

# Synergistic Potential of Cinnamon Extracts, Antibiotics, and Alum on Selected Clinical Microbes.

## ABSTRACT

Antibiotics are natural or synthetic substances used to inhibit or kill susceptible microorganisms. Due to the increase rate of bacterial resistance to antibiotics, there is the need to venture in other methods involving the coalescence effect of antibiotics and other substances such as cinnamon and alum to effectively eradicate these pathogenic organisms using less concentrated conventional antibiotics. The minimum inhibitory concentration (MIC) and the minimum bactericidal concentration (MBC) of the selected antimicrobial agents were determined using the broth dilution method, following standards and procedures set by the national council of clinical laboratory standards (NCCLS). The synergistic potentials of all the antimicrobics used were analyzed by the checkerboard assay. The MIC values of 25, 0.25, 0.75, 25mg/ml were obtained for cinnamon extracts, gentamicin, penicillin G, and aq. Alum respectively against *S. aureus*. For *E. coli O157H:H7*, the MIC values of cinnamon extracts, gentamicin, penicillin G, and aq. Alum were 12.5, 0.25, 0.75 and 25mg/ml respectively, while the MIC values of cinnamon extracts alum and fluconazole for *C. albicans* were 12.5, 25 and 0.05mg/ml respectively. There is a reduction in the MIC values for the various combinations against the same clinical microbes. All the antimicrobics used in this study inhibited the growth of the clinical microbes. All the microbes used in this study inhibited the tested microbes in dose dependent manner. The combinations of these agents resulted in reduction in MIC and MBC values, with an additive, indifference, and antagonistic effects using checkerboard assay which resulted to a fractional inhibitory concentration index which ranged from 0.56 to 7.9. Therefore, application of cinnamon, alum, and their combinations with conventional antibiotics can be used as treatment measure against infections caused by *S. aureus*, *E. coli O157H:H7*, and *C. albicans*.

**Keywords:** Alum, Cinnamon extracts, fractional inhibitory concentration index, minimum bactericidal concentration, synergistic effects.

## INTRODUCTION

Antimicrobial agents are substance that are used to suppress the growth of microorganisms while antibiotics are natural or synthetic agents that are used to inhibit or kill microorganisms (AL-Khikani et al., 2023). Antibiotics having the ability to kill microbes are known as Bactericidal agents, whereas those therapeutic agents having the ability to inhibit/stop the growth and proliferation of microorganisms are known as Bacteriostatic agents (Tosun et al., 2022). The misuse and overuse of antibiotics in recent years has increased the number of microbes that are resistant to antibiotics which is a major challenge to clinicians and public health practitioners (Becerril et al., 2012). The world health organization (WHO) has compiled a list of multidrug

resistant microbes with a call for urgent discovery of new therapeutic agents (Miethke et al., 2021). Initially the synthesis of antimicrobial agent was mainly based on single natural substance which was obtained from several sources such as trees/plants, animal, and microorganisms (fungi and bacteria) (Ali 2018).

Cinnamon powder is a spice obtained from the bark of the tree *Cinnamomum zeylanicum*. It is used as a preservative due its antimicrobial properties. Cinnamon is mostly found in countries such as China, Sri Lanka Indonesia, and India (Ravindran et al., 2004). cinnamaldehyde and eugenol are the two major antimicrobial compounds found in cinnamon resulting to its antimicrobial activity against a variety of Gram negative and Gram-positive bacteria as well as fungi. Cinnamaldehyde which is one of the major bioactive molecules in cinnamon is electronegative in nature and can interfere with cellular biological processes, especially the nitrogen containing compounds including the nucleic acids and proteins (Nabavi et al., 2022). Also, cinnamon and its essential oil (EO) has been shown to inhibit bacterial growth through the following mechanisms: ATPase, Biofilm formation membrane porin, anti-quorum sensing effects, cell division inhibitions, and alterations of the cell membrane, these mechanisms could cause the reduction in susceptibility of bacteria to antibiotics, decreasing the dose of antibiotic necessary for the effective treatment, hence a decrease in the toxic side effect of the drug (EL Atkiet al., 2019; Nabavi et al., 2022). However, there is an increased keen interest in using active compound from organic and inorganic material as source of active ingredient for the synthesis of antimicrobial agents. cinnamon tree has shown a promising source of natural active compound necessary for the synthesis of new antimicrobial drugs (Cava-Roda et al., 2021). This is due to the fact that many studies have been devoted on it, and it has been shown that cinnamon and its derivatives have a strong antimicrobial effect on microbes (Fadli et al., 2012).

Furthermore, it has also been shown that the combine effect of cinnamon and antibiotic produces a better and increased antimicrobial effect on tested clinical microbes.

Alum is an inorganic compound formed from natural substances, Alum is generally known as a hydrated double salt of aluminum with general formulae  $XAl(SO_4)_2 \cdot 12H_2O$ . Alum can exist in nature as pure and impure form (Ali, 2018). Alum is a compound formed from the combination of two or more separate elements, there are various types of alum for example potash alum is a combination of potassium, aluminum, sulphur, and oxygen, also known as potassium aluminum sulphate (Amadi, 2020). It is used as antiseptic and as astringent in several food preparatory and preserving processes/procedures as well as water purification (Amadi *et al.*, 2016). Alum can also be used in the following ways: acts as mordant (binder) use in dyeing, fixation of dye to cotton, as well as other fabrics, as fire extinguisher, as astringent in medicine, and adjuvant in vaccine production. The antimicrobial effect of alum has been reported and documented (Bnyanet *al.*, 2014; Amadi *et al.*, 2016).

Also, its medical benefits as immune enhancer, used as adjuvant in vaccine production such as hepatitis A & B (Bestoon, 2012; Amadi *et al.*, 2016). Often times a single antimicrobial agent is not sufficient to effectively eradicate an infectious agent, therefore in order to achieve the desired antimicrobial affect, a combination of drugs or agents is used to produce the desired antimicrobial effect not obtained with a single antimicrobial agent (Amadi *et al.*, 2016). Finally, there is a limited research report on the coalescence of cinnamon, and antibiotics which shows a strong synergistic effect against several clinical microbes (EL Atki *et al.*, 2019; Umar *et al.*, 2017) However, to the best of my knowledge there is no available research data about the combine antimicrobial effect of cinnamon, antibiotics, and Alum against *Staphylococcus aureus*, *E. coli O157H:H7* and *Candida albicans*. Hence the aim of this research is to investigate the

possible synergistic antimicrobial effects of cinnamon, antibiotic (gentamicin and penicillin G) and Alum against *Staphylococcus aureus*, *E. coli* 0157H:H7 and *Candida albicans*.

## MATERIALS AND METHODS

### Experimental Design

Crude cinnamon bark extracts (aqueous and ethanolic) of the plant *Cinnamomum zeylanicum* were prepared in a Soxhlet assembly using hot extraction method. Analytical grade alum was purchased from the chemical store with the required standard concentration and was diluted serially using distilled water as solvent. The antimicrobials (antifungal and antibiotics) - Gentamicin, Penicillin G and Fluconazole were purchased from the pharmaceutical store. The antimicrobials (crude aqueous and ethanolic cinnamon bark extracts, aqueous alum, Gentamicin, Penicillin G, and Fluconazole) were then screened, and their antimicrobial activity were evaluated using Broth dilution assay to determine the Minimum inhibitory concentration (MIC[mg/ml]), minimum microcidal (bactericidal/fungicidal) concentration [mg/ml] against selected clinical microbes ( *S. aureus* = Gram positive, *E. coli* 0157H:H7 = Gram negative, *C. albicans* = fungi). The MIC values obtained were used to calculate the fractional inhibitory concentration index (FICI) of the antimicrobial agents, which were subsequently used to classify the antimicrobial effects as synergistic, additive, indifference or antagonistic.

### Preparation and Extraction of Cinnamon (*Cinnamomun Zeylanicum*) Bark

About 300 grams of *Cinnamomun zeylanicum* was purchased from the Mile 3 market, Port Harcourt, Nigeria. After sterilization of the blender by autoclaving at 121°C for 15 minutes, allowed to cool, and thereafter used to grind the *Cinnamomun zeylanicum* bark into powder form, 100 grams of the powdered *Cinnamomun zeylanicum* was weighed out and was used for

ethanol extraction while another 100 grams was used for the aqueous extraction. The extraction was performed using the Soxhlet extractor at the ratio of 50 mg: to 1mL, that is 2000 mL of ethanol was used for 100grams (100,000mg) of the *Cinnamomun zeylanicum* powder. Furthermore, ethanol available in the extract was removed/ eliminated by the use of rotary evaporator, leaving a sediment of crude extract with a concentration of 50mg/ml. the crude extract was then stored at an appropriate temperature until it was used on the selected clinical isolates. The same procedures and method were used for the aqueous extraction. same volume of solvent and mass of *Cinnamomun zeylanicum* but in place of ethanol, distilled water was used.

#### **Preparation of Aqueous Alum**

Analytical grade alum was purchased from the chemical store,manufactured by Vicker's Laboratories LTD. Bunley in Wharfedale west Yorkshire England. 5000mg was dissolved in 50ml of distilled water to produce an initial concentration of 100 mg /ml. This concentration was further diluted serially. The serially diluted alum with varying concentrations was tested on the selected clinical isolates to ascertain the antimicrobial properties of alum on the selected pathogens.

#### **Sources of Selected Clinical Microbes**

The Microorganism used this study were *S. aureus* = Gram positive, *E. coli* O157H:H7 = Gram negative, *C albicans*= fungi) obtained from different site of the body of an infected patient. *S. aureus* isolated from topical wound infection, *E. coli* O157H:H7 was isolated from stool sample, and *C albicans* was isolated from oral thrush patent. These clinical isolates were procured from the University of Port Harcourt Teaching Hospital (UPTH). The Selected Clinical Microbes were cultured on different media where colonial morphology were used to confirm the identity of the microorganisms.

### **Culturing of *E. coli* O157H:H7**

*Escherichia coli* serotype O157H:7H was inoculated on sorbitol MacConkey agar prepared accurately according to the manufacturer's instruction. Sorbitol MacConkey agar used is a selective, differential media which was used for the detection and differentiation of the *Escherichia coli* serotype O157H:7H from other strains of *Escherichia coli*. The media was autoclaved at 121°C for 15 minutes, and was poured on a sterile petri dishes, allowed to solidify. The plates were dried and the organisms was inoculated by streak plate method. The inoculated plates were incubated at 37°C for 24 hours. The growth of colorless colonies was an indication of *Escherichia coli* serotype O157H:H7. However, other strains of *Escherichia coli* other than O157H:H7 were pink colonies growth due to the ability to ferment sorbitol. The media inhibits the growth of other Gram-positive bacteria such as *Staphylococcus aureus*.

### **Culturing of *Candida albicans***

Sabouraud Dextrose agar was used for the growth and maintenance of the pathogenic fungi *Candida albicans* using the cultural morphology and color. The Sabouraud dextrose agar was prepared according to the manufacturer's instructions, incorporated in the agar was 50mg of chloramphenicol and 5mg of gentamicin to inhibit bacteria growth. The media was autoclaved at 121°C for 15 minutes and was poured on a sterile petri dishes then allowed to cool at room temperature. The organisms were inoculated by streak plate method and was incubated for 24 hours at 37°C. the result showed creamy pasty colonies with good growth for *Candida albicans*, and there was no growth for *Escherichia coli* and *Staphylococcus aureus*. Other morphological and biochemical investigations was performed to confirmed the identity of the fungi.

### **Determination of Minimal Inhibitory Concentration**

Mueller Hinton broth was prepared according to the manufacturer's instruction, it was autoclaved at 121°C for 15 minutes, and 0.5ml each of the sterilized broth was pipette into different test tubes. Exactly 0.5ml of the antimicrobial agent was added to the broth in the first test tube, the pipette was discarded, with a fresh pipette, the content of the first test tube was mixed thoroughly and 0.5ml was transferred into the second test tube containing 0.5ml of the sterile broth, again the pipette was discarded and a fresh pipette was used to mixed the content of the second test tube, and 0.5 ml was transferred to the third test tube; the dilution process was continued throughout the test tube number. However, the antimicrobial agent was not added to the test tubes labelled positive growth control and the sterility control. After the content of the last test tube was mixed, 0.5 ml was removed from the tube and was discarded. So that the content of all the test tube be uniformed, that is, 0.5ml in all test tubes.

From a plate culture of the clinical isolates a suspension of the organism was prepared in 5ml of normal saline which is equivalent to 0.5 MarFarland standard. With the aid of a sterile pipette 0.1ml of the microbe in broth was transferred to 9.9ml of normal saline. With a fresh pipette the content of the test tube was mixed, and 0.1ml of the inoculum were transferred to the test tubes containing the antibiotics, and to the test tube labelled growth control; however, none was added to the test tube labelled sterility control. The test tubes were mixed by gently shaking the test tubes. The tubes were incubated at 37°C for a duration of 18-24 hours. After the said duration elapsed, the tubes with the lowest concentration of the microbial agent with no visible growth was considered. And the concentration of the tube was reported as the minimal inhibitory concentration of the antimicrobial agent, as there was visible growth on the positive growth control tube and no growth on the sterility control tube.

#### **Determination of Minimal Bactericidal Concentration.**

The minimal lethal concentration was determined by the transfer of sample from the tubes with no visible growth from the test tube of the minimal inhibitory concentration test onto a sterile agar medium on Petri dishes by streak plate method, and thereafter incubated for 24- 48 hours at 37°C. the lowest concentration or the highest dilution of antimicrobial agent that do not show any visible growth on the agar plate was considered the minimal bactericidal concentration.

### **Evaluation of Interaction Between the Antimicrobials**

This checkerboard assay was used to evaluate the interactions between cinnamon extracts, antibiotics, and alum. The procedure was performed according to the methods conducted by Moody et al., 2003 with modifications AL ALK et al., 2019. Briefly, six concentrations of alum, and five concentrations of each cinnamon extracts were prepared in a sterilized tubes by serial dilutions. Subsequently, Alum at decreasing concentrations going from 100mg/ml through 3.13mg/ml were added introduced horizontally into the tubes. In the same manner the cinnamon extracts at decreasing concentrations starting from 50mg/ml through 3.13mg/ml were introduced vertically. The final volume in each tube was 8ml, comprising of 1ml of each alum and cinnamon extracts, as well as 6ml of Mueller Hinton (MH) media consisting of 10<sup>6</sup>CFU/ml of the selected clinical microbe suspension. The tubes were incubated at 37°C for 24 hours. The analysis of the consortiums were obtained by calculating the FICI (AL ALK et al., 2019; Ali et al., 2018). The same methods and procedures were repeated by combining Alum or Cinnamon extracts with different concentrations of Gentamicin (0.25, 0.13, 0.06, and 0.03mg/ml), Penicillin G (1.50, 0.75, 0.37, 0.19) and fluconazole (0.2, 0.1, 0.05, 0.025, 0.013mg/ml), against *S. aureus*, *E. coli* O157H:H7, *C albicans*. The fractional inhibitory concentration was calculated using the following formula.



$$FICI = FIC_A + FIC_B$$

$$= A/MIC_A + B/MIC_B$$

Where A and B are the minimal inhibitory concentration of antimicrobial A and B combined in the same tube, whereas  $MIC_A$  and  $MIC_B$  were the minimal inhibitory concentration of each of the two antimicrobial agents respectively. The value of the fractional inhibitory concentration index was used to classify the interaction between the two antimicrobial agents combined together using the following statements as guideline (Wendakoon, & Sakaguchi 1995).

1. The two antimicrobial agents have a synergistic effect when the fractional inhibitory concentration index is less than 0.5 ( $FICI \leq 0.5$ )
2. When the value of the fractional inhibitory concentration index is between 0.5 and 1, the two antimicrobial agents have an additive effect.
3. The two antimicrobial agents have an indifferent effect when the fractional inhibitory concentration index is greater than 1 and less than 4 ( $1 < FICI < 4$ )
4. The combination of two antimicrobial agents was considered antagonistic when the fractional inhibitory concentration index is greater than 4 ( $FICI \geq 4$ )

## RESULTS

### Minimum Inhibitory Concentration MIC

The MIC values of the crude cinnamon extracts for *S. aureus* were 25mg/ml, 0.25mg/ml for gentamicin, 25mg/ml for Aqueous Alum and 0.19mg/ml for Penicillin G. The lowest MIC value was with Penicillin G against *S. aureus* (0.19mg/ml), and the highest MIC value for *S. aureus* were 25mg/ml with cinnamon extracts and Aqueous Alum. For *E. coli* O157H:H7, the MIC values ranged from 0.25 to 25mg/ml for all the antimicrobial agents used, with the lowest MIC value of 0.25mg/ml for gentamicin and the highest MIC value of 25mg/ml with Aqueous Alum

and cinnamon extracts. The recorded MIC value of the antimicrobial agents against *C. albicans* was between 0.05 and 25 mg/ml, Fluconazole showed the lowest MIC value of 0.05 mg/ml and the highest was 12.5 mg/ml with Aqueous Alum for the uncombined antimicrobial agents. The combinations of these antimicrobial agents resulted in lower MIC values when compared to the individual antimicrobial agents.

**Table 1: MIC and MBC values of different concentrations of aqueous and ethanolic *cinnamomum zeylanicum* extracts on *S. aureus*, *E. coli* O157H:H7, and *C. albicans*.**

Antimicrobial Agent	Organisms	MIC and MBC values mg/ml	
		MIC	MBC
Aqueous Extract	<i>S. aureus</i>	25	50
	<i>E. coli</i> O157H:H7	12.5	25
	<i>C. albicans</i>	12.5	12.5
Ethanolic Extract	<i>S. aureus</i>	25	50
	<i>E. coli</i> O157H:H7	12.5	25
	<i>C. albicans</i>	12.5	12.5

**Table 2: MIC and MBC values of different concentrations of gentamicin, penicillin G, and fluconazole Against *S. aureus*, *E. coli* O157H:H7, and *C. albicans*.**

Antimicrobial Agent	Organisms	MIC and MBC values mg/ml	
		MIC	MBC
Gentamicin	<i>S. aureus</i>	0.25	0.5
	<i>E. coli</i> O157H:H7	0.25	0.5
Penicillin G	<i>S. aureus</i>	0.19	0.37
	<i>E. coli</i> O157H:H7	0.75	1.5
Fluconazole	<i>C. albicans</i>	0.05	0.1

**Table 3: MIC and MBC values of different concentrations of Alum on *S. aureus*, *E. coli* O157H:H7, and *C. albicans*.**

antimicrobial agent	Organisms	MIC and MBC values mg/ml	
		MIC	MBC

<b>Alum</b>	<i>S. aureus</i>	25	50
	<i>E. coli O157H:H7</i>	25	50
	<i>C. albicans</i>	25	12.5

**Table 4: MIC and MBC values of different combinations and concentrations of gentamicin, penicillin G, and fluconazole on *S. aureus*, *E. coli O157H:H7*, and *C. albicans*.**

Antimicrobial Agent		Organisms MIC and MBC values mg/ml	
		MIC	MBC
<b>Gentamicin + Penicillin G</b>	<i>S. aureus</i>	0.13+0.75	0.25+1.15
	<i>E. coli O157H:H7</i>	0.08+0.37	0.13+0.75
	<i>C. albicans</i>	-	-
<b>Gentamicin + Fluconazole</b>	<i>S. aureus</i>	0.13+0.10	0.25+0.20
	<i>E. coli O157H:H7</i>	0.13+0.10	0.25+0.20
	<i>C. albicans</i>	0.08+0.05	0.13+0.10
<b>Penicillin G + Fluconazole</b>	<i>S. aureus</i>	0.75+0.10	1.50+0.20
	<i>E. coli O157H:H7</i>	0.75+0.10	1.50+0.20
	<i>C. albicans</i>	0.37+0.05	0.75+0.10

**Table 5: MIC and MBC values of different combinations and concentrations of gentamicin, penicillin G, fluconazole and aqueous Alum on *S. aureus*, *E. coli O157H:H7*, and *C. albicans*.**

Antimicrobial Agent		Organisms MIC and MBC values mg/ml	
		MIC	MBC
<b>Gentamicin + Alum</b>	<i>S. aureus</i>	0.08+25	0.13+50
	<i>E. coli O157H:H7</i>	0.08+25	0.13+50
	<i>C. albicans</i>	0.13+50	0.25+100
<b>Penicillin G</b>	<i>S. aureus</i>	0.37+25	0.75+50

+ Alum	<i>E. coli</i> O157H:H7	0.37+25	0.75+50
	<i>C. albicans</i>	0.75+50	1.50+100
Fluconazole + Alum	<i>S. aureus</i>	0.10+50	0.20+100
	<i>E. coli</i> O157H:H7	0.10+50	0.20+100
	<i>C. albicans</i>	0.05+25	0.10+50

**Table 6: MIC and MBC values of different combinations and concentrations of gentamicin, penicillin G, fluconazole and aqueous extract of *cinnamomun zeylanicum* on *S. aureus*, *E. coli* O157H:H7, and *C. albicans*.**

Antimicrobial Agent	Organisms	MIC and MBC values mg/ml	
		MIC	MBC
Gentamicin + Aqueous Cinnamon Extract	<i>S. aureus</i>	0.08+12.5	0.13+25
	<i>E. coli</i> O157H:H7	0.08+12.5	0.13+25
	<i>C. albicans</i>	0.13+25	0.25+50
Penicillin G + Aqueous Cinnamon Extract	<i>S. aureus</i>	0.19+6.25	0.37+12.5
	<i>E. coli</i> O157H:H7	0.37+12.5	0.75+25
	<i>C. albicans</i>	0.75+25	1.50+50
Fluconazole + Aqueous Cinnamon Extract	<i>S. aureus</i>	0.75+25	1.50+50
	<i>E. coli</i> O157H:H7	0.75+25	1.50+50
	<i>C. albicans</i>	0.37+12.5	0.75+25

**Table 7: MIC and MBC values of different combinations and concentrations of gentamicin, penicillin G, fluconazole and ethanolic extract of *cinnamomun zeylanicum* on *S. aureus*, *E. coli* O157H:H7, and *C. albicans*.**

Antimicrobial Agent	Organisms	MIC and MBC values mg/ml	
		MIC	MBC
Gentamicin + Ethanolic Cinnamon Extract	<i>S. aureus</i>	0.06+12.5	0.13+25
	<i>E. coli</i> O157H:H7	0.06+12.5	0.13+25
	<i>C. albicans</i>	0.13+25	0.25+50
Penicillin G	<i>S. aureus</i>	0.19+6.25	0.37+12.5

<b>Ethanollic Cinnamon Extract</b>	<b>+</b> <i>E. coli O157H:H7</i>	0.37+12.5	0.75+25
	<i>C. albicans</i>	0.75+25	0.20+50
<b>Fluconazole</b> <b>+</b> <b>Ethanollic Cinnamon Extract</b>	<i>S. aureus</i>	0.10+25	0.20+50
	<i>E. coli O157H:H7</i>	0.05+12	0.10+25
	<i>C. albicans</i>	0.03+6.25	0.05+12.5

**Table 8: MIC and MBC values of different combinations and concentrations of aqueous alum, aqueous and ethanolic extract of *cinnamomunzeylanicum* on *S. aureus*, *E. coli O157H:H7*, and *C. albicans*.**

Antimicrobial Agent	Organisms	MIC and MBC values mg/ml	
		MIC	MBC
<b>Ethanollic Cinnamon Extract</b> <b>+</b> <b>Alum</b>	<i>S. aureus</i>	25+50	50+100
	<i>E. coli O157H:H7</i>	12.5+25	25+50
	<i>C. albicans</i>	6.25+12.5	12.5+25
<b>Aqueous Cinnamon Extract</b> <b>+</b> <b>Alum</b>	<i>S. aureus</i>	25+50	50+100
	<i>E. coli O157H:H7</i>	12.5+25	25+50
	<i>C. albicans</i>	6.25+12.5	12.5+25

#### **Minimum Bactericidal Concentration (mg/ml)**

The highest dilution of antimicrobial agent that yielded no antimicrobial growth as evidence of absent turbidity on the broth. The range of MBC in this investigation was 0.1 to 50mg/ml for all the antimicrobial agents conducted alone. The lowest MBC value of 0.1mg/ml was for Fluconazole against *C. albicans*, and the highest (50mg/ml) for *S. aureus* with cinnamon extracts.

#### **Fractional Inhibitory Concentration Index (FICI).**

Fractional inhibitory concentration index is the sum of the fractional inhibitory concentration FICs of each drug tested when used in combination. FIC is calculated by dividing each drug's MIC in combination by each drug's MIC when used alone. The fractional inhibitory concentration index showed

a ranged of 0.56 to 7.9. gentamicin + penicillin G showed a FICI of 0.56 as the lowest with *E. coli* O157H:H7, and the highest FICI was 7.9 with a combination of Fluconazole + aqueous cinnamon extract against *C. albicans*. The result for this present study showed additive effect ( $0.5 < \text{FICI} < 1$ ) Indifferent effect ( $1 < \text{FICI} < 4$ ), and antagonistic effect ( $\text{FICI} \geq 4$ ). This present study has also shown that combination of cinnamon extracts antibiotics (gentamicin, and penicillin G) and Aqueous Alum resulted in an additive, indifferent and antagonist effects against *S. aureus*, *E. coli* O157H:H7, and *C. albicans*.

**Table 9: Fractional inhibitory concentration index (FICI) of various combinations of gentamicin, penicillin and fluconazole against *S. aureus*, *E. coli* O157H:H7, and *C. albicans*.**

AntimicrobialAgent	Organisms	FIC <sub>A</sub>	FIC <sub>B</sub>	FICI	S, A, I, AN
<b>Gentamicin + Penicillin G</b>	<i>S. aureus</i>	0.52	0.5	1.02	I
	<i>E. coli</i> O157H:H7	0.3	0.2	0.56	A
	<i>C. albicans</i>	-	-	-	NT
<b>Gentamicin + Fuconazole</b>	<i>S. aureus</i>	0.52	-	0.52	NT
	<i>E. coli</i> O157H:H7	0.52	-	0.52	NT
	<i>C. albicans</i>	-	1	1.00	NT
<b>Penicillin G + Fuconazole</b>	<i>S. aureus</i>	0.5	-	0.5	NT
	<i>E. coli</i> O157H:H7	1	-	1.0	NT
	<i>C. albicans</i>	-	1	1.0	NT

Key: FIC<sub>A</sub> = Fractional inhibitory concentration of drug A, FIC<sub>B</sub> = Fractional inhibitory concentration of drug B, A=Additive, S=Synergistic, I = indifference, AN = antagonistic, FICI= Fractional inhibitory concentration index.

**Table 10: Fractional inhibitory concentration index (FICI) of various combinations of gentamicin, penicillin, fluconazole and ethanolic cinnamon extract against *S. aureus*, *E. coli* O157H:H7, and *C. albicans*.**

AntimicrobialAgent	Organisms	FIC <sub>A</sub>	FIC <sub>B</sub>	FICI	A, S, I, AN
<b>Gentamicin+ Ethanolic Cinnamon Extract</b>	<i>S. aureus</i>	0.24	0.48	0.72	A
	<i>E. coli</i> O157H:H7	0.24	0.96	1.2	I
	<i>C. albicans</i>	-	1	1.0	NT
<b>Penicillin G+ Ethanolic Cinnamon</b>	<i>S. aureus</i>	1	0.25	1.25	I
	<i>E. coli</i> O157H:H7	0.493333	0.96	1.453333	I

<b>Extract</b>	<i>C. albicans</i>	-	2	2.0	NT
<b>Fluconazole+</b>	<i>S. aureus</i>	-	1	1.0	NT
<b>Ethanollic Cinnamon</b>	<i>E. coli O157H:H7</i>	-	0.96	0.96	NT
<b>Extract</b>	<i>C. albicans</i>	0.6	0.5	1.1	I

Key: FIC<sub>A</sub> = Fractional inhibitory concentration of drug A, FIC<sub>B</sub>= Fractional inhibitory concentration of drug B, A=Additive, S=Synergistic, I = indifference, AN = antagonistic, FICI= Fractional inhibitory concentration index.

**Table 11: Fractional inhibitory concentration index (FICI) of various combinations of gentamicin, penicillin, fluconazole and Aqueous Alum against *S. aureus*, *E. coli O157H:H7*, and *C. albicans*.**

Antimicrobial	Agent	Organisms	FIC <sub>A</sub>	FIC <sub>B</sub>	FICI	S, A, I
<b>GENTAMICIN+</b> <b>ALUM</b>		<i>S. aureus</i>	0.32	0.5	0.82	A
		<i>E. coli O157H:H7</i>	0.32	0.5	0.82	A
		<i>C. albicans</i>	-	2	2.13	NT
<b>PENICILLIN G+</b> <b>ALUM</b>		<i>S. aureus</i>	1.947368	0.5	2.44	I
		<i>E. coli O157H:H7</i>	0.493333	0.5	0.99	A
		<i>C. albicans</i>	-	2	2.0	NT
<b>FLUCONAZOLE+</b> <b>ALUM</b>		<i>S. aureus</i>	-	1	1.1	NT
		<i>E. coli O157H:H7</i>	-	1	1.1	NT
		<i>C. albicans</i>	0.05	1	1.05	I

Key: FIC<sub>A</sub> = Fractional inhibitory concentration of drug A, FIC<sub>B</sub>= Fractional inhibitory concentration of drug B, A=Additive, S=Synergistic, I = indifference, AN = antagonistic, FICI= Fractional inhibitory concentration index.

**Table 12: Fractional inhibitory concentration index (FICI) of various combinations of gentamicin, penicillin, fluconazole and Aqueous cinnamon extract against *S. aureus*, *E. coli O157H:H7*, and *C. albicans*.**

AntimicrobialAgent	Organisms	FIC <sub>A</sub>	FIC <sub>B</sub>	FICI	S, A, IAN
<b>Gentamicin+</b> <b>Aq. Cinnamon Extract</b>	<i>S. aureus</i>	0.32	0.5	0.82	A
	<i>E. coli O157H:H7</i>	0.32	1	1.32	I
	<i>C. albicans</i>	-	2	2.0	NT
<b>Penicillin G+</b>	<i>S. aureus</i>	1	0.25	1.25	I

<b>Aq Cinnamon Extract</b>	<i>E. coli</i> O157H:H7	0.49	1	1.49	I
	<i>C. albicans</i>	-	2	2.0	I
<b>Fluconazole+ Aq. Cinnamon Extract</b>	<i>S. aureus</i>	-	0.5	0.5	NT
	<i>E. coli</i> O157H:H7	-	0.5	0.5	NT
	<i>C. albicans</i>	7.4	0.5	7.9	AN

Key: FIC<sub>A</sub> = Fractional inhibitory concentration of drug A, FIC<sub>B</sub>= Fractional inhibitory concentration of drug B, A=Additive, S=Synergistic, I = indifference, AN = antagonistic, FICI= Fractional inhibitory concentration index.

**Table 13: Fractional inhibitory concentration index (FICI) of various combinations of ethanolic cinnamon extract, Alum, and Aqueous cinnamon extract against *S. aureus*, *E. coli* O157H:H7, and *C. albicans*.**

AntimicrobialAgent	Organisms	FIC <sub>A</sub>	FIC <sub>B</sub>	FICI	S, A, I
<b>AQ. CINNAMON EXTRACT + ALUM</b>	<i>S. aureus</i>	1	1	2	I
	<i>E. coli</i> O157H:H7	1	0.5	1.5	I
	<i>C. albicans</i>	0.5	0.5	1	A
<b>Eth. Cinnamon Extract + Alum</b>	<i>S. aureus</i>	1	1	2	I
	<i>E. coli</i> O157H:H7	1	0.5	1.5	I
	<i>C. albicans</i>	0.5	0.5	1	A

Key: FIC<sub>A</sub> = Fractional inhibitory concentration of drug A, FIC<sub>B</sub>= Fractional inhibitory concentration of drug B, A=Additive, S=Synergistic, I = indifference, AN = antagonistic, FICI= Fractional inhibitory concentration index

### Discussions

The MIC and the MBC values of the aqueous and ethanolic cinnamon extracts was greater against *S. aureus* 25mg/ml and 50mg/ml respectively, and the lowest was with *C. albicans* at 12.5mg/ml, when the same concentrations of antimicrobial agents were applied. Aqueous and ethanolic cinnamon extracts were more active against *C. albicans* with lower MIC value than *S. aureus*. different organisms have different MIC value when cinnamon was used as an antimicrobial agent. This is in accordance with EL Atkiet *al.*, 2019 which showed that the



cinnamon essential oil (EO) resulted to a lower MIC values ranging from 19.53ug/ml to 4.4ug/ml. Also, Raeisi *et al.*, 2015 showed that the MIC value of cinnamon essential oil against *E. coli* and *S aureus* was 2500ug/ml.

when gentamicin was tested on *S. aureus* and *E. coli O157H:H7* it resulted to an MIC and MBC values of 0.25mg/ml and 0.50mg/ml respectively. While Penicillin G against *S. aureus* and *E. coli O157H:H7* resulted to an MICs of 0.19mg/ml and 0.75mg/ml and MBCs of 0.37mg/ml and 1.50mg/ml respectively. Fluconazole against *C. albicans* resulted to an MIC and MBC of 0.05mg/ml and 0.1ng/ml respectively. Hence the study conducted by Jaccobset *al.*, 1998; Dilnessa, 2019; Glajzneret *al.*, 2022 showed that the MIC and methicillin susceptible *Staphylococcus aureus* are between 0.06-128ug/ml, and the MIC of methicillin resistant *Staphylococcus aureus* for penicillin G is between 32-128ug/ml. The MIC of gentamicin methicillin susceptible *Staphylococcus aureus* are between 0.06-16ug/ml, and between 0.06-64ug/ml for methicillin resistant *Staphylococcus aureus*. susceptible *S. aureus* to penicillin G, also means that it is susceptible to other beta lactam antibiotics. Furthermore, a study conducted by Gomes *et al.*, 2019 showed an MIC value between 0.4 -24ug/ml for penicillin G against all strains of *S. aureus*. A study conducted by sabeet *al.*, 2020 showed that the MIC of gentamicin against *S. aureus* is 0.02mg/ml, same MIC with *E. coli*. Landman *et al* 2010 also showed that MIC(90) of gentamicin was 32mg/L against most isolates of *E coli*.

Aqueous Alum was active against the three selected clinical isolates used the MICs of aqueous alum against *S. aureus* and *E. coli O157H:H7* and *C. albicans* were 25mg/ml, 25mg/ml and 12.5mg/ml respectively, and the MBCs aqueous alum against *S. aureus* and *E. coli O157H:H7* and *C. albicans* were 50mg/ml, 50mg/ml and 25mg/ml respectively this showed that the MIC and MBC of alum was low against *C. albicans* compared to *S. aureus*, this is in accordance with

a study conducted by Ali , 2018 revealed that the MIC and MBC of Aqueous Alum against *S. aureus* was 20%, the MIC of alum against *E. coli* was 10% while the MBC was 20%. The same study by Ali , 2018; Lee et al 2013, also revealed that the MIC and MBC of aqueous Alum against *C. albicans* was 10%. The MIC of alum against *E. coli* was reported to be 10mg/ml (Shahriari et al.,2017). Ahmed, 2011 Showed that Alum has different MIC value for different pathogens, and that the MIC value of alum against *Proteus mirabilis* was 0.8g/ml.

### **Interactions Between the Antimicrobials**

The combination of penicillin G, gentamicin, and fluconazole resulted in the additive and indifferent effects with a FICI ranging from 0.56 to 1.02, lowest FICI of 0.56 was with gentamicin and penicillin G against *E. coli O157H:H7* and the highest FICI was 1.37 with the combination of penicillin G and gentamicin against *E. coli O157H:H7*.

For the combination of ethanolic cinnamon extracts with gentamicin, penicillin G and Fluconazole, there was an additive and indifferent effects against the tested clinical isolates with a FICI ranging from 0.72 to 1.45, the lowest FICI of 0.72 obtained from the combination of gentamicin + ethanolic cinnamon extracts against *S aureus*, and the highest FICI of 1.45 obtained from the combination of Penicillin G + Ethanolic cinnamon extracts against *E. coli O157H:H7*. Reports from Utcharyakiatet *et al.*, 2016; Yap *et al.*, 2013 & Mahadletet *et al.*, 2012, showed that combinations of *cinnamomumzeylanicum* and antibiotics expresses additive and synergistic effects against a variety of microorganisms. This is in confirmation with the study conducted by Guerra et al., 2012, that a combination of *cinnamomun zeylanicum* with antibiotics shows an additive effect.

The combinations aqueous Alum Gentamicin Penicillin G and fluconazole tested on the selected clinical isolates resulted to both additive and indifferent effects with a FICI between 0.82 to 2.44. fluconazole + aqueous Alum resulted in an indifferent effect against *C. albicans* with a FICI of 1.05 while the combination of gentamicin + aqueous Alum and combinations of Penicillin G + aqueous Alum produced both additive and indifferent effects against *S. aureus*, *E. coli* O157H:H7.. For the combination aqueous cinnamon extracts with Gentamicin, Penicillin G, and fluconazole against *S. aureus*, *E. coli* O157H:H7, and *C. albicans* produced an additive, indifferent, and antagonistic effects with a FICI ranging from 0.82 to 7.9. The combination of fluconazole + aqueous cinnamon extracts resulted to an antagonistic effect against *C. albicans* with FICI of 7.9. Furthermore, the combinations aqueous Alum, with Ethanolic Aqueous cinnamon extracts produced an additive and indifferent effects against the three tested clinical isolates used in this study, with a FICI ranging from 1 to 2. The combinations produced an additive effect against *C. albicans* and indifferent effects against *S. aureus* and *E. coli* O157H:H7. EL Atkiet *al.*, 2019 showed that the combination of cinnamon essential oil (EO) with streptomycin resulted in an additive effect against strains of *E. coli* and *S. aureus* with FICI of 1.0 to 0.75 respectively. The same study also revealed that the coalescence effects of cinnamon EO+Ampicillin resulted in an indifferent effect against *E. coli* with a FICI of 1.2. this showed that the combination of cinnamon extracts and different conventional antibiotics produced different effects ranging from synergistic to indifferent effects depending on the organism and antibiotics used. Mahadleket *al.*, 2012, also showed that the combination of cinnamon EO + Ciprofloxacin resulted to an additive effect against *S. aureus* using the checkerboard assay. Furthermore, Lu *et al.*, 2011; Utchariyakiat *et al.*, 2016 revealed that the

combination of cinnamon EO + Clove displayed both additive and indifferent effects against a variety of bacteria; both clinical and foodborne bacteria.

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