 10 and 20 mg/kg of thymol have shown a significant increase in the levels by $P < 0.05$ compared with those of the diabetic group. NO, LPO, and TNF-α values have a significant decrease in the levels by $P < 0.05$ compared to the diabetic group.

4. Discussion

A large part of the world’s population suffers from diabetes whose major complications include hyperglycemia and diabetic neuropathy. Inflammation and hyperglycemia are the two events that alter gene expressions, thus affecting the cellular proteins, hence leading to progressive changes like diabetic complications and pathological changes. With the current growth status, diabetic neuropathy will be the prime cause of terminal stage renal diseases globally with intolerable consequences and costs for healthcare in developed countries [50, 51].

In the current treatment scenario, antidepressant drugs have been used in the treatment and regulation of neuropathic pain, which leads to the maintenance of sustained levels of neurotransmitters (norepinephrine and serotonin). As these hormones have shown a reduction in the central
nervous system’s pain pathways, they have shown good results in regulation and sensitivity to pain [52, 53]. Spectroscopic analysis has confirmed the presence of thymol (Figures 9–14). In the current study, the effects of thymol isolated from Trachyspermum ammi have been investigated for doses of 10 and 20 mg/kg on diabetes and diabetic neuropathy induced by streptozotocin in Wistar rats. The tests were also conducted for behaviour analysis, food, water consumption, lipid profile analysis, nociception, response latency, neuromuscular coordination, and histopathological analysis of the liver and sciatic nerve. A decrease in the blood glucose levels was noted by doses of 10 and 20 mg/kg of thymol. The reduction in blood glucose levels must be due to the stimulation of insulin secretion from beta cells that stimulate glucose metabolism and restore the remaining cells—this might be the best possible mechanism suggested by the standard drug glibenclamide [54].

This experiment has also shown protective effects of the isolated compound to treat diabetic neuropathy, which is a prominent effect of prolonged diabetes conditions. A loss of pain perception is noted in people who have diabetes due to extensive nerve damage; hence, there is the induction of peripheral neuropathy [55, 56]. The study revealed a delayed response in diabetic control group animals due to loss of perception and extensive damage to nerves, induced by prolonged diabetes. A gradual healing action was noted in treated groups which were most significant in groups receiving 20 mg/kg of thymol. In comparison, the diabetic groups have shown a trend of high sensitivity responses in nerves to delayed responses during week 12 due to extensive nerve damage. The treatment groups have also demonstrated a significant reduction in the paw withdrawal during thermal and mechanical hyperalgesia also reported a loss of in nociception in diabetic animals must be due to the development of DN, which is the characterization of nerve degradation, leading to loss in the perception of pain [55, 58]. When an animal starts developing nerve damage, it tends to become lethargic and shows delayed responses due to progression of pain and development of diabetic neuropathy [59, 60]. The results obtained from the tail-flick test, thermal hyperalgesia, and cold hyperalgesia also reported a loss of perception of pain, hot or cold due to intense nerve damage in diabetic rats, and improved results in treated groups due to their nerve damage healing action. The doses of 10 and 20 mg/kg have also shown improvement in the liver and sciatic nerve tissues, hence expressing the neuro-protection and antidiabetic action of isolated thymol.

Loss of nociception in diabetic animals must be due to the development of DN, which is the characterization of nerve degradation, leading to loss in the perception of pain [55, 58]. When an animal starts developing nerve damage, it tends to become lethargic and shows delayed responses due to progression of pain and development of diabetic neuropathy [59, 60]. The results obtained from the tail-flick test, thermal hyperalgesia, and cold hyperalgesia also reported a loss of perception of pain, hot or cold due to intense nerve damage in diabetic rats, and improved results in treated groups due to their nerve damage healing action. The doses of 10 and 20 mg/kg have also shown improvement in the liver and sciatic nerve tissues, hence expressing the neuro-protection and antidiabetic action of isolated thymol.

In the analysis of vasa nervorum (longitudinal) in the sciatic nerves of Wistar rats on 12-week prominent signs of...
inflammation and damage, lipid degenerated axons showing focal peripheral axonal loss were noted in the diabetic group while the treated groups receiving 10 and 20 mg/kg thymol have shown minimal axonal degeneration which is a sign of healing and neuroprotective action of the drug. These healing signs are the recoveries of the damage induced by prolonged diabetes conditions. Similarly, the cells in the diabetic group have shown fatty necrosis and ballooning degradation, which represents disruptive cells in the liver. At the same time, the treated groups (10 and 20 mg/kg thymol) have shown normal hepatic cells
with narrow sinusoids which are healthy liver cells. They signify the recovery of cells from diabetes. Hence, the sciatic nerve and liver cell histopathology hints on antidiabetic and neuroprotective thymol on Wistar rats.

Positive findings were recorded in thermal hyperalgesia, writhing, cold hyperalgesia responses, mechanical hyperalgesia, grip strength, spontaneous locomotor (exploratory) test, neuromuscular coordination tests, and histopathological and lipid profile analysis. The neuroprotective results were quite significant at the end of treatment that is by the end of 12 weeks. Uncontrollable lipid profiles have been the prime complications of diabetes mellitus, which is noted in 40% of the diabetes cases. Hypertriglyceridemia is a condition very commonly found in patients suffering from diabetes mellitus and is also seen in situations of insulin resistance either due to reduced catabolism or reduced production of lipoproteins [61].

Endothelial damages are caused due to the presence of superoxide and endogenous enzymes. An increase in aldose reductase and protein kinase C has shown connections with pain perceptions. Antioxidant protection is done by SOD which transforms the superoxide anions into H2O2. Endogenous enzymes SOD and MDA closely reciprocate to the oxidative stress. Elevation in the stress levels as depicted by MDA leads to lipid membrane disruption. This causes rearrangement of bonds in unsaturated fatty acids created by tissue damage. Increased levels of lipid peroxidase are linked to oxidative stress levels, which are decreased during the drug treatment process. Nitric oxide is an intracellular messenger, and it has a significant role in the pathological process as it combines with ROS (reactive oxygen species), an antioxidant nature and healing action of thymol isolated from the herbal extract were signified and confirmed.

5. Conclusion
In conclusion, the treatment of thymol concentrations of 10 mg/kg and 20 mg/kg has ameliorated diabetes and conditions of diabetic neuropathy in STZ rats. Hence, it is obvious that thymol’s oral dosing isolated from Trachyspermum ammi has improved the nociceptive latency, glucose levels, and lipid profiles and has subsequently shown healing effects in the liver and sciatic nerve of the treated groups where 20 mg/kg of thymol is more potent than its lower concentration group. The future aspects of this work suggest a detailed study on the mechanism of action to be performed to get better insights into thymol’s role in the treatment of neuropathy, diabetes, and other metabolic disorders.

Abbreviations

| Abbreviation | Description |
|--------------|-------------|
| DPN          | Diabetic polyneuropathy |
| DN           | Diabetic neuropathy |
| CPCSEA       | Committee for the Purpose of Control and Supervision of Experiments on Animals |
| DMSO         | Dimethyl sulfoxide |
| OECD         | Organisation for Economic Co-operation and Development |
| DNA          | Deoxyribonucleic acid |
| DM           | Diabetes mellitus |
| TMB          | 3,3′,5,5′-Tetramethylbenzidine |
| UV spectroscopy | Ultraviolet spectroscopy |
| TLC          | Thin Layer Chromatography |
| IAEC         | Institute Animal Ethics Committee |
| FTIR         | Fourier-transform infrared spectroscopy |
| HPLC         | High-Performance Liquid Chromatography |
| HRMS         | High-resolution mass spectrometry |
| NMR          | Nuclear Magnetic Resonance |
| STZ          | Streptozotocin |
| SOD          | Superoxide dismutase |
| MDA          | Malondialdehyde |
| NO           | Nitric oxide |
| LPO          | Neutral Lipid Peroxidase |
Na⁺K⁺ATPase: Membrane-bound inorganic phosphate

TNF-α: Tumor necrosis factor alpha

Pyridyl porphyrin: Fe(III)tetrakis-2-(N-triethylene glycol monomethyl ether.

Data Availability

Data will be available on request to the corresponding author.

Conflicts of Interest

The authors declare that they have no conflicts of interest.

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