ESSENTIAL OILS AND METHYLGLYOXAL: A POSSIBLE ALTERNATIVE TREATMENT FOR ANTIBIOTIC RESISTANT BACTERIAL INFECTIONS

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ABSTRACT

Objective: Essential oils are of significant interest in today’s world of healthcare because these compounds have a variety of medicinal properties. In this study, we evaluated the in vitro antibiotic role of essential oils as a possible alternative treatment in combating Methicillin-resistant Staphylococcus aureus (MRSA).

Methods: In conjunction with carrier oils, three essential oils (cassia, cinnamon bark, and thyme), as well as methylglyoxal were tested on MRSA using the Kirby-Bauer disc diffusion method.

Results: The minimum inhibitory concentration of each tested essential oil and methylglyoxal in carrier oil was determined to be 25% essential oil and 75% carrier oil mixture. This concentration worked much more effectively than the standard antibiotic, vancomycin, which is currently used to treat MRSA infections.

Conclusion: Antibacterial emollients made from naturally occurring products like essential oils can be cost-effective alternatives to antibiotics. The results of this research show that these emollients are more effective against MRSA than standard antibiotics in cell culture.

Keywords: Methicillin-resistant Staphylococcus aureus, Essential oils, Cassia, Cinnamon bark, Thyme, Methylglyoxal, Antibiotic resistance

INTRODUCTION

Antibiotics have long been the “go-to” treatment for pathogenic bacterial infections [1]. Common antibiotics, such as penicillin and methicillin, have been used since the mid-1900’s. These antibiotics, once effective medications for treating bacterial infections, have rapidly become less useful due to the highly adaptive genomes of pathogenic bacteria [2]. Bacterial genomes contain plasmids, which are small, circular, extracellular pieces of DNA that allow the bacterium to evolve quickly and adapt to its environment [3]. As antibiotics were being used, and oftentimes, misused, many bacterial species developed a resistance gene against the antibiotic within its plasmid DNA [4].

Antibiotic resistant bacteria are receiving national attention in the United States. The White House and the CDC are working tirelessly to find ways to help prevent the antibiotic resistant bacteria pandemic from worsening by releasing an annual update on the matter to the public. In the latest 2014 update, the White House released this statistic: “[The CDC estimates that] annually at least two million illnesses and 23,000 deaths are caused by antibiotic-resistant bacteria in the United States alone” [5]. Scientists have been researching for years for effective ways to treat such infections like MRSA that are resistant to common antibiotics, resulting in the creation of newer classes of FDA-approved antibiotics which are costly and possibly short-lived in efficacy as it has been seen with earlier antibiotics.

Antibiotics can kill bacteria, both helpful and pathogenic, within an organism. Broad-spectrum antibiotics have severe side effects in many patients who are taking them. These side effects include nausea, vomiting, diarrhea, numbness, urinary retention, and even the possibility of becoming more susceptible to other infections due to the fact that the body’s microbiota (like E. coli, Lactobacillass species) are being destroyed along with the pathogenic bacteria. With options growing thin and unpleasant, some scientists have turned towards natural products, such as essential oils, for therapy.

Plants and their derivatives have been used for medicinal purposes as far back as the Egyptian civilizations in 2800 BC [6]. Essential oils are chemical compounds that can be hydrocarbons or oxygenated compounds like esters, aldehydes, alcohols, and phenols that are derived from plants. For therapeutic purposes, essential oils are usually diluted in a carrier before being applied to the skin to prevent irritation [6]. In this study, the essential oils cassia, cinnamon bark, and thyme as well as a key ingredient in Australia’s popular Manuka honey, methylglyoxal, were being tested for efficacy as emollients with several carrier oils in treating MRSA infections.

These essential oil compounds showed promise against antibiotic resistant bacteria in previous studies conducted at Monmouth University [7-8]. These studies showed that essential oil emollients could be excellent alternative treatments for topical infections caused by antibiotic-resistant bacteria.

MATERIALS AND METHODS

Bacterial strains and growth conditions

MRSA samples were supplied by Dr. Albert Rojtman, Chief of Clinical Microbiology at Jersey Shore University Medical Center in Neptune, NJ, USA. The anonymous samples were originally cultured on blood agar plates. For experimentation, the bacteria were grown under aerobic conditions on Mueller Hinton II agar media over a 24 hour incubation period at 37°C.

Obtaining essential oils

Methylglyoxal (40% in H2O) was obtained by mail order from Sigma Aldrich® of St. Louis, MO, USA. Cassia, cinnamon bark, and thyme were obtained by mail order from doTERRA Essential Oils® of Pleasant Grove, UT, USA.

Obtaining carrier oils

Jojoba oil was obtained by mail order from Sigma Aldrich® of St. Louis, MO, USA. Coconut oil was obtained by mail order from doTERRA Essential Oils® of Pleasant Grove, UT, USA. Liquid Lanolin was obtained by mail order from Now® Solutions of Bloomingdale, IL, USA. Olive oil (Filippo Berio®) was purchased at the local supermarket. All four carrier oils are safe for use on skin.

Conclusion: Antibacterial emollients made from naturally occurring products like essential oils can be cost-effective alternatives to antibiotics. The results of this research show that these emollients are more effective against MRSA than standard antibiotics in cell culture.

Keywords: Methicillin-resistant Staphylococcus aureus, Essential oils, Cassia, Cinnamon bark, Thyme, Methylglyoxal, Antibiotic resistance
Standard antibiotics utilized

Vancomycin susceptibility test discs (30 μg) and rifampin susceptibility test discs (5 μg) were obtained by mail order from Becton, Dickinson and Company of Franklin Lakes, NJ, USA.

Preparation of Mueller Hinton II agar medium

Mueller Hinton II agar was purchased by mail order from Becton, Dickinson and Company of Franklin Lakes, NJ, USA. Mueller Hinton II agar was chosen as a nutritive source for growing the MRSA bacteria as well as its suitability for performing susceptibility testing. The media was prepared as per manufacturer’s instructions under sterile conditions. Prepared media plates were stored at 4 °C until ready for use.

Kirby-bauer disc diffusion

A single MRSA colony from the blood agar plate provided by JSUMC was reconstituted in tryptic soy nutrient broth to match a 0.5 McFarland standard 1 x 10^8 colony forming unit per milliliter (CFU/ml) volume. The broth culture was then swabbed onto the Mueller Hinton II agar plates.

Sterile blank 6 mm paper discs were placed in the center of each swabbed plate. A volume of 5 μl of each emollient was pipetted onto the blank disc. Vancomycin (30 μg) and rifampin (5 μg) antibiotic susceptibility test discs were also placed on swabbed plates to act as controls. After a 24 hour incubation period at 37 °C, the diameters of zones of inhibition around each disc were measured in millimeters [9]. The experiments were performed in triplicate and the zone of inhibition measurements were averaged to yield a mean zone of inhibition.

RESULTS AND DISCUSSION

Antibiotic discs of both vancomycin (30 μg) and rifampin (5 μg) were both tested against the MRSA in every experiment. Vancomycin averaged a zone of inhibition of 19 mm and rifampin averaged a zone of inhibition of 36 mm (table 1). Rifampin is used as an adjuvant only in very severe cases of MRSA when biofilm formation is suspected, so vancomycin is still the current standard of care [10]. However, it is apparent that vancomycin is losing its effectiveness against MRSA because a growing number of MRSA infections are showing antibiotic treatment failures.

Table 1: Mean diameter of zone of inhibition of standard antibiotics used to treat MRSA

| Standard antibiotic | Mean diameter of zone of inhibition (mm) |
|---------------------|----------------------------------------|
| Vancomycin          | 19                                     |
| Rifampin            | 36                                     |

In order to effectively determine the minimum inhibitory concentration (MIC) of the essential oils, the experiments began by testing a concentration of 100% of each carrier oil against the MRSA to determine that the carriers chosen did not show any antibacterial effect of their own against the MRSA. A volume of 5 μl of fractionated coconut oil, jojoba oil, lanolin oil, and olive oil was pipetted onto a sterile blank paper disc and tested in triplicate against the MRSA. All proved to have no antibacterial properties because there was no zone of inhibition surrounding each carrier oil disc in every trial (table 2).

Table 2: Mean diameter of zone of inhibition of carrier oils at 100% concentration used against MRSA

| Essential oil       | Mean diameter of zone of inhibition (mm) |
|---------------------|----------------------------------------|
| Fractionated coconut oil | 0.0                                   |
| Jojoba oil          | 0.0                                    |
| Lanolin oil         | 0.0                                    |
| Olive oil           | 0.0                                    |

Once the carrier oils were proved to have no bactericidal properties, 5 μl of the essential oils were tested at 100% concentration to show that each essential oil chosen did, in fact, show antibacterial properties specific to MRSA. Each of the three essential oils, cassia, cinnamon bark, and thyme, as well as methylglyoxal, the main component of Australia’s popular Manuka honey, all formed large zones on inhibition that were larger than the zones of inhibition of both tested antibiotics (table 3).

Table 3: Mean diameter of zone of inhibition of essential oils at 100% concentration used against MRSA

| Essential oil       | Mean diameter of zone of inhibition (mm) |
|---------------------|----------------------------------------|
| Cassia              | 40.3                                   |
| Cinnamon bark       | 38.7                                   |
| Methylglyoxal       | 43.3                                   |
| Thyme               | 37.3                                   |

The essential oil concentration was halved to 50% for the first dilution trial. The remaining 50% of the volume contained one of the four carrier oils. Each essential oil was tested in combination with each of the four carriers against the MRSA. Dilution was necessary because although in this experiment emollients were not being tested on skin, essential oils at their full concentration are normally known to be very irritant on the skin and would not be a feasible treatment on an open wound, like a MRSA infection.

Table 4a: Mean diameter of zone of inhibition of 50% essential oil and 50% carrier oil emollient against MRSA

| Essential oil | Carrier oil  | Mean diameter of zone of inhibition (mm) |
|---------------|--------------|----------------------------------------|
| Cassia        | Coconut      | 36.7                                   |
|               | Jojoba       | 44                                      |
|               | Lanolin      | 35                                      |
|               | Olive        | 38.3                                   |
| Cinnamon bark | Coconut      | 31.3                                   |
|               | Jojoba       | 40.3                                   |
|               | Lanolin      | 33.7                                   |
|               | Olive        | 36.3                                   |
| Methylglyoxal | Coconut      | 41.7                                   |
|               | Jojoba       | 43                                      |
|               | Lanolin      | 38                                      |
|               | Olive        | 42.3                                   |
| Thyme         | Coconut      | 7.7                                     |
|               | Jojoba       | 14                                      |
|               | Lanolin      | 16.7                                   |
|               | Olive        | 12.7                                   |
Hence, the goal of finding the lowest dilution concentration of essential oil emollient that would still be deemed more effective than the standard antibiotic. At a 50% dilution, both cassia, cinnamon bark, and methylglyoxal all worked significantly better than vancomycin based on measured zone of inhibition. Of those three, cassia and methylglyoxal worked the best, working as well or better than even the strong rifampin antibiotic. Thyme, an essential oil that worked well on its own, did not work well in conjunction with carrier oils (table 4a) and therefore was excluded from further dilution trials.

Cassia, cinnamon bark, and methylglyoxal were chosen to continue the dilution trials. The essential oil concentration was halved once again to 25%. The remaining 75% of the emollient consisted of a carrier oil. Cassia, cinnamon, and methylglyoxal mixed with lanolin showed the weakest results. Although the results for lanolin carrier oil emollients were less effective overall than rifampin, this emollient still worked better than vancomycin. The strongest results were from cassia-jojoba oil, cassia-olive oil, methylglyoxal-olive oil, and methylglyoxal-coconut oil emollients, all of which continued to work more effectively than even rifampin (table 4b).

| Essential oil | Carrier oil | Mean diameter of zone of inhibition (mm) |
|---------------|-------------|-----------------------------------------|
| Cassia        | Coconut     | 32                                      |
|               | Jojoba      | 39.7                                    |
|               | Lanolin     | 35.7                                    |
|               | Olive       | 39.3                                    |
| Cinnamon bark | Coconut     | 25.7                                    |
|               | Jojoba      | 29.7                                    |
|               | Lanolin     | 25.3                                    |
|               | Olive       | 31.7                                    |
| Methylglyoxal | Coconut     | 39                                      |
|               | Jojoba      | 35.3                                    |
|               | Lanolin     | 31                                      |
|               | Olive       | 38.7                                    |

In a second trial, the essential oil concentration was halved to 12.5% with the remaining 87.5% as a carrier oil. In this trial, the results were clearly disappointing. The cassia and cinnamon bark emollients worked about as effectively as vancomycin. Methylglyoxal still had a slight advantage over vancomycin, but none of the emollients were nearly as effective as rifampin (table 4c).

| Essential oil | Carrier oil | Mean diameter of zone of inhibition (mm) |
|---------------|-------------|-----------------------------------------|
| Cassia        | Coconut     | 20.3                                    |
|               | Jojoba      | 23.6                                    |
|               | Lanolin     | 20                                      |
|               | Olive       | 20.3                                    |
| Cinnamon bark | Coconut     | 13                                      |
|               | Jojoba      | 22.6                                    |
|               | Lanolin     | 19.3                                    |
|               | Olive       | 18                                      |
| Methylglyoxal | Coconut     | 27.3                                    |
|               | Jojoba      | 29                                      |
|               | Lanolin     | 21                                      |
|               | Olive       | 24.3                                    |

At 25% dilution, all of the emollients worked significantly better than the vancomycin antibiotic. Several of the emollients at this concentration continued to exceed the results of rifampin. The results from the lowest dilution concentration (12.5%) were unremarkable. Of the 25% dilutions, four had results that particularly stood out: cassia-jojoba oil, cassia-olive oil, methylglyoxal-olive oil, and methylglyoxal-coconut oil. All four of these emollients surpassed not only the vancomycin standard but also the rifampin antibiotic (fig. 1). Thus, the MIC of these essential oils were concluded to be 25%.

The structures of methylglyoxal and cinnamaldehyde, which is the main active ingredient that makes up 75-90% of cassia, contain aldehyde structures (fig. 2a, 2b). Aldehydes have been known to confer antimicrobial activity and have been an important component of laboratory disinfectants for years because of their ability to quickly and effectively kill pathogenic bacteria. Aldehydes act by inhibiting and preventing peptide and protein synthesis, and a bacterial cell that cannot make more proteins cannot persist [11]. The fact that these two essential oils contain aldehyde structures may help explain their antibacterial activity.
CONCLUSION
Antibacterial emollients made from naturally occurring products like essential oils can be cost-effective alternatives to antibiotics. As a natural product that would be used on the skin, it could also potentially prevent the unpleasant side effects that go along with taking traditional antibiotics. This can also provide a possible alternative for those who prefer more natural methods of treatment, and for underserved populations that do not have access to or cannot afford other means of care. Cassia and methylglyoxal have proven themselves in this study to have strong bactericidal properties at concentrations as low as 25% in carrier oil emollients, working even better than the current standard of care, vancomycin.

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CONFLICTS OF INTERESTS
Declared none

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