Trachyspermum ammi (L.) Sprague ex Turrill Seed Extracts Lack Antibacterial Activity

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ABSTRACT

Introduction: *Trachyspermum ammi* leaves and seeds have been used as dietary supplements and to treat multiple pathogenic diseases in several traditional healing systems. Despite this, their therapeutic properties have been poorly studied. **Methods:** Chloroform, ethyl acetate, hexane, methanol and water extracts of *T. ammi* were prepared and tested and the activity was compared to that of standard antibiotics for their *in vitro* antibacterial activity against ten human bacterial pathogens. The antibacterial activity was studied by standard disc diffusion assays and the activity was recorded as zones of inhibition. **Results:** All *T. ammi* seed extracts were ineffective at inhibiting the growth of all of the gram-positive and gram-negative bacteria pathogens screened against. **Conclusion:** Despite their use in traditional healing systems to treat some pathogenic diseases, *T. ammi* seed extracts were completely ineffective bacterial growth inhibitors. However, these

extracts were screened against a limited panel of bacteria and further testing against other pathogens is required.

Key words: Ajwain, Bishops weed, Apiaceae, Antibacterial activity, Traditional medicine, Medicinal plants.

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INTRODUCTION

Traditional plant derived medicines are used by multiple ethnic groups globally for a variety of therapeutic purposes, including for the treatment of diseases caused by microbial pathogens. Interestingly, the potential of plant extracts to inhibit the growth of pathogenic bacteria has gained considerable momentum in recent studies.¹⁻⁵ Most studies have focused on African,⁶⁻⁸ Asian,⁹⁻¹¹ South American¹² and Middle Eastern plants.^{5,13,14} Despite this, the capability of traditional plants to offer useful pharmaceutical benefits remains poorly explored. Approximately 5-10% of the 300,000-500,000 plant species across the world have been studied for therapeutic bioactivities.⁹ With plenty of plant species yet to be screened, it is paramount that plant selection processes narrow down the field to highlight the species with higher likelihood of providing viable new drug leads. One of the main selection criteria is to select plants which have documented uses as traditional medicines based upon their ethnobotanical usage. Another important selection criteria is based upon examining plants for which medicinal capacity is well established, or on species taxonomically related to species used traditionally. The investigation of traditional medicinal plants may lead to the discoveries of natural therapies.

The recent evolution of bacterial pathogens to strains that are either totally or extremely resistant to conventional antibiotics¹⁵ has rendered many clinical antibiotics of substantially lower efficacy, or in some cases has rendered them completely ineffective. There is now an urgent need to look for new and more effective antibiotic therapies. There are currently limited therapeutic options available for many diseases that are caused by resistant bacterial pathogens and the situation is expected to worsen in the future as the bacteria exchange resistance genes, rendering further pathogens resistant to clinical antibiotics. According to World Health Organization (WHO) the development of new antibiotic therapies is one of the most serious challenges currently faced by the medical community. Traditional medicines have a great capacity for the development of new antimicrobial drugs, explaining the considerable recent increase in interest in this discipline.^{16,17}

Trachyspermum ammi (L.) Sprague ex Turrill (commonly known as ajwain, bishops weed or carom) (Figure 1) is an annual herb that is native to Egypt and is widely cultivated in Afghanistan, Iran, Iraq, India and Pakistan. *T. ammi* belongs to family Apiaceae and is a highly valued medicinal seed. Analysis of ajwain seeds revealed that it contains fiber (11.9%), tannins, glycosides, carbohydrates (38.6%), moisture (8.9%), fat (18.1%), protein (15.4%), flavone, saponins and mineral matter (7.1%) containing phosphorus, calcium, iron and nicotinic acid.¹⁸ The seed contains around 2-4.4% of brown colored oil called ajwain oil and the primary component of this oil is thymol, which is widely used in the treatment of gastrointestinal diseases.¹⁹ The non-thymolic fraction contains γ -terpinene, dipentene, α - and β -pinenes, carvacrol and α -terpinene.²⁰ It is widely grown in areas where soil contains more salts.^{21,22} *T. ammi* is a 60-90 cm tall branched annual herb. Due to the therapeutic and commercial importance of *T. ammi*, this study was designed to screen *T. ammi* seed extracts for the ability to restrict the growth of a panel of gastrointestinal and autoimmune bacterial pathogens.

MATERIALS AND METHODS

Plant material and extraction

T. ammi seeds were purchased from Noodles Emporium, Australia (an online herbalist) and were originally sourced from Egypt. A voucher sample (GU-TANE19a) is stored in the School of Environment and Science, Griffith University, Australia. Individual 1 g masses of the seed were extracted individually in five different tubes each containing 50mL of methanol, deionized water, ethyl acetate, chloroform or hexane overnight at room temperature. The extract was then filtered through the Whatman number 1 filter paper and subsequently dried at 60°C under vacuum. The resultant pellet was weighed to determine extraction yield and dissolved in 100 μ L of dimethyl sulfoxide (DMSO). The volume was then increased to 10mL with the addition of deionized water. The extract was filtered through 0.22 μ m filter (Millipore) to remove microbial contaminants and stored at the room temperature until use.

Figure 1: *Trachyspirum ammi* (a) whole plant, (b) leaves, (c), flowers and (d) seeds.

Antibacterial screening Test microorganisms

Clinical isolate microbial strains of gram-negative bacteria (*Escherichia coli*, *Shigella sonnei*, *Aeromonas hydrophilia*, *Salmonella newport*, *Proteus mirabilis*, *Klebsiella pneumonia*, *Acinetobacter baylyi*, *Pseudomonas aeruginosa* and gram-positive bacteria (*Bacillus cereus*, *Enterococcus faecalis*, *Streptococcus pyogene*) were obtained from Ms. Michelle Mendell and Ms. Jane Gifkins, Griffith University. All the bacterial stock culture was sub-cultured and grown to log phase at 37°C. The cultures were maintained in nutrient broth (Oxoid, Australia) at 4°C until use.

Evaluation of antimicrobial activity

Antimicrobial activity of the *T. ammi* seed extracts was determined using a disc diffusion assay.²³ Briefly, 100µL of each of the microbial suspension in the log phase was spread onto individual nutrient agar plates. The extracts were screened for antimicrobial activity using 6mm sterilized filter paper discs. Each disc was infused with 10µL of the individual seed extracts and allowed to dry before incubated at 37°C for 24h. The diameter of the zone of inhibition (ZOIs) were measured to the closest possible whole millimeters. Each assay was performed in triplicate ($n=3$). Mean values ($\pm$ SEM) are reported in the study. Standard discs of tetracycline (10µg) and ampicillin (2µg) were used as a positive control to compare antibacterial activity. Filter disc infused with the 10µL of deionized water were included on each plate as negative controls.

Statistical analysis

Data is expressed as the mean $\pm$ SEM of two independent experiments, each with internal triplicates ($n=6$). One-way ANOVA is used to calculate the statistical between the control and treated groups, with a P value < 0.01 is consider to be statistically significant.

RESULTS

Liquid extraction yields

The extracted yield of 1g of dried seeds in methanol, water, ethyl acetate, chloroform and hexane is shown in the Table 1. The extracts were re-suspended in 10mL of deionized water (having 1 % DMSO), resulting in the extract concentration shown in Table 1.

Table 1: The mass of dried extracted material and the concentration after resuspension in deionized water.

	Mass of extracted material (mg)	Concentration of re-suspended extract (mg/ml)
Methanol	31.5	3.15
Water	144	14.4
Ethyl acetate	1.5	0.15
Chloroform	9	0.9
Hexane	12.55	1.25

Figure 2: Growth inhibitory activity of *T. ammi* seed extracts and reference antibiotics against gram-negative bacterial species (a = *E. coli*, b = *S. sonnei*, c = *A. hydrophilia*, d = *S. newport*, e = *P. mirabilis*, f = *K. pneumonia*, g = *A. baylyi*, h = *P. aeruginosa*) measured as ZOIs (mm) $\pm$ SEM. M= methanol, W= water, E= ethyl acetate, C= chloroform, H= hexane, A = ampicillin (2µg); T = tetracycline (10µg); NC = negative control. All assays were completed twice, each with internal triplicates ($n=6$) and the results are expressed as mean zones of inhibition (mm) $\pm$ SEM.

Antimicrobial activity

In order to determine the growth inhibitory activity of the *T. ammi* seeds extract, 10µL aliquots of each extract were screened in the disc diffusion assay. The *T. ammi* seeds extract were ineffective at inhibiting the growth of all gram-negative (Figure 2) and gram-positive (Figure 3) tested bacterial species. On the other hand, both the positive control antibiotics (ampicillin and tetracycline) were effective growth inhibitors except the *K. pneumonia* which is totally resistant against both ampicillin and tetracycline. The zone of inhibition of up to 17mm is recorded for tetracycline against *E. coli*. Ampicillin is found to be resistant against *A. hydrophilia*, *A. baylyi* and *P. aeruginosa*. Similarly, *B. cereus* and *S. pyogene* were resistant to ampicillin, against whilst *E. faecalis* showed some activity against ampicillin.

Figure 3: Growth inhibitory activity of *T. ammi* seed extracts and reference antibiotics against gram-positive bacterial species (a = *B. cereus*, b = *E. faecalis*, c = *S. pyogenes*) measured as ZOIs (mm) $\pm$ SEM. M= methanol, W= water, E= ethyl acetate, C= chloroform, H= hexane, A= ampicillin (2 μ g); T= tetracycline (10 μ g); NC = negative control. All assays were completed twice, each with internal triplicates ($n=6$) and the results are expressed as mean zones of inhibition (mm) $\pm$ SEM.

DISCUSSION

Due to the continuous evolution in bacterial resistance to many antibiotics, the development of new antibiotic therapies is the utmost priority of medical science.^{15,22} A simultaneous decline in the discovery of new antibiotics by traditional ways has raised interest in re-examining medicinal plants for new antibiotic therapies.²⁴ *T. ammi* belongs to the family of Apiaceae and is a highly valued medicinally important seed. The Apiaceae plant family includes celery, carrot and parsley. A common feature of these species is that they contain a group of bioactive aliphatic C₁₇ called polyacetylenes. Polyacetylenes possess potent antimicrobial effects against panels of bacteria, fungi, mammalian cells.²⁵ They also display anti-inflammatory, anti-platelet-aggregatory and neurotoxic effects.²⁵ *T. ammi* also possesses hypolipidemic,²⁶ antimicrobial,²⁷ antihypertensive, antispasmodic, hepatoprotective, bronchodilating²⁸ and digestive effects.²⁹

A single assay technique was used to screen for the antibacterial activity in this study. Disc diffusion assays were used in this study as it is a rapid method and has been widely used in other studies. The disc diffusion assay relies on the diffusion of molecule through the aqueous environment of an agar gel, this procedure may be affected by the solubility of the extract compounds in aqueous environment. Chloroform, ethyl acetate, hexane, methanol and water extracts of *T. ammi* were used in the disc diffusion assays in this study. Notably, the lower polarity extracts would contain lower polarity compounds which may be unable to diffuse readily through the agar gel. Therefore, false negatives may have been obtained against those extracts. Other methods, including liquid dilution assays may have provided a more accurate measurement of the

antibacterial activity of those extracts and such studies are planned in the future. Furthermore, liquid dilution assays are generally considered to be more sensitive than solid phase inhibition assays.

Notably, our study tested against a limited panel of bacterial pathogens. It is possible that inhibitory activity may have been detected if the extracts were tested against an extended panel of bacteria. Furthermore, the literature on the traditional uses of this species as a medicine is incomplete. The species is listed as useful in the treatment of several pathogenic diseases, yet the specific uses are not stated. As gastrointestinal and autoimmune diseases can be caused by multiple pathogen types, these extracts may have inhibitory properties against fungal, protozoal and viral pathogens, as well as against tumour cells and further testing is required to more completely evaluate the therapeutic potential of *T. ammi*.

CONCLUSION

Chloroform, ethyl acetate, hexane, methanol and water extracts of *T. ammi* displayed no antibacterial activity in the disc diffusion and liquid dilution assays against the panels of human pathogenic bacteria.

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CONFLICT OF INTEREST

The authors declare no conflicts of interest

ABBREVIATIONS USED

DMSO: Dimethyl sulfoxide; **LC₅₀**: The concentration required to achieve 50 % mortality; **MIC:** minimum inhibitory concentration; **ZOI:** zone of inhibition.

SUMMARY

- T. ammi* seed extracts were screened for the ability to block the growth of a panel of human bacterial pathogens.
- No inhibitory activity was evident against any of the bacterial species tested.
- Toxicity of the *T. ammi* extracts was determined using the Artemia nauplii toxicity bioassay.
- All extracts were non-toxic.

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