

## Original Research Article

### **Integrating *in Silico* and *in Vitro* approaches to screen the Antidiabetic Properties from *Trachyspermumammi* leaves**

#### **Abstract**

**AIM:** The purpose of this study was to explore the possible mechanisms through which *Trachyspermumammi* may be beneficial in managing diabetes by *in vitro* methods and to predict the potential bioactive constituent/s responsible for its anti-diabetic activity through *in silico* docking study.

**STUDY DESIGN:** This study engaged *in vitro* and *in silico* studies to investigate the enzyme inhibition assay and molecular docking studies of phytochemical constituents when compare with standard treatment.

**PLACE AND DURATION OF STUDY:** Gokaraju Rangaraju college of pharmacy, Bachupally, Hyderabad, India. December 2021-2022.

**METHODOLOGY:** Methanolic leaf extract of *Trachyspermumammi* was prepared using maceration extraction. Plant extract was evaluated for its *in vitro* anti-diabetic activity. Further, docking screening was performed using Mcule software to predict potential moiety which may be responsible for its anti-diabetic activity.

**RESULTS:** Maceration extraction resulted in extractive yield of 16.15%. The presence of phenolic and flavonoid components in extract ( $135.59 \pm 8.87$  GAE/gm and  $62.1 \pm 4.17$  quercetin/gm) dried extract respectively. *In vitro* anti-diabetic assays revealed that anti-hyperglycaemic activity of this plant can be attributed to its high efficiency to inhibit  $\alpha$ -amylase (69.7%) and glucosidase (62.23%) enzymatic activity and protein glycation assay (95.23%), which are well established targets for the management of diabetes. Further, through docking studies we predicted that Apigenin, luteolin, Germacrene D and  $\gamma$ -Tocopherol present in this plant might be responsible for the anti-diabetic properties exhibited by this plant. These results provide a scientific justification for the traditional anti-diabetic use of this plant

**CONCLUSION:** It may control diabetes through lowering dietary glucose uptake. Predicted anti-diabetic molecules need to be screened further for the management of hyperglycemia. It may conclude that *Trachyspermumammi* is an important vegetable with a potent source of natural

antioxidants and antidiabetic activity justifying its traditional use in green therapeutics.

**Keywords:** *Trachyspermum ammi*, antidiabetic activity, phenolics, flavonoids, Mucule, molecular docking.

## 1. INTRODUCTION

Diabetes is a chronic metabolic condition. disorder typified by excess amount of blood sugar, also known as glucose, which causes catastrophic collateral harm to the kidneys, eyes, blood arteries, heart, and nerves over time. Type 2 diabetes, which frequently affects adults and develops when the body becomes insulin resistant or is unable to produce enough insulin, is the most prevalent. The rate of type 2 diabetes has increased substantially across nations of any socioeconomic status over the last thirty years [1].

In the present research, we looked in to the anti-diabetic properties of crude extracts from *Trachyspermum ammi* leaves in vitro. The primary goal of this experimental investigation is to determine the anti-diabetic effectiveness of a Methanol leaf extract using *invitro*- amylase inhibition assays, assay regarding  $\alpha$ - glucosidase inhibition, and assay regarding protein glycation inhibition. The natural compounds work by preventing glucose absorption by blocking hydrolyzing enzymes of carbs such as pancreatic amylase. Inhibiting this enzyme's activity slows carbs digestion and lengthens overall digesting time for carbohydrate, resulting in a decrease in glucose absorption rate and, as a result, a dulling of the postprandial plasma glucose spike. Several indigenous medicinal plants have a strong potential for suppressing the activity of the  $\alpha$ -amylase enzyme [2]. Alpha-glucosidases are digestive enzymes that metabolise carbs into glucose. Inhibiting the action of alpha-glucosidases with synthetic medicines is one technique for treating type 2 diabetes [3]. Chronically elevated glucose levels causes protein glycation and the slow generation of advanced glycation end products (AGEs) in numerous human tissues, which contributes to the onset of long-term diabetes problems. The production of AGEs occurs under normal biological settings, but it grows more rapid when blood sugar levels are raised. Depositions of AGEs are associated with a variety of illnesses. Diabetes-related problems such as diabetic retinopathy, nephropathy, cataract, and atherosclerosis are facilitated by chronic hyperglycemia resulting in AGEs [4].

Acarbose is a complicated oligosaccharide that inhibits alpha-amylase present in pancreas and membrane-bound intestinal alpha-glucosidase hydrolase in an effective, reversible manner. Acarbose reduces glucose absorption by delaying carbohydrate digestion, resulting in lower postprandial blood glucose concentrations [5]. In the Maillard's process, aminoguanidine acts as a scavenger of reactive carbonyl intermediates. Production of AGEs is suppressed by Aminoguanidine which additionally decreases retinal, diabetic renal, vascular and neurological problems [6].

*Trachyspermum ammi* L., a member of the *Apiaceae* family, is a highly prized seed spice with significant therapeutic significance. Thymol, the primary ingredient in this oil, has been utilised to treat bronchial issues, loss of appetite and gastrointestinal conditions [7]. The primary goal of the current investigation is to introduce different in vitro methodologies for doing diabetes research.

## **2. METHODOLOGY**

### **2.1 Collection and drying of plants**

In January, a collection of *Trachyspermum ammi* Linn leaves was made in Hyderabad, Telangana, and they were determined to be, verified and confirmed by a qualified botanist. The leaves are roughly blended in a mixer grinder after being shade-dried for about 1 week. The blended material was either saved or gathered for extraction.

### **2.2 Plant extract preparation**

Maceration extraction principle This is an extraction technique where solvent for extraction is poured on top of coarsely powdered plant material, such as leaves, stem bark, or root bark, until the drug material is thoroughly covered. The solvent is then allowed to evaporate for at least 3 to 7 days after which the container is tightly closed [8]. The leaves of *Trachyspermum ammi* are extracted using methanol as a solvent.

### **2.3 Phytochemical analysis**

To determine the secondary metabolites present in *Trachyspermum ammi* leaves methanolic extract of the leaves were subject to preliminary phytochemical screening.

### **2.4 Quantitative analysis**

#### **2.4.1 Total Phenolic Contents (TPC)**

The Folin Ciocalteu (FC) reagent is used to calculate TPC. Add 5 µl of FC reagent to 5 µl of plant extract, and incubated at 22 °C. 50 µl of 20% sodium carbonate should be added. After 1hr 30 min 22 °C, the absorbance at 765 nm should be adjusted. The dry weight of gallic acid is used as a reference to calculate the total phenolic content (mg/g). TPC is determined using a regression equation in Excel with the formula  $y=mx+c$  and various META concentrations.

#### **2.4.2 Total Flavonoid Contents (TFC)**

Aluminium chloride ( $AlCl_3$ ) method is used for determining TFC. Mix two hundred microliters of Methanolic Extract *Trachyspermum ammi* with 5 µl of distilled water, seventy five microliters of 5% sodium nitrite, and 150 micro liters of 10%  $AlCl_3$  after incubating at 25 °C for 5 minutes. The remaining 500 µl of the 1 mol/L NaOH solution were added to the reaction mixture. The absorbance at 510 nm was measured after 15 minutes of standing. Quercetin is used as a benchmark to assess the total flavonoid content (mg/g) of dry weight. TPC is determined using an Excel regression equation with the formula  $y=mx+c$  with varying META concentrations [9].

## **2.5 Acute Toxicity Studies**

The OECD 425 recommendations were followed for conducting the acute toxicity studies. The current study was completed in the animal facility approved by CPCSEA Gokaraju Rangaraju College of Pharmacy's in Bachupally, Hyderabad, India (Registration Number 1175/PO/ERe/S/08/CPCSEA).

### **2.5.1 Experimental animals**

We bought 200–250 g Wistar albino rats from Jeeva Life Sciences in Hyderabad. care and maintenance for animals were taken according to the guidelines approved by CPCSEA.

### **2.6 *In-vitro* methods for Antidiabeticactivity**

**2.6.1  $\alpha$ -Amylase inhibition assay:** The colon's  $\alpha$ -Amylase enzyme is inhibited, delaying the transformation of starch and sugars into monosaccharides prior to ingestion. As a result, it lowers blood sugar levels after a meal and glucose absorption. The sample concentrations (10–50  $\mu\text{g/ml}$ ), 10 mL of  $\alpha$ -amylase which is (0.5 microgram/milliliter) concentration, twenty mL of starch solution, are combined with hundred mL of 0.2 M phosphate buffer (pH – 6.9). After standing at 37°C for 5 minutes, the reaction was stopped by adding 2 ml of the 3, 5-dinitrosalicylic acid reagent. Using 10 mL of distilled water, a tray of ice is heated for fifteen minutes at 100 °C before the volume is changed. The activity of  $\alpha$ -amylase, colour intensity at 540 nm is estimated using a spectrophotometer. The equation was used to calculate the % inhibition of enzyme activity. Three separate runs of each test were completed. The standard used was acarbose as reference [10].

### **2.6.2 $\alpha$ -Glucosidase inhibition assay:**

Enzymes glucosidases are that increase conversion into simple sugars from starch. This facilitates digestion of carbohydrates from food, starch as well which increases blood glucose levels in people and causes sugar to be produced for intestinal absorption. Each sample concentration (10–50 mg/ml) receives a 10-min fifty microliters of  $\alpha$ -glucosidase (Two U/ml) pre-incubated, and then fifty microliters of PB (pH 7.4) were incubated at thirty seven °C. The reaction was then provided with 50  $\mu\text{L}$  of p NPG (20 mM) and at 37 degrees Celsius it was incubated for thirty min. Acarbose was used as a reference and the procedure was repeated three times. Utilising a spectrophotometer, the absorbance is constant at 540 nm [10].

### **2.6.3 Protein glycation inhibition assay**

Reducing sugars and free amino groups are used in the non-enzymatic process of glycation, which results in reversible precursors that go through modifications to form AGEs. It was also acknowledged that the production of AGEs is influenced by a variety of metabolites, such as fructose, trioses, and dicarbonyl compounds, which participate much in the glycation reaction more quickly compared to

glucose. Since glucose is most prevalent blood sugar and is present in high concentrations in diabetes, research on glucose and protein glycation has been extensive.

#### **2.6.4 Procedure**

Phosphate buffer is prepared in such way that 0.1 M  $\text{NaH}_2\text{PO}_4$  with pH 6 is combined with 0.2 M disodium hydrogen phosphate with the pH of 9, the overall pH is adjusted to 6.9 by addition of NaCl solution. Ten milliliter of PB (200 millimolar concentration) with 7.4 as pH, five milliliter of BSA twenty milligram/milliliter in PB, 1 milliliter of plant extract 15.6-500 gram/milliliter, and 4 milliliter of 1000 millimolar fructose with PB are kept at room temperature for a week. Blank is made only with the extract of plant at the different concentrations, in contrast to blank control, that is made totally with BSA. Buffer is added to test tubes along with aminoguanidine at concentrations of 1.25, 0.75, and 0.25 mg/ml. A fluorescence spectrometer is used to detect the fluorescence emission of each pair at wavelengths of 355 nm and 440 nm, respectively [11].

#### **2.7 *In silico*/Molecular Docking**

From ligand design to drug development, molecular docking has been utilised to address a wide range of problems. By identifying the basic ingredient of the scoring functions for protein-protein docking between the two molecules, it models the interaction of protein-ligand complexes. The underlying principle of docking is that molecules with high affinity for protein-binding sites typically have molecular structures comparable to those of the binding sites. The form and spatial arrangement of the hydrophobic and hydrogen-bonding residues that surround the binding site are frequently used to describe the morphology of the binding site [12]. Proteins with PDB IDs 2QV4, 4W93 (for the amylase enzyme), 3L4J (for the glucosidase enzyme), and 5HY8 (for protein glycation) are docked against 12 botanical compounds acquired from GC-MS investigations for antidiabetic efficacy.

##### **2.7.1 Docking simulations**

**Preparation of ligands:** ligand is constructed and the chemical structures of molecules were both illustrated. Molecule's 2D ligands depicted docking on the ligand imported side.

**Preparation of protein:** The proteins with the PDB IDs 2QV4, 4W93 (for the amylase enzyme), 3L4J (for the glucosidase enzyme), and 5HY8 (for protein glycation) were initially retrieved in PDB format from the RCSB protein bank website. The SBD site sphere's attributes are prepared and recorded using the discovery studio visualizer.

**2.7.2 Protein-ligand interactions:** Through docking simulations, the binding position of drug candidates to their protein targets is predicted. Molecule was used to develop docking simulation experiments.

**2.7.3 Ligand docking and scoring:** 3L4J (X= 43.922180, Y= 100.994820, Z= 32.862640), 5HY8 (X= -17.511549, Y= 15.878669, Z= 66.503238), and 2QV4 (X= 13.758000, Y= 58.583058, Z= 22.465406) proteins were uploaded with sphere attributes, and the ligand structures had been docked against the corresponding proteins. According to docking, few of the compounds have strong affinity for the proteins

3L4J, 2QV4, 4W93, and 5HY8. With the use of mCULE Docking, flexible glide-ligand docking was able to enhance protein ligand interactions. The docked compounds show a docking score. The proteins 2QV4, 4W93, 3L4J, and 5HY8 interact with chemicals found in *Trachyspermum ammi* in the ways shown below.

**2.7.4 Visualization and analysis:** The final docking places were pictured using the discovery studio visualizer. The ligand interactions were visualized in order to comprehend the binding interactions between ligands and proteins. The best docked forms were chosen using the glide score technique. The binding will be increasingly beneficial as the value decreases. Additionally, the docked ligand locations and various ligand receptor interactions were examined.

### **2.8 Statistical analysis:**

Values are presented as the Mean SEM, with n = 6. ANOVA was used for the statistical analysis, which was followed by Dunnett's test. Comparisons were made between each group and the standard groups, control and disease-control, significance is denoted by the following symbols: standard (A=p<0.0001), control group (\*=p<0.001), a=p<0.0001 and b = p<0.001), and ns- nonsignificant.

## **3. RESULTS AND DISCUSSION**

Antioxidant and antidepressant properties for Methanolic extract of *Trachyspermum ammi* leaves were investigated. The complete set of research's findings is shown below.

### **3.1 Preparation of methanolic extract of *Trachyspermum ammi* leaves**

*Trachyspermum ammi* leaves were macerated to create a methanolic extract. The equation below is used to calculate the extract's percent yield.

$$\begin{aligned} \text{\% yield of extract} &= \text{Amount of extract obtained} / \text{Amount of powder used} \times 100 \\ &= 32.3/200 \times 100 \\ &= 16.15 \text{ \% w/w.} \end{aligned}$$

Quantity obtained was 16.15 % w/w.

### **3.2 Acute toxicity studies**

2000 milligram/kilogram body weight., per oral dose methanolic extract of *Trachyspermum ammi* leaf was administered to female albino mice. At 2000 milligram/kilogram body weight., the plant extract exhibited no symptoms of mortality or morbidity. All of the animals were safe even after fourteen days of observation. The biological analysis was performed at doses of 2 hundred and 4 hundred mg/kg body weight, orally.

### **3.3 Quantitative analysis**

#### **3.3.1 Total phenolic content (TPC)**

The calibration curve's linear regression equation and coefficient of linear regression are, respectively,  $y = 0.00090x - 0.04190$  and  $R^2 = 0.98710$ . Where x = absorbance, y = concentration for gallic acid solution in (g/ml). META has a TPC of  $135.59 \pm 8.87$  mg GAE/gm given in table 1.

**Table 1:** calibration data for Gallic acid

S.no	Concentration (µg/mL)	Absorbance
1.	0.100	0.0650
2.	0.200	0.1240
3.	0.300	0.1940
4.	0.400	0.2970
5.	0.500	0.3720
6.	0.600	0.4950

### 3.3.2 Total Flavonoid content (TFC)

The calibration curve's equation for linear regression and linear regression coefficient were, respectively,  $y = 0.0015x + 0.0517$  and  $R^2 = 0.9442$ , where  $x$ =absorbance,  $y$ =concentration of the quercetin solution (µg/mL). Methanolic extract of *Trachyspermum ammi* had a TFC of  $62.1 \pm 4.17$  mg quercetin/gm represented in table 2.

**Table 2:** calibration data of Quercetin

S.no	Concentration (µg/mL)	Absorbance
1.	0.100	0.1250
2.	0.200	0.3570
3.	0.300	0.6130
4.	0.400	0.7240
5.	0.500	0.7950
6.	0.600	0.9120

## 3.4 ANTIDIABETIC ACTIVITY

### *In-vitro* models

**3.4.1 α-Amylase inhibition assay:** To calculate the  $IC_{50}$  values of the test extract and acarbose, the α-amylase inhibition assay is used. Increasing concentration lead to a rise in the percent inhibition given in table 3. Test extract and acarbose have respective  $IC_{50}$  values of 33.950 microgram/milliliter and 24.830microgram/millilitre.

**Table 3:** *In-vitro* α-amylase inhibition assay of *Trachyspermum ammi*

S. no	Compounds	Concentrations (µg/mL)	% Inhibition	$IC_{50}$ (µg/mL)
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		10.0	17.84 ± 0.510	
		20.0	30.19 ± 0.400	
<b>1.</b>	<b>Test Extract</b>	30.0	44.18 ± 0.670	33.95
		40.0	55.28 ± 0.620	
		50.0	69.7 ± 0.880	
		10.0	27.33 ± 0.950	
		20.0	40.26 ± 0.310	
<b>2.</b>	<b>Acarbose</b>	30.0	65.46 ± 0.490	24.83
		40.0	71.43 ± 0.660	
		50.0	80.92 ± 0.820	

Experiment was performed for 3 times, and Mean±SEM was used to represent the percentage of inhibition. α-Amylase is a crucial digestion enzyme that is mostly found in saliva and pancreatic juice [13]. Inhibit and target this enzyme to lower elevated blood sugar levels during the night. Since the quantity of the extract affects the amount of enzyme inhibition, the amount of enzyme blocked appears to be dose-dependent. Amylase and glucosidase activities are strongly inhibited by β-pinene [14]. The pancreatic -amylase enzyme of humans is inhibited by fucosterol [15].

#### **3.4.2 α-Glucosidase inhibition assay:**

The 50% inhibition concentration values of the acarbose and test extract are determined using the α-glucosidase inhibition assay. Test extract and acarbose have IC<sub>50</sub> values of 32.040 microgram/milliliter and 25.620 microgram/milliliter, respectively given in table 4.

**Table 4:** *In-vitro* antidiabetic activity of META using α-glucosidase inhibition assay

S. no	Samples	Concentration (µg/mL)	% Inhibition	IC <sub>50</sub> (µg/mL)
		10.0	15.630±0.670	
<b>1.</b>	<b>Test Extract</b>	20.0	31.980±0.650	
		30.0	46.810±0.750	32.04
		40.0	57.400±0.530	
		50.0	62.230±0.420	
		10.0	17.060±0.360	
		20.0	39.020±0.660	
<b>2.</b>	<b>Acarbose</b>	30.0	58.070±0.420	25.62
		40.0	67.640±0.180	
		50.0	72.340±0.130	

Experiment was performed for 3 times, and Mean±SEM was used to represent the percentage of

inhibition.

The  $\alpha$ -Glucosidase enzyme, which is found on the mucosal brush border of the small intestine, also functions as a crucial digestive enzyme. Its function is to convert and disintegrate complicated carbohydrates into tiny, intuitive, and easily absorbed ones. Its inhibition is a useful method for postponing glucose uptake and lowering elevated postprandial blood glucose levels, and this may slow the course of diabetes [16]. The greatest quantities of thymol exhibit the highest inhibitory action for  $\alpha$ -glucosidase rather than  $\alpha$ -amylase [17].

### **3.4.3 Protein glycation inhibition assay:**

The rate of inhibition was determined to be  $92.85\pm 0.87$  and  $85.71\pm 0.87$ , respectively, in the protein glycation inhibitory assay of the extract and aminoguanidine at 250 g/mL. We found that concentration affects the % inhibition of the extract and standard given in table 5.

**Table 5:** *In-vitro* antidiabetic activity of *Trachyspermummammi* using protein glycation inhibition assay

S. no	Samples	Concentration ( $\mu\text{g/mL}$ )	% Inhibition
1.	<b>Test Extract</b>	0.15	21.420 $\pm$ 0.670
		0.30	41.260 $\pm$ 0.840
		0.60	53.380 $\pm$ 0.540
		0.120	75.390 $\pm$ 0.570
		0.250	92.850 $\pm$ 0.870
		0.500	95.230 $\pm$ 0.850
2.	<b>Aminoguanidine</b>	0.250	85.710 $\pm$ 0.870
		0.750	099 $\pm$ 0.630
		0.1250	1090 $\pm$ 0.740

Experiment was performed for 3 times, and MEAN $\pm$ SEM was used to represent the percentage of inhibition.

### **3.5 Molecular docking**

**Table 6:** Molecular docking studies of isolated compounds from *Trachyspermummammi* against respective proteins

**2QV4**



To predict the affinity and behavior of small molecules and drug candidates, computational molecular docking is commonly used to predict the binding orientation to their protein targets [20]. We used molecular docking to model various bioactive compounds isolated from the Jasmine plant that is known to inhibit glucosidase. Hydrogen bonds play a crucial role in the structure and interaction of protein-protein or ligand-receptor complexes. Hydrogen bonds are important in drug design to ensure that the drug is unique to the protein target. The presence of such associated hydrogen and hydrophobic bonds indicates a positive response to the inhibition of the  $\alpha$ -amylase enzyme. As a result, phytochemicals found in various plant extracts may be responsible for these extracts' antidiabetic properties.

#### 4. CONCLUSION

The results of this investigation suggest that *Trachyspermumammi* leaf extract has strong anti-diabetic effect. A novel study method is necessary to elucidate *Trachysperumammi*'s mode of action.

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